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Appendix 1: Report purpose

Document purpose	To disseminate the outcomes, recommendations and proposed next steps of
	phase 2 of the Asthma Audit Development Project (AADP)
Title	Asthma Audit Development Project (AADP): Phase 2 final report
Authors	Andrews R, McMillan V, Nasser S, Quint J, Roberts CM (on behalf of the Asthma
	Audit Development Project (AADP) steering committee)
Publication date	August 2018
Audience	Healthcare professionals; NHS managers, chief executives and board members;
	service commissioners; policymakers; voluntary organisations; patient support
	groups; asthma patients and their families/carers; and the public.
	The successful provider of the National Asthma and COPD Audit Programme.
Description	This is the second of the Asthma Audit Development Audit (AADP) reports,
	released by the Royal College of Physicians, London. This report details the
	activities, outcomes and recommendations from phase 2 of the Asthma Audit
	Development Project and proposed next steps in the implementation of a
	National Asthma Audit in England and Wales.
	The report is relevant to anyone with an interest in asthma. It enables lay
	people, as well as experts, to understand the outcomes, recommendations and
	next steps proposed.
Supersedes	None
Related	Asthma Audit Development Project (AADP): Phase 1 final report
publications	https://www.rcplondon.ac.uk/projects/outputs/asthma-audit-development-
	<u>project-information-and-resources</u>
Contact	asthmaauditdevelopmentproject@rcplondon.ac.uk

Appendix 2: Report preparation

This report was prepared and written by the following, on behalf of the Asthma Audit Development Project (AADP) steering committee. Full Steering Committee membership included in Appendix 3.

Ms Rachael Andrews

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Associate Director, Care Quality Improvement Department, Royal College of Physicians, London; Programme Clinical Lead, National COPD Audit Programme; and clinical academic lead for population health, UCL Partners.

Ms Sophie Robinson

Programme Coordinator, National COPD Audit Programme, Care Quality Improvement Department, Royal College of Physicians, London

Mr Philip Stone

Research Assistant in Statistics/Epidemiology, National Heart & Lung Institute, Imperial College London

Appendix 3: Current members of the AADP steering committee

- Shuaib Nasser, Clinical Lead, Asthma Audit Development Project (AADP), Care Quality Improvement Department, Royal College of Physicians, London, Consultant in Allergy and Asthma, Addenbrookes Hospital, Cambridge University Hospitals NHS Foundation Trust
- Rachael Andrews, Project Manager, Asthma Audit Development Project (AADP), Care Quality
 Improvement Department, Royal College of Physicians, London
- Catherine Broadbent, Asthma UK, Senior Insight Analyst
- James Calvert, British Thoracic Society, Consultant Respiratory Physician, North Bristol NHS
 Trust
- **Toby Capstick**, Royal Pharmaceutical Society of Great Britain, Lead Respiratory Pharmacist, Leeds Teaching Hospital NHS Trust
- **Teena Chowdhury**, Operations Director Audit and Accreditation, Care Quality Improvement Department, Royal College of Physicians, London
- Anita Critchlow, National Paediatric Respiratory and Allergy Nurses Group (NPRANG) chair,
 Respiratory Nurse Specialist, Sheffield Children's Hospital NHS Foundation Trust
- Luke Daines, Primary Care Respiratory Society UK (PCRS-UK), Academic Clinical Fellow in General Practice, Usher Institute of Population Health Sciences and Informatics, University of Edinburgh
- **Erol Gaillard**, Royal College of Paediatrics and Child Health, Senior Lecturer in Child Health and Honorary Consultant in Paediatric Respiratory Medicine
- **Lizzie Grillo**, Association for Chartered Physiotherapists in Respiratory Care (ACPRC) chair, Therapy Lead and Specialist Physiotherapist, Royal Brompton Hospital
- **Kevin Gruffydd-Jones**, Respiratory Lead, Royal College of General Practitioners; Clinical Policy Lead, Primary Care Respiratory Society UK; Honorary Lecturer at University of Bath
- Natalie Harper, Association of Respiratory Nurse Specialists (ARNS), Respiratory Advanced Nurse Practitioner, Dorset County Hospital NHS Foundation Trust
- Petronella Hutchinson, Patient Representative
- Richard Iles, Respiratory Paediatrics Consultant, Evelina London Children's Hospital
- **Jeff Keep**, Royal College of Emergency Medicine, Consultant and Honorary Senior Lecturer in Emergency Medicine & Major Trauma, King's College Hospital
- Viktoria McMillan, National COPD Audit Programme Manager, Care Quality Improvement Department, Royal College of Physicians, London
- Daniel Menzies, NHS Wales representative, Consultant Respiratory Physician, Glan Clwyd Hospital
- Andrew Menzies-Gow , Difficult Asthma Registry, Respiratory Medicine Consultant, Royal Brompton Hospital
- Jenni Quint, Clinical Senior Lecturer in Respiratory Epidemiology, Occupational Medicine and Public Health, National Heart and Lung Institute, Imperial College London; and Honorary Consultant Physician in Respiratory Medicine, Royal Brompton Hospital, London
- James Riordan, National COPD Audit Programme Coordinator, Care Quality Improvement Department, Royal College of Physicians, London
- **C Michael Roberts**, National COPD Audit Programme, Programme Clinical Lead; and clinical academic lead for population health, UCL Partners

- Carol Roberts, Chief Executive Officer, PrescQIPP Community Interest Company
- **Sophie Robinson**, National COPD Audit Programme Coordinator, Care Quality Improvement Department, Royal College of Physicians, London
- Robert Spaight, Clinical Audit and Research Manager, East Midlands Ambulance Service NHS
 Trust
- **Simon Standen**, College for Paramedics, Consultant Paramedic Emergency Care, Yorkshire Ambulance Service NHS Trust
- Carol Stonham MBE, Senior Nurse Practitioner Respiratory, Gloucestershire CCG and Primary Care Respiratory Society UK (PCRS-UK) Lead Nurse
- Ralph Sullivan, Clinical Lead for Patient Online, Royal College of General Practitioners
- Ian Woolhouse, Clinical Director Audit and Accreditation, Care Quality Improvement Department, Royal College of Physicians, London

Appendix 4: Participating hospitals in the AADP pilot

Hospitals that submitted clinical and organisational data for the adult and/or paediatric pilot datasets. Those with a * additional submitted a case study on participating in the AADP pilot.

Trust	Hospital	Adult	Paediatric
Alexander of Dec Marine and Little and	Singleton Hospital	✓	✓
Abertawe Bro Morgannwg University Health Board	Morriston Hospital	✓	✓
neatti board	Princess of Wales Hospital	✓	✓
Cambridge University Hospitals NHS Foundation Trust	Addenbrookes Hospital	✓	✓
Cardiff and Vale University Health Board	University Hospital Llandough	✓	✓ (Minimal org submission)
curdin and vale offiversity fleaten board	University Hospital Wales	✓	✓ (Minimal org submission)
Colchester Hospital University NHS Foundation Trust	Colchester General Hospital		✓
Countess of Chester Hospital NHS Foundation Trust	Countess of Chester Hospital	✓	✓
Dartford and Gravesham NHS Trust	Darent Valley Hospital	✓	✓
Derby Teaching Hospitals NHS Foundation Trust	Royal Derby Hospital	✓	√ *
East Kent Hospitals University NHS	Queen Elizabeth The Queen Mother	✓	
Foundation Trust	William Harvey Hospital	✓	
Friedry Hoolth Foundation Trust	Frimley Park Hospital	✓	✓
Frimley Health Foundation Trust	Wexham Park Hospital	✓	✓
Gloucestershire Hospitals NHS Foundation Trust	Gloucestershire Royal Hospital	✓	
Guy's and St Thomas' NHS Foundation Trust	Evelina Children's Hospital		✓
Lancashire Teaching Hospitals NHS Foundation Trust	Royal Preston Hospital	✓ (Clinical data only)	
Maidstone and Tunbuidge Wells NUS Trust	Tunbridge Wells Hospital	✓	✓
Maidstone and Tunbridge Wells NHS Trust	Maidstone Hospital	✓	✓
Manchester University Foundation Trust	Wythenshawe Hospital	✓	✓
Mid-Yorkshire Hospitals NHS Trust	Pinderfields General Hospital	✓	✓
Norfolk and Norwich University Hospital	Norfolk and Norwich	√	✓
NHS Foundation Trust	University Hospital		,
Northumbria Healthcare NHS Foundation	Northumbria Specialist	✓	✓
Trust	Emergency Hospital Oxford University Hospitals		
Oxford University Hospitals NHS	NHS Foundation Trust,		√
Foundation Trust	Children's hospital		,
Portsmouth Hospitals NHS Trust	Queen Alexandra Hospital	√	
Royal Bournemouth and Christchurch	·		
Hospitals NHS Foundation Trust	Royal Bournemouth Hospital	√	

Royal Free London NHS Foundation Trust	Hampstead Hospital	✓	
Royal United Hospitals Bath NHS Foundation Trust	Royal United Hospital, Bath	√ *	✓
Sandwell & West Birmingham Hospitals	City Hospital	✓	✓
NHS Trust	Sandwell Hospital	✓	✓
Sheffield Children's NHS Foundation Trust	Sheffield Children's Hospital		✓
Southend University Hospital NHS Foundation Trust	Southend University Hospital		✓ (Clinical data only)
St Helen's and Knowsley Teaching Hospitals NHS Trust	Whiston Hospital	✓	✓
Tameside and Glossop Integrated Care NHS Foundation Trust	Tameside General Hospital	✓	✓
Taunton and Somerset NHS Foundation Trust	Musgrove Park Hospital	✓	✓ (Minimal org submission)
The Royal Wolverhampton NHS Trust	Newcross Hospital	✓	✓
University Hospital of North Midlands	Royal Stoke University Hospital	✓	✓ (Clinical data only)
University Hospitals Birmingham NHS Foundation Trust	Queen Elizabeth Hospital, Birmingham	√ *	
University Hospitals Bristol NHS	Bristol Royal Hospital for		√
Foundation Trust	Children		·
University Hospitals of Leicester NHS Trust	Leicester Children's Hospital		✓ (Clinical data only)
West Hertfordshire Hospitals NHS Trust	Watford Hospital	√ *	✓

Appendix 5: Consultation respondents

Secondary care clinical datasets: Public consultation

Name	On behalf of:
Angela Cooper	University Hospital of North Midlands,
Aligeia Coopei	Royal Stoke University Hospital
Carol Still Maidstone and Tunbridge Wells NHS Trust	
Jeremy Hull Oxford University Hospitals NHS Foundation Trust	
Kathrun Crahh	Royal Bournemouth and Christchurch Hospitals,
Kathryn Crabb	Royal Bournemouth Hospital
Paul Gilliatt	Northampton General Hospital NHS Trust
Rachel Evens	Abertawe Bro Morgannwg University Health Board
Sara Lock	Whittington Health NHS Trust
Sarah Elkin Imperial College Healthcare NHS Trust	
Coomo Brii	North West Anglia NHS Foundation Trust,
Seema Brij	Peterborough City Hospital
Sharon Sturney	Royal United Hospitals Bath NHS Foundation Trust,
Silaron Sturney	Royal United Hospital, Bath
Simon Gompertz	University Hospitals Birmingham NHS Foundation Trust,
Sillion dompertz	QE Hospital Birmingham
Will McConnell	Dorset County Hospital NHS Foundation Trust,
vviii ivicconnen	Dorset County Hospital

Primary care queries: Targeted consultation

Name	On behalf of:
Deirdre Siddaway	Primary Care Respiratory Society (PCRS)
Duncan Keeley	Royal College of General Practitioners (RCGP)
Durush Attar	Primary Care Respiratory Society (PCRS)
Imran Rafi	Royal College of General Practitioners (RCGP)
John Robinson	Royal College of General Practitioners (RCGP)
Kevin Gruffydd-Jones	Royal College of General Practitioners (RCGP)
Natalie Harper	Association for Respiratory Nurse Specialists (ARNS)
Ralph Sullivan	Royal College of General Practitioners (RCGP)
Val Gerrard	Primary Care Respiratory Society (PCRS)
Wendy Preston	Association for Respiratory Nurse Specialists (ARNS)

Appendix 6: Other acknowledgements

Name	Organisation
Andrew Cumella	Asthma UK
Pooja Panchasara	National Clinical Audit for Rheumatoid and Early Inflammatory Arthritis
Catherine Broadbent	Asthma UK
Paul Davies	NHS Business Services Authority, Prescription Information Services
Margaret Dockey	NHS Business Services Authority, Prescription Information Services
Daisy Ellis	Asthma UK
Emma Fernandez	Royal College of Emergency Medicine
Focus Group attendees	Asthma UK patient and carer representatives
Natasha Dunkley	Health Research Authority
Noel Baxter	Primary Care Respiratory Society, National COPD Audit Programme
Noei baxtei	Clinical Lead
Sam McIntyre	Royal College of Emergency Medicine
Tim Bunning	Crown Informatics

Appendix 7: Secondary care audit: Adult clinical dataset

Secondary Care (Adult)

Clinical audit (adult) data collection sheet

Version: FINAL 1.2 AADP 310118

This dataset should be used and completed for adult asthma patients (age 16 years and over on the date of admission) who have been <u>admitted to hospital adult services</u> with a primary diagnosis of an acute exacerbation of asthma.

Case definition

Any adult (16 and above) who is admitted to hospital with a primary diagnosis of an acute exacerbation of asthma and treated within an adult unit/ward.

Item	Question and answer options	Helpnotes and rationale
1. Gene	ric questions	
1.1	NHS number	Rationale
		Enables linkage
	//	
1.2	Date of birth	Helpnotes
		Do not include asthma patients under the age of 16 or
	/(DD/MM/YYYY)	between the ages of 16-18 (on date of admission) who
		have been treated on a paediatric unit/ward. Please use
		the Secondary Care (Paediatric) Pilot 2017 dataset (6 – 18
		years) for these patients.
		Rationale
		Enables linkage and investigation into demographics of
		asthma patients.
1.3	Home postcode	Rationale
		Enables linkage and investigation into deprivation
	/	demographics.
1.4	Patient GP practice	Rationale
	(Drop down list)	Enables linkage
1.5	Gender (Select one only)	Helpnotes
		Select one only.
	☐ Male	
	☐ Female	Rationale
	☐ Other	Enables investigation into asthma demographics.
1.6	What was the smoking status of the patient, as documented	Helpnotes
	for the current admission? (Tick all that apply)	Tick all that apply.
	☐ Never smoked	Please enter the smoking status as recorded during the
	☐ Ex-smoker	admission.
]		

	☐ Current smoker	If smoking status was not documented, please select 'not
	☐ Vaping only	recorded'.
	□ Not recorded	If the patient stopped smoking at least 4 weeks prior to
	_ 100110001000	the admission, please enter 'ex-smoker'.
		If the patient has stopped within 4 weeks, mark as a
		smoker.
		This question does not just refer to tobacco smokers,
		regular cannabis smokers should also be counted as
		current smokers.
		carrent smokers.
		Rationale
		BTS Adult Asthma Audit
		Included in dataset
		BTS guideline
		Parents with asthma should be advised about the
		dangers, to themselves and to their children with asthma,
		of smoking, and be offered appropriate support to stop
		smoking.
		Clinicians should be aware that higher doses of inhaled
		corticosteroids may be needed in patients who are
		smokers or ex-smokers.
		NRAD
		A history of smoking and/or exposure to second-hand
		smoke should be documented in the medical records of
		all people with asthma. Current smokers should be
		offered referral to a smoking-cessation service.
		This question embraces NICE 2011 QS 5, NICE 2013
		(Smoking: Supporting People to Stop) QS43.
		https://www.nice.org.uk/guidance/qs43.
1.7	Was the patient regularly exposed to smoke at home?	Helpnotes
	(Select one only)	Select one only.
	☐ Yes	This question aims to ascertain if the patient has
	□ No	prolonged or acute exposure to second hand smoke.
	☐ Not recorded	These are associated with an increased risk of developing
		asthma and exacerbations respectively. Please select yes
		if the patient is exposed to second hand smoke in the
		home at least weekly.
		If the exposure stopped at least 4 weeks prior to the
		admission, please enter 'No'.
		Rationale
		NRAD
		A history of smoking and/or exposure to second-hand
		smoke should be documented in the medical records of
		all people with asthma. Current smokers should be
		offered referral to a smoking-cessation service.

2. Pre-ho	ospital care and arrival at hospital	
2.1	In the 12 hours prior to arrival at hospital, were systemic steroids administered to the patient? This includes	Helpnotes Select one only.
	administration by primary care professionals, ambulance or first responder crew. (Select one only) Yes	Only select 'Yes' if systemic steroids were administered within 12 hours prior to arrival to hospital for this exacerbation.
	□ No □ Not known	Administration since the patient became symptomatic for this exacerbation of asthma. Should include administration by healthcare professionals including primary care, ambulance crew and first responder crew. Rationale
		See rationale given for systemic steroids in question 4.2.
2.2	In the 4 hours prior to arrival at hospital, was nebulised β2 agonist administered to the patient? This includes administration by primary care professionals, ambulance or first responder crew. (Select one only)	Helpnotes Select one only. Only select 'Yes' if nebulised β2 agonist was administered within 4 hours prior to arrival to hospital for this
	☐ Yes ☐ No ☐ Not known	exacerbation. Administration since the patient became symptomatic for this exacerbation of asthma. Should include administration by healthcare professionals including primary care, ambulance crew and first responder crew.
		BTS guidance on nebulised salbutamol in acute severe asthma: Give salbutamol 5 mg by oxygen driven nebuliser. Rationale Nebulised salbutamol is recommended for acute severe asthma by BTS guidance. In moderate acute asthma salbutamol can be administered via pMDI and spacer but patients in this category are less likely to require admission.
2.3	Prior to arrival at hospital, was oxygen administered to the patient? This includes administration by primary care professionals, ambulance or first responder crew. (Select one only) Yes No Not known	Helpnotes Select one only. Only select 'Yes' if oxygen was administered prior to arrival to hospital for this exacerbation. Administration since the patient became symptomatic for this exacerbation of asthma. Should include administration by healthcare professionals including
		primary care, ambulance crew and first responder crew. Rationale See rationale given for oxygen in question 4.2.

2.4	Did the patient arrive by ambulance? (Select one only)	Helpnotes
		Select one only
	□ Yes	
	□ No	
	□ Not known	
2.5	What was the date and time of arrival at your hospital?	Helpnotes
	, ,	dd/mm/yyyy
	Date of arrival// (dd/mm/yyyy)	24 hour clock 00:00
	Time of arrival / (24hr clock 00:00)	
		Arrival date/time refers to the patient's first contact with
		the hospital. The point of arrival is most often the
		Emergency Department (ED) or Medical Assessment Unit
		(MAU), although patients do occasionally come straight
		from home/elsewhere into other wards, and you must
		include these cases also. The time is best determined
		from the ambulance transfer sheet, the ED record or the
		MAU, ward arrival record.
		Rationale
		To ensure that analysis can take place against individual
		time-pressured metrics (administration of steroids, β2
		agonists etc) and enables investigation into length of stay
		etc.
3. Acute	observations	
3. Acute 3.1	observations What was the first recorded heart rate for the patient	Helpnotes
		Helpnotes Whole number. Range 30-150 BPM.
	What was the first recorded heart rate for the patient	· ·
	What was the first recorded heart rate for the patient following arrival at hospital?	· ·
	What was the first recorded heart rate for the patient following arrival at hospital?	Whole number. Range 30-150 BPM.
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation.
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation.
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions.
3.1	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM)beats/min (BPM)	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset
	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM) beats/min (BPM) What was the first recorded respiratory rate for the patient	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset Helpnotes
3.1	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM) beats/min (BPM) What was the first recorded respiratory rate for the patient following arrival at hospital? (Whole number. Range 0-60	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset
3.1	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM) beats/min (BPM) What was the first recorded respiratory rate for the patient	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset Helpnotes
3.1	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-150 BPM) beats/min (BPM) What was the first recorded respiratory rate for the patient following arrival at hospital? (Whole number. Range 0-60	Whole number. Range 30-150 BPM. Record the first measured heart rate upon arrival at hospital for this exacerbation. The number of times the heart beats per minute. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset Helpnotes Whole number. Range 0-60 BPM.

		The number of times the person breathes per minute.
		Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions. BTS Adult Asthma Audit Included in dataset
3.3	What was the first recorded oxygen saturation (SpO ₂)	Helpnotes
	measurement for the patient following arrival at hospital?	Whole number. Range 60 – 100%.
	(Whole number. Range 60 – 100%)	Record the first measured SpO ₂ upon arrival at hospital for
	%	this exacerbation.
	☐ Not recorded	
		If 'Not recorded' go straight to Q.3.4
2 2 1	Was this measurement taken whilst the nations was on	Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation RCEM dataset Included in audit questions BTS Adult Asthma Audit Included in dataset BTS guideline Give controlled supplementary oxygen to all hypoxaemic patients with acute severe asthma titrated to maintain an SpO ₂ level of 94–98%. Do not delay oxygen administration in the absence of pulse oximetry but commence monitoring of SaO ₂ as soon as it becomes available.
3.3.1	Was this measurement taken whilst the patient was on	Helpnotes
	supplementary oxygen? (Select one only)	Select one only.
	☐ No, on room air	Do not answer if Q3.3 = 'Not recorded'.
	☐ Yes	
	☐ Not recorded	
3.4	What was the first recorded peak flow measurement (PEF)	Helpnotes
	for the patient following arrival at hospital?	Whole number. Range 30-800.
	(Whole number. Range – 30-800)	

		The best pre-bronchodilator value should be recorded in
	L/min (on arrival)	L/min. Record the first measured peak flow (PEF) upon
	☐ Patient too unwell	arrival at hospital for this exacerbation.
	☐ Not recorded	
		PEF on arrival to hospital must be completed for all
		patients, unless they are too unwell.
		patients, amess they are tee arms
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of
		their symptoms receive an objective measurement of
		severity at the time of presentation.
		BTS guideline
		Throughout the guideline peak flow is one of the
		recommended measurement systems referred to for
		assessing asthma severity.
		RCEM dataset
		Included in audit questions.
		BTS Adult Asthma Audit
		Included in dataset
	What was the patient's previous best PEF?	Helpnotes
	(Whole number. If not recorded, enter predicted. Range –	Whole number. If not recorded, enter predicted. Range
	30-800)	30-800.
	L/min	Previous best according to Personalised Asthma Action
	☐ Not recorded	Plan (PAAP), patient notes or the patient themselves is to
		be given to accompany PEF on arrival. If previous best is
	If previous best PEF = 'Not recorded' please give	not available, predicted should be entered.
	predicted PEF:	not a tanadic) production on our and of circles.
	•	Rationale
	L/min	
		See rationale for question 3.4.
Saction 4	: Acute treatment	
4.1	Was oxygen prescribed for the patient at any point during	Helpnotes
4.1		Helpnotes Select one for prescribed and one for administered only.
4.1	Was oxygen prescribed for the patient at any point during	
4.1	Was oxygen prescribed for the patient at any point during this admission?	
4.1	Was oxygen prescribed for the patient at any point during this admission?	Select one for prescribed and one for administered only.
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only)	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed ☐ Yes	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes	Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription.
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered	Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription. Rationale
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription. Rationale RCEM dataset
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription. Rationale RCEM dataset O2 should be given on arrival to maintain sats of 94-98%
4.1	Was oxygen prescribed for the patient at any point during this admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes	Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription. Rationale RCEM dataset

		Give supplementary oxygen to all hypoxaemic patients with acute severe asthma to maintain an SpO ₂ level of 94-98%. BTS/SIGN 2.6.1 BTS Adult Asthma Audit Included in dataset Enables timings of administration to be measured.
4.2	When was the patient first administered systemic steroids	Helpnotes
	(including oral or IV) following arrival at hospital?	dd/mm/yyyy
	(Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered	24 hour clock 00:00
	alone).	Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not
	Data of stansida / / /dd/nam/www.l	administered.
	Date of steroids / (dd/mm/yyyy) Time of steroids / (24hr clock 00:00)	Date and time should not be before date and time of
	□ Not recorded	arrival.
	□ Not administered	
		Please record the date and time of the first administration
		of systemic steroids. ie. prednislone given orally,
		hydrocortisone IV or (rarely) IM steroids such as
		depomedrone or kenalog upon arrival at hospital for this
		exacerbation.
		Rationale
		NICE quality standard 25, statement 8: Treatment of
		acute asthma
		People aged 5 years or older presenting to a healthcare
		professional with severe or life threatening acute
		exacerbation of asthma receive oral or intravenous
		steroids within 1 hour of presentation.
		Also, BTS guideline.
		To enable analysis of time to steroid administration.
		RCEM dataset Included in dataset
		If not already given before arrival to the ED, steroids
		should be given as soon as possible (within 60 minutes of
		arrival for severe and 4 hours of arrival for moderate).
		BTS guideline
		People aged 5 years or older presenting to a healthcare
		professional with a severe or life-threatening acute
		exacerbation of asthma receive oral or intravenous
		steroids within 1 hour of presentation.
		Give steroids in adequate doses to all patients with an
		acute asthma attack. BTS/SIGN 2.6.1

		BTS Adult Asthma Audit
		Included in dataset
		Enables timings of administration to the measured.
4.3	When was the patient first administered nebulised β2	Helpnotes
	agonist following arrival at hospital?	dd/mm/yyyy
	(Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered	24 hour clock 00:00
	alone).	Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not
	uionej.	
		administered.
	Date of administration// (dd/mm/yyyy)	
	Time of administration / (24hr clock 00:00)	Date and time should not be before date and time of
	☐ Not recorded	arrival.
	☐ Not administered	
		Please record the time and date of the first administration
		of nebulised β2 agonist (eg salbutamol) upon arrival at
		hospital for this exacerbation.
		nospital for this exacerbation.
		BTS guidance on nebulised salbutamol:
		Give nebulised β2 agonist more frequently eg salbutamol
		5 mg up to every 15-30 minutes or 10 mg per hour via
		continuous nebulisation (requires special nebuliser).
		Rationale
		BTS guideline
		_
		In hospital, ambulance and primary care, nebulisers for
		giving nebulised β2 agonist bronchodilators should
		preferably be driven by oxygen BTS/SIGN 9.3.2
		In patients with severe asthma that is poorly responsive
		to an initial bolus dose of β2 agonist, consider continuous
		nebulisation with an appropriate nebuliser. BTS/SIGN
		9.3.2
		RCEM dataset
		Included in dataset
		High dose nebulised β2 agonist bronchodilator should be
		given 10 minutes of arrival at ED.
		Enables timings of administration to be measured.
Continu	F. Povious and discharge	
	5. Review and discharge	Halamataa
5.1	What was the date of discharge from your hospital?	Helpnotes
		dd/mm/yyyy
	Date of discharge / / (dd/mm/yyyy)	24 hour clock 00:00
	Time of discharge / (24hr clock 00:00)	Date should not be before date and time entered into
		sections 1, 2, 3 or 4.
		The date of discharge is usually found at the end of the
		admission record, or on the discharge summary.
		If the patient is discharged onto another hospital, an early
		discharge scheme, hospital at home or community

		asthma scheme, please give the date of discharge from
		your hospital and not the scheme. If the patient self-
		discharged, use date of self-discharge.
		Rationale
		Enables length of stay to be measured.
		BTS Adult Asthma Audit
		Included in dataset
		RCEM dataset
		Included in dataset
		moduce in dataset
5.2	What was the date of death, if the patient died as an	Helpnotes
	inpatient?	dd/mm/yyyy
	mpatient.	24 hour clock 00:00
	Date of death// (dd/mm/yyyy)	24 Hour clock oo.oo
	Time of death/_ (24hr clock 00:00)	Date should not be before date and time of dates and
	Time of death/_ \(\(\frac{1}{2} \) = \(\	times entered into sections 1, 2, 3 or 4.
		times entered into sections 1, 2, 3 or 4.
		Rationale
		BTS Adult Asthma Audit
		Included in dataset
		RCEM dataset
		Included in dataset
		Enables investigation into mortality rates within hospital.
5.3	Was a discharge bundle completed for this admission?	Helpnotes
	(Select one only. If yes, select one only for BTS discharge	Select one only. If yes, select one only for BTS discharge
	bundle)	bundle
	☐ Yes	A structured way of improving discharge processes and
	□ No	care leading to improved patient outcomes. Based on
	☐ Self-discharge	evidence based clinical interventions or actions.
	If yes, was this a BTS discharge bundle?	BTS care bundle for asthma
	☐ Yes	https://www.brit-thoracic.org.uk/standards-of-
	□ No	care/quality-improvement/care-bundle-for-asthma/
	•	
		If 'No' or 'Self-discharge' are selected please still complete
		what elements of a discharge bundle were complete for
		this patient in Q 5.3.1.
		Rationale
		NICE quality standard 25, statement 9: Specialist review
		People admitted to hospital with an acute exacerbation of
		asthma have a structured review by a member of the
		specialist respiratory team before discharge
		BTS Adult Asthma Audit
		Included in dataset

5.3.1	Which of the following elements were undertaken as	Helpnotes
	part of the patient's discharge?	Tick all that apply.
	(Tick all that apply)	
		If any of the discharge elements have not been completed
	Inhaler technique	and/or are not applicable please do not select that the
	☐ Inhaler technique checked	component has been completed.
	4	
	Assessment of medication and adherence	If 'No' or 'self-discharge' are selected (question 5.3)
	☐ Medication issued/classes reviewed	please still complete which elements of a discharge
	☐ Doses (including ICS) reviewed	bundle were completed for this patient (if any).
		buildle were completed for this patient (if unity).
	(increasing/decreasing as necessary)	Do not complete smoking cessation if patient is not
	☐ Importance of adherence to preventer medication	current smoker.
	discussed with patient/family	current smoker.
		Follow up requests
	Personal Asthma Action Plan	Communication directly with a named individual
	☐ Reviewed/Modified existing	responsible for asthma care within the practice, by means
	<u>OR</u>	
	☐ Issued new	of fax or email counts as a request for follow-up.
		If the patient has been asked and/or been provided with
	Triggers and exacerbating factors	the necessary information they need to make/request the
	$\ \square$ Triggers and exacerbating factors documented	follow up appointment(s) themselves within the
		recommended timeframe please select that the
	Smoking cessation	component was completed.
	☐ Referral to stop smoking services	
	☐ Smoking cessation discussed/prescribed	TAPES =
	(validation = if not current smoker to grey out)	Technique and Medication + Action Plan + Environment +
	(vaniation ly not carrent sincial to givey easy	S ubsequent care
	Follow up requests	An acronym used to describe the elements of the British
	☐ In the community within 2 working days	Thoracic Society (BTS) discharge bundle.
	☐ Specialist review within 4 weeks	https://www.brit-thoracic.org.uk/document-library/audit-
	□ Specialist review within 4 weeks	and-quality-improvement/asthma-care-bundle/care-
		bundle-statement/
		Bundle Statement 1:
		All patients (or family members/carers administering
		medicines) should have their inhaler technique assessed
		prior to discharge.
		Bundle Statement 2:
		All patients should have their medications assessed. The
		importance of medication adherence to good asthma
		control should be reinforced to patients (and/or any
		family members or carers administering medicines) prior
		to discharge.
		Bundle Statement 3:
		A written asthma action plan for how to manage care
		should be provided to patients and families/carers.
		Bundle Statement 4:
		Triggering and exacerbating factors in the patient's overall
		environment should be considered.
		Bundle Statement 5:

Subsequent care: follow-up in the community to be arranged within 2 working days plus specialist care according to criteria* within 2 weeks.

* BTS/SIGN British guideline on the management of asthma, sections 8.6.3.

'It is essential that the patient's primary care practice is informed within 24 hours of discharge from the emergency department or hospital following an asthma attack. Ideally this communication should be directly with a named individual responsible for asthma care within the practice, by means of fax or email.'

Rationale

NICE quality standard 25, statement 3: Written

personalised action plans

NICE quality standard 25, statement 4: Inhaler technique NICE quality standard 25, statement 6: Assessing asthma control

NICE quality standard 25, statement 10: Follow up on Primary Care

RCEM dataset

Evidence of assessment before discharge:

The patient's inhaler technique

The patient's inhaler type is satisfactory GP or clinical follow up arranged according to local policy for discharged patients within 2 working days.

BTS guideline

BTS/SIGN 2.4

BTS/SIGN 2.5 and 7.1.

BTS/SIGN 2.2

BTS/SIGN 2.2

BTS/SIGN 8.6.3

(Page 43)

BTS Adult Asthma Audit

Included in dataset

NRAD

Please see NRAD recommendations in Why Asthma Still Kills on:

Personal asthma action plan (PAAP) which details triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency.

Follow-up arrangements for patients who have attended ED or out of hours services.

Factors that trigger or exacerbate asthma.

An assessment of recent asthma control should be undertaken at every asthma review.

Non-adherence to preventer inhaled corticosteroids.

		Health professionals' awareness of factors that increase
		risk of asthma attacks.
		Patient self-management.
		Parents and children, and those who care for or teach
		them, should be educated about managing asthma.
5.4	Was the patient prescribed prednisolone at discharge?	Helpnotes
	(Select one only. If yes, enter mgs per day and number of	Select one only. If yes, enter mgs per day AND number of
	days)	days
		Range 5-60 (mgs) and 1-30 (days).
	☐ Yes	
	□ No	Please record the daily dose and length of the course of
	☐ No, completed course prior to discharge	prednisolone in days. Do not include dose and days prior
	☐ No, patient on maintenance dose of steroids	to discharge.
	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
	If yes,	If self-discharge, select 'No'.
	mgs per day (<i>Range – 5-60</i>)	
	days (Range – 1-30)	Rationale
	==	RCEM dataset
		Included in dataset
		Discharged patients should have oral prednisolone
		prescribed.
		BTS guideline
		Give Steroids in adequate doses for all acute attacks.
		BTS/SIGN 2.6.1
		Continue prednisolone 40–50mg daily for at least 5 days
		or until recovery.
		BTS/SIGN 8.3.3
		BTS Adult Asthma Audit
		Included in dataset

Appendix 8: Secondary care audit: Adult organisation dataset

NEW questions included in the pilot only.

Secondary Care (Adult)

Organisational audit (Adult) data collection sheet

(Version – AADP Final 1.2: 31 January 2018)

All items are compulsory unless specified.

Section 1	Admissions – Number and beds	
	Admission numbers	
1.3	How many emergency coded admissions did your hospital admit in XXXX for asthma:	Whole number only.
	Number Number	This refers to all emergency asthma coded admissions only. A stay in hospital of less than 4 hours would be classed as a non-admission and would not be included. XXXX refers to 1 April – 31 March XXXX inclusive. This may be identified from your hospital's discharge/death codes – ICD10 codes. Consider asking your IT or clinical coding department to help with this. We have already approached your audit department to facilitate the derivation of this metric. Asthma should be the primary diagnosis. The main codes that will be relevant are: J45.2- Mild intermittent J45.3- Mild persistent J45.5- Severe persistent J45.9- Other and unspecified asthma J46 - Acute severe asthma This additionally includes all sub-component codes within these (eg J45.20 uncomplicated, J45.21 with (acute) exacerbation), J45.22 with status asthmaticus). See links below for full list:
		http://www.icd10data.com/ICD10CM/Codes/J00-J99/J40- J47/J45-
1.4	What number of the total emergency asthma coded respiratory admissions (response to 1.3) were discharged from your dedicated respiratory ward(s) in XXXX: Number	Whole number only. This is the number of asthma coded admissions discharged from your dedicated respiratory wards during XXXX. We have already approached your audit department to facilitate the derivation of this metric.
Section 3	Access to specialist staff and 7-day working	
	Access to specialist staff	
3.4	Does your hospital have a designated, named clinical lead for asthma? (Select one only) Yes	Select one only. NRAD
	• No	Every NHS hospital and general practice should have a

		designated, named clinical lead for asthma services,
		responsible for formal training in the management of acute
		asthma.
3.4.1	If yes, is this person responsible for formal training in	Select one only.
	the management of acute asthma? (Select one only)	
	• Tyes	NRAD
	• No	Every NHS hospital and general practice should have a
	_	designated, named clinical lead for asthma services,
		responsible for formal training in the management of acute
		asthma.
	7 day working	
3.9	On which days is a respiratory nurse(s) available to	Tick all that apply.
	review asthma patients: (Tick all that apply)	
	No respiratory nurse(s) available to review	This could be a respiratory nurse specialist, Asthma nurse or
	asthma patients	clinical nurse specialist.
	Monday	
	• Tuesday	If these days can vary, select the days this is most likely to or
	• Wednesday	'usually' happen(s) on.
	•	
	• ☐ Friday	
	• Saturday	
	• Sunday	
Section 5	Integrating care	
5.3	Are you a severe asthma service? (Select one only)	Select one only.
	• Tyes	,
	• No	NICE quality statement 11: Difficult asthma
		People with difficult asthma are offered an assessment by a
		multidisciplinary difficult asthma service.
		NRAD
		Patients with asthma must be referred to a specialist asthma
		service if they have required more than two courses of
		systemic corticosteroids, oral or injected, in the previous 12
		months or require management using British Thoracic Society
		(BTS) stepwise treatment 4 or 5 to achieve goal.
		BTS guideline
		Patients with difficult asthma should be systematically
		evaluated, including:
		- confirmation of the diagnosis of asthma, and
		- identification of the mechanism of persisting symptoms
		and assessment of adherence to therapy.
		This assessment should be facilitated through a dedicated
		multidisciplinary difficult asthma service, by a team
		experienced in the assessment and management of difficult
		asthma.
		https://www.england.nhs.uk/publication/specialised-
		respiratory-services-adult-severe-asthma/

5.3.1	If no, do you have a referral pathway to a severe asthma	Select one only.
	service? (Select one only)	
	• Tyes	See rational above.
	• No	
Section 6	Patient and carer engagement	
6.1	How often is there a formal survey seeking	Select one only.
	patient/carer views on respiratory services undertaken?	
	(Select one only)	Excludes the Friends and Family Test.
	 Continuous (every patient) 	Excludes the Friends and Farmy Festi
	More than 4 times a year	
	•	
	• 1-2 times a year	
	Less than once a year	
	• Never	
6.2	Do you have a strategic group for respiratory services?	Select one only.
	(Select one only)	
	• Yes	This group is defined as consisting of senior clinical and
	• No	management representatives, who meet regularly, set and
	Not known	review targets, implement the stroke strategy and make plans
		for the future of the service.
6.2.1	If yes, does this group have a patient representative?	Select one only.
	(Select one only)	
	• Yes	A patient or carer who attends to provide a user perspective of
	• No	care.
	Not known	
6.3	Are patients able to access their own electronic records	Select one only.
0.0	at your hospital, as part of a collaborative self-	
	management scheme? (Select one only)	This question aims to establish if patients have access to their
	• Tyes	own electronic records routinely, outside of the FOI request
	• No	process, as part of a self-management scheme.
		process, as part of a seif management seneme.
	Not known	
C 4	le communications comice limbed with a specific matient	Colort and only
6.4	Is your respiratory service linked with a specific patient	Select one only.
	support or engagement group? (Select one only)	This are included assumed as the Dritish Long Franchism
	•	This can include groups such as the British Lung Foundations
	• No	'Breathe Easy' groups.
	Not known	
Section 7	Transitional care	
7.1	Do your processes for transitioning young people from	Tick all that apply.
	paediatric to adult services include ensuring that:	
	(Tick all that apply)	NICE guideline: Transition from children's to adults' services
	we do not have any formal transition	for young people using health or social care services
	arrangements	https://www.nice.org.uk/guidance/ng43/chapter/Implementat
	 the young person has a full record of their 	<u>ion-getting-started</u>
	condition	
	 their GP is sent the same record 	Ready Steady Go programme
	 the young person has a transition plan and 	A suite of resources designed to deliver high-quality transition

that they have worked on this with both paediatric and adult clinicians

 the young person has a named case worker to assist in signposting for them and their family for young people (YP) across all subspecialties.

https://www.nice.org.uk/sharedlearning/implementing-transition-care-locally-and-nationally-using-the-ready-steady-go-programme

BTS guideline

TRANSITION TO ADULT-BASED HEALTHCARE

Transition to adult services is important for all adolescents with asthma, irrespective of the asthma severity. No studies on transition of adolescents with asthma to adult services were identified although there are many studies looking at transition of adolescents with chronic illness. Few studies compare different approaches and many recommendations come from consensus statements rather than randomised controlled trials.

It is important that the process of transition is co-ordinated and it is recommended that a healthcare professional be identified to oversee transition and either link with a counterpart in adult services or remain involved until the young person is settled within adult services.

In the initial period after transition to adult services in secondary care, adolescents are best seen by one consultant to build their confidence and encourage attendance.

PREPARATION FOR TRANSITION

Transition should be seen as a process and not just the event of transfer to adult services.

It should begin early, be planned, involve the young person, and be both age and developmentally appropriate (see Table 15).

British guideline on the management of asthma Table 15:

Recommendations for organising transition services (Table 15)

- Young people should be given the opportunity to be seen without their parents/ carers.
- Transition services must address the needs of parents/carers whose role in their child's life is evolving at this time.
- Transition services must be multidisciplinary and multiagency. Optimal care requires a co-operative working relationship between adult and paediatric services, particularly where the young person has complex needs with multiple specialty involvement.
- Co-ordination of transitional care is critical. There should be an identified coordinator who supports the young person until he or she is settled within the adult system.
- Young people should be encouraged to take part in transition/support programmes and/or put in contact

	with other appropriate youth support groups. The
	involvement of adult physicians prior to transfer supports
	attendance and adherence to treatment.
	Transition services must undergo continued evaluation.

Appendix 9: Secondary care audit: Paediatric clinical dataset_1-5 year olds

Secondary Care (Paediatric, 1-5 years)

Clinical audit (Paediatric, 1-5 year olds) data collection sheet Version: FINAL 1.2 AADP 220118

This dataset should be used and completed for paediatric asthma patients (age 1-5 years on the date of admission) who have been <u>admitted to hospital paediatric services</u> for asthma as per the case definition below.

Case definition (1 – 5 year olds)

Paediatric patients:

- Over 12 months old at the time admission to hospital, and
- · Admitted to a paediatric unit/ward, with a primary diagnosis of:
 - a) wheezing (ICD-10 R06.2) which is responsive to salbutamol, or
 - b) asthma

Item	Question and answer options	Helpnotes/Rationale
1. Gen	eric questions	
1.1	NHS number	Rationale
		Enables linkage
	//	
1.2	Date of birth	Helpnotes
		1-5 years
	/(DD/MM/YYYY)	Do not include patients who are under the age of 1 or above
		5 years old. The Secondary Care (paediatric 6-18 years)
		dataset should be used for any paediatric patients 6 years or
		above.
		Rationale
		Enables linkage and investigation into demographics of
		asthma.
1.3	Home postcode	Rationale
		Enables linkage and investigation into deprivation
	/	demographics.
1.4	Patient GP practice	Rationale
	(Drop down list)	Enables linkage
1.5	Gender (Select one only)	Select one only
	☐ Male	Rationale
	☐ Female	Enables investigation into asthma demographics.
	☐ Other	

1.6	Was the patient regularly exposed to smoke at home? (Select one only)	Helpnotes Select one only.
	☐ Yes ☐ No ☐ Not recorded	This question aims to ascertain if the patient has prolonged or acute exposure to second hand smoke. These are associated with an increased risk of developing asthma and exacerbations respectively. Please select yes if the patient is exposed to second hand smoke in the home at least weekly. If the exposure stopped at least 4 weeks prior to the admission, please enter 'No'.
		NRAD A history of smoking and/or exposure to second-hand smoke should be documented in the medical records of all people with asthma. Current smokers should be offered referral to a smoking-cessation service.
2. Pre-	hospital care and arrival at hospital	
2.1	In the 12 hours prior to arrival at hospital, were systemic steroids administered to the patient? This includes administration by primary care professionals, ambulance or first responder crew. (Select one only) Yes No Not known	Helpnotes Select one only. Only select 'Yes' if systemic steroids were administered within 12 hours prior to arrival at hospital for this exacerbation. Administration since the patient became symptomatic for this exacerbation of asthma. Should include administration by healthcare professionals including primary care, ambulance crew and first responder crew. Children between 1-5 years of age should only be given systemic steroids in the event of a severe exacerbation of asthma. Rationale See rationale given for systemic steroids in question 4.2.
2.2	In the 4 hours prior to arrival at hospital, were high dose β2 agonists administered to the patient? This includes administration by primary care professionals, ambulance or first responder crew. (Select one only) Yes No	Helpnotes Select one only. Only select 'Yes' if β2 agonists were administered within 4 hours prior to arrival at hospital for this exacerbation. Administration since the patient became symptomatic for this exacerbation of asthma. Should include administration by healthcare professionals including primary care, ambulance crew and first responder crew.

		BTS guidance on high dose β2 agonist: Children (2-5 years): Moderate attack: β2 agonist 2–10 puffs via spacer ± facemask (given one puff at a time inhaled separately using tidal breathing) Acute severe: β2 agonist 10 puffs via spacer ± facemask or nebulised salbutamol 2.5 mg Life threatening: Nebulised β2 agonist: salbutamol 2.5 mg plus ipratropium bromide 0.25 mg nebulised Rationale Nebulised salbutamol is recommended for acute severe asthma by BTS guidance. In moderate acute asthma salbutamol can be administered via pMDI and spacer but
		patients in this category are less likely to require admission.
2.3	Prior to arrival at hospital, was oxygen administered to	Helpnotes
	the patient, for this exacerbation? This includes	Select one only.
	administration by primary care professionals, ambulance or first responder crew. (Select one only) Yes	Only select 'Yes' if oxygen was administered prior to arrival at hospital for this exacerbation .
	□ No	Administration since the patient became symptomatic for
	□ Not known	this exacerbation of asthma. Should include administration
		by healthcare professionals including primary care ,
		ambulance crew and first responder crew.
		Rationale See rationale given for oxygen in question 4.1.
2.4	Did the patient arrive by ambulance? (Select one only)	Helpnotes
2.7	bid the patient drive by difficulties. (Select one omy)	Select one only
	□ Yes	,
	□ No	
	☐ Not known	
2.5	What was the date and time of arrival at your hospital?	Helpnotes
		dd/mm/yyyy
	Date of arrival / (dd/mm/yyyy) Time of arrival / _ (24hr clock 00:00)	24 hour clock 00:00
		Arrival date/time refers to the patient's first contact with the
		hospital. The point of arrival is most often the Emergency
		Department (ED) or Medication Assessment Unit (MAU),
		although patients do occasionally come straight from home/elsewhere into other wards, and you must include
		these cases also. The time is best determined from the
		ambulance transfer sheet, the ED record or the MAU ward
		arrival record.
		Rationale

		To ensure that analysis can take place against individual time
		pressured metrics (administration of steroids, β2 agonists
		etc) and enables investigation into length of stay etc.
3. Acut	e observations	
3.1	What was the first recorded heart rate for the patient	Helpnotes
	following arrival at hospital?	Whole number. Range 30-250 BPM.
	(Whole number. Range 30-250 BPM)	
		Record the first measured heart rate upon arrival at hospital
	beats/min (BPM)	for this exacerbation.
		The number of times the heart beats per minute.
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of
		their symptoms receive an objective measurement of severity
		at the time of presentation.
		RCEM dataset
		Included in audit questions
		BTS guideline
		Includes pulse (heart) rate in diagrams for assessing and
		recording asthma severity.
		No NICE support.
		BTS Paediatric Asthma Audit
		Included in dataset
3.2	What was the first recorded respiratory rate for the	Helpnotes
	patient following arrival at hospital?	Whole number. Range 0-80 BPM.
	(Whole number. Range 0-80 BPM)	
		Record the first measured respiratory rate upon arrival at
	breaths/min	hospital for this exacerbation.
		The number of times the person breathes per minute.
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of
		their symptoms receive an objective measurement of severity
		at the time of presentation.
		RCEM dataset
		Included in audit questions
		BTS guideline
		Includes respiratory rate in diagrams for assessing and
		recording asthma severity.
		No NICE support.
		BTS Paediatric Asthma Audit
		Included in dataset

What was the first recorded oxygen saturation (Sp0₂) measurement for the patient following arrival at hospital? (Whole number. Range 60 − 100%) % Not recorded Was this measurement taken whilst the patient was on supplementary oxygen? (Select one only)	Helpnotes Whole number. Range 60 – 100%. Record the first measured SpO ₂ upon arrival at hospital for this exacerbation. If 'Not recorded' go straight to section 4. Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation RCEM dataset Included in audit questions BTS Paediatric Asthma Audit Included in dataset BTS guideline Children with life-threatening asthma or SpO ₂ <94% should receive high-flow oxygen via a tight fitting face mask or nasal cannula at sufficient flow rates to achieve normal saturations of 94-98%. Helpnotes Select one only.
☐ No, on room air ☐ Yes ☐ Not recorded	Do not answer if Q3.3 = 'Not recorded'.
l n 4: Acute treatment	
Was oxygen prescribed for the patient at any point during admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes No	Helpnotes Select one for prescribed and one for administered only. Patients likely to require oxygen at any point during their admission should have it formally prescribed. This is to ensure that all patients admitted with an exacerbation of asthma have a prescription of oxygen available to them at all times during their in-patient stay and includes patients who do not use the prescription. Rationale RCEM dataset O2 should be given on arrival to maintain sats of 94-98% O2 should be prescribed on arrival to maintain sats 94-98% BTS guideline Give supplementary oxygen to all hypoxaemic patients with acute severe asthma to maintain an SpO ₂ level of 94-98%. BTS/SIGN 2.6.1
	measurement for the patient following arrival at hospital? (Whole number. Range 60 – 100%) % Not recorded Was this measurement taken whilst the patient was on supplementary oxygen? (Select one only) No, on room air Yes Not recorded 14: Acute treatment Was oxygen prescribed for the patient at any point during admission? (Select one for prescribed and one for administered only) Prescribed Yes No Administered Yes

		BTS Paediatric Asthma Audit
		Included in dataset
		Enables timings of administration to be measured.
4.2	Was the patient administered systemic steroids (oral or	Helpnotes
	IV) following arrival at hospital? (Select one only)	Select one only.
	☐ Yes	Can include prednisolone given orally, hydrocortisone IV or
	□ No	(rarely) IM steroids such as Depomedrone or Kenalog upon
		arrival at hospital for this exacerbation.
	☐ Not recorded	Children between 1-5 years of age should only be given
		systemic steroids in the event of a severe exacerbation of
		asthma.
		If not indicated select 'No'.
		Battanala
		Rationale
		NICE quality standard 25, statement 8: Treatment of acute
		asthma
		People aged 5 years or older presenting to a healthcare
		professional with severe or life threatening acute
		exacerbation of asthma receive oral or intravenous steroids
		within 1 hour of presentation.
		Also, BTS guideline.
		To enable analysis of time to steroid administration.
		RCEM dataset
		Included in dataset
		If not already given before arrival to the ED, steroids should
		be given as soon as possible (within 60 minutes of arrival for
		severe and 4 hours of arrival for moderate).
		BTS guideline
		People aged 5 years or older presenting to a healthcare
		professional with a severe or life-threatening acute
		exacerbation of asthma receive oral or intravenous steroids
		within 1 hour of presentation.
		Give steroids in adequate doses to all patients with an acute
		asthma attack. BTS/SIGN 2.6.1
		Give oral steroids early in the treatment of acute asthma
		attacks in children
		Use a dose of 10 mg prednisolone for children under 2 years
		of age, 20 mg for children aged 2–5 years and 30–40 mg for
		children older than 5 years. Those already receiving
		maintenance steroid tablets should receive 2 mg/ kg
		prednisolone up to a maximum dose of 60 mg.
		Repeat the dose of predisolone in children who vomit and
		consider intravenous steroids in those who are unable to
		retain orally ingested medication.
		BTS/SIGN 9.8.4
		BTS Paediatric Asthma Audit

		Included in dataset
4.3	When was the patient first administered β2 agonist via	Helpnotes
	nebuliser or inhaler + spacer following arrival at hospital?	dd/mm/yyyy
	(Enter date and time <u>OR</u> Not recorded <u>OR</u> Not	24 hour clock 00:00
	administered alone).	Enter ONE of: Date and time OR Not recorded OR Not
	,	administered.
	Date of administration// (dd/mm/yyyy)	danimister edi.
	Time of administration / (24hr clock 00:00) ☐ Not recorded	Date and time should not be before date and time of arrival.
	☐ Not administered	Please record the time and date of the first administration of
		β2 agonist (eg salbutamol) upon arrival at hospital for this
		exacerbation. A pMDI + spacer is the preferred option for
		children except for life-threatening attacks.
		BTS guidance on high dose β2 agonist:
		Children (2-5 years):
		Moderate attack: β2 agonist 2–10 puffs via spacer ± facemask
		(given one puff at a time inhaled separately using tidal
		breathing)
		Acute severe: β2 agonist 10 puffs via spacer ± facemask or
		nebulised salbutamol 2.5 mg
		Life threatening: Nebulised β2 agonist: salbutamol 2.5 mg
		plus ipratropium bromide 0.25 mg nebulised
		Rationale BTS guideline In hospital, ambulance and primary care, nebulisers for giving nebulised β2 agonist bronchodilators should preferably be driven by oxygen BTS/SIGN 9.3.2 In patients with severe asthma that is poorly responsive to an initial bolus dose of β2 agonist, consider continuous nebulisation with an appropriate nebuliser. BTS/SIGN 9.3.2 RCEM dataset Included in dataset High dose nebulised β2 agonist bronchodilator should be given within 10 minutes of arrival at ED. BTS Paediatric Asthma Audit Included in dataset Enables timings of administration to be measured.
5. Revi	ew and discharge	
5.1	What was the date of discharge from your hospital?	Helpnotes
		dd/mm/yyyy
	Date of discharge / / (dd/mm/yyyy)	24 hour clock 00:00
	Time of discharge / (24hr clock 00:00)	Date and time should not be before date and time entered
	, , ,	into sections, 1, 2, 3 or 4.
		The date of discharge is usually found at the end of the

If the patient is discharged onto another hospital, an early discharge scheme, hospital at home or community asthma scheme, please give the date of discharge from your hospital and not the scheme. If the patient was discharged by their parent of guardian, use parental-discharge date. Rationale Enables length of stay to be measured. BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date of death / / (dd/mm/yyyy) Time of death / (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset Enables investigation into mortality rates within hospital.
Enables length of stay to be measured. BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date of death/ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset
Enables length of stay to be measured. BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date of death/ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset
BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset Solve Included in dataset Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date of death/ (dd/mm/yyyy) Time of death/ (24hr clock 00:00) Rationale BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset RCEM dataset Included in dataset
Included in dataset RCEM dataset Included in dataset 8.2 What was the date of death, if patient died as an inpatient? Date of death / (dd/mm/yyyy) Time of death / (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
RCEM dataset Included in dataset 5.2 What was the date of death, if patient died as an inpatient? Date of death/ _ / (dd/mm/yyyy) Time of death/ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
5.2 What was the date of death, if patient died as an inpatient? Date of death / (dd/mm/yyyy) Time of death / _ (24hr clock 00:00) Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
inpatient? dd/mm/yyyy 24 hour clock 00:00 Date of death/ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
inpatient? dd/mm/yyyy 24 hour clock 00:00 Date of death/ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
Date of death/ (dd/mm/yyyy) Time of death/ (24hr clock 00:00) Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
Date of death/_ (dd/mm/yyyy) Time of death/_ (24hr clock 00:00) Date and time should not be before date and time entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
Time of death/_ (24hr clock 00:00) Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
Rationale BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
BTS Adult Asthma Audit Included in dataset RCEM dataset Included in dataset
Included in dataset RCEM dataset Included in dataset
RCEM dataset Included in dataset
Included in dataset
Enables investigation into mortality rates within nospital.
5.3 Was a discharge bundle completed for this admission? Helpnotes
(Select one only. If yes, select one only for BTS discharge Select one only. If yes, select one only for BTS discharge
bundle) bundle.
☐ Yes A structured way of improving discharge processes and care
□ No leading to improved patient outcomes. Based on evidence-
☐ Parental-discharge based clinical interventions or actions.
If yes, was this a BTS discharge bundle? BTS care bundle for asthma
Yes https://www.brit-thoracic.org.uk/standards-of-care/quality-
□ No <u>improvement/care-bundle-for-asthma/</u>
If (Not an (Demontal disabage) are salarted places still
If 'No' or 'Parental-discharge' are selected please still
complete what elements of a discharge bundle were
complete for this patient in Q 5.3.1.
Rationale
Rationale NICE quality standard 25, statement 9: Specialist review
NICE quality standard 25, statement 9: Specialist review

		As below
5.3.1	Which of the following elements were undertaken as	Helpnotes
	part of the patients discharge? (Tick all that apply)	Tick all that apply.
	Inhaler technique	If any of the discharge elements have not been completed
	☐ Inhaler technique checked	and/or are not applicable please do not select that the component has been completed.
	Assessment of medication and adherence	
	☐ Medication issued/classes reviewed	If 'No' or 'Parental-discharge' are selected (question 5.3)
	☐ Doses reviewed (increasing/decreasing as	please still complete which elements of a discharge bundle
	necessary)	were completed for this patient (if any).
	$\ \square$ Importance of adherence to preventer	
	medication discussed with patient/family	Smoking
		Parents with asthma should be advised about the dangers, to
	Personal Asthma Action Plan	themselves and to their children with asthma, of smoking,
	☐ Reviewed/Modified existing	and be offered appropriate support to stop smoking. Advice can include schemes such as Step Right Out
	<u>OR</u>	http://www.stopsmokingleic.co.uk/step-right-out-sign-up/.
	☐ Issued new	intep.//www.stopsmokingieic.co.uk/step-right-out-sigh-up/.
	Triangue and accordanting fratering	Follow up requests
	Triggers and exacerbating factors	Communication directly with a named individual responsible
	☐ Triggers and exacerbating factors documented	for asthma care within the practice, by means of fax or email
	Smoking cessation	counts as a request for follow-up.
	Advice given to or discussion had with parents	If the patient/guardian has been asked and/or been provided
	about smoking	with the necessary information they need to make the follow
	about smoking	up appointment(s) themselves within the recommended
	Following up requests	timeframe please select that the component was completed.
	☐ In the community within 2 working days	
	☐ Paediatric asthma clinic within 4 weeks	TAPES =
	☐ Paediatric respiratory specialist review if there	Technique and Medication + Action Plan + Environment +
	have been life threatening features	Subsequent care An acronym used to describe the elements of the British
		Thoracic Society (BTS) discharge bundle.
		https://www.brit-thoracic.org.uk/document-library/audit-
		and-quality-improvement/asthma-care-bundle/care-bundle-
		statement/
		Bundle Statement 1:
		All patients (or family members/carers administering
		medicines) should have their inhaler technique assessed prior
		to discharge.
		Bundle Statement 2:
		All patients should have their medications assessed. The
		importance of medication adherence to good asthma control
		should be reinforced to patients (and/or any family members
		or carers administering medicines) prior to discharge.
		Bundle Statement 3:
		A written asthma action plan for how to manage care should

be provided to patients and families/carers.

Bundle Statement 4:

Triggering and exacerbating factors in the patient's overall environment should be considered.

Bundle Statement 5:

Subsequent care: follow-up in the community to be arranged within 2 working days plus specialist care according to criteria* within 2 weeks.

* BTS/SIGN British guideline on the management of asthma, sections 8.6.3.

'It is essential that the patient's primary care practice is informed within 24 hours of discharge from the emergency department or hospital following an asthma attack. Ideally this communication should be directly with a named individual responsible for asthma care within the practice, by means of fax or email.'

Rationale

NICE quality standard 25, statement 3: Written personalised action plans

NICE quality standard 25, statement 4: Inhaler technique NICE quality standard 25, statement 6: Assessing asthma control

NICE quality standard 25, statement 10: Follow up on

Primary Care

RCEM dataset

Evidence of assessment before discharge:

The patient's inhaler technique

The patient's inhaler type is satisfactory GP or clinical follow up arranged according to local policy for discharged patients within 2 working days.

BTS guideline

BTS/SIGN 2.4

2.5 and 7.1

BTS/SIGN 2.2 8.6.3

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BTS Paediatric Asthma Audit

Some elements included in dataset

NRAD

Please see NRAD recommendations in Why Asthma Still Kills on:

Personal asthma action plan (PAAP) which details triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency.

Follow-up arrangements for patients who have attended ED or out of hours services.

Factors that trigger or exacerbate asthma.

An assessment of recent asthma control should be

		undertaken at every asthma review. Non-adherence to preventer inhaled corticosteroids. Health professionals' awareness of factors that increase risk of asthma attacks. Patient self-management. Parents and children, and those who care for or teach them, should be educated about managing asthma. BTS Paediatric Asthma Audit Included in dataset
5.4	Was the patient prescribed prednisolone at discharge? (Select one only) Yes No No, completed course prior to discharge No, patient on maintenance steroids	Helpnotes Select one only. If parental discharge, select 'No'. Rationale RCEM dataset Included in dataset Discharged patients should have oral prednisolone prescribed. BTS guideline Give Steroids in adequate doses for all acute attacks. BTS/SIGN 2.6.1 Use a dose of 10 mg prednisolone for children under 2 years of age, 20 mg for children aged 2–5 years and 30–40 mg for children older than 5 years. Those already receiving maintenance steroid tablets should receive 2 mg/kg prednisolone up to a maximum dose of 60 mg. Treatment for up to three days is usually sufficient, but the length of course should be tailored to the number of days necessary to bring about recovery. Tapering is unnecessary unless the course of steroids exceeds 14 days. BTS/SIGN 9.8.4 BTS Paediatric Asthma Audit Included in dataset

Appendix 10: Secondary care audit: Paediatric clinical dataset_6-18 year olds

Secondary Care (Paediatric, 6 – 18 years)

Clinical audit (Paediatric, 6 – 18 years old) data collection sheet Version: FINAL 1.2 AADP 310118

This dataset should be used and completed for paediatric asthma patients (6-18 years old on the date of admission) who have been <u>admitted to hospital paediatric services</u> for asthma as per the case definition below.

Case definition

Paediatric patients between 6 and 18 years old:

· Admitted to a paediatric unit/ward, with a primary diagnosis of acute exacerbation of asthma.

Item	Question and answer options	Helpnotes/Rationale
1. Gen	eric questions	
1.1	NHS number	Rationale Enables linkage
1.2	Date of birth	Helpnotes
1.2	/(DD/MM/YYYY)	Do not include patients under the age of 6. The Secondary Care (paediatric – 1-5 year) dataset should be completed for any patients below 6 years of ages. Include patients between the ages of 16 and 18 who are treated on a paediatric ward/unit.
		For patients between the ages of 16 and 18 who are treated on an adult ward/unit please use the Secondary Care (adult) dataset.
		Rationale Enables linkage and investigation into demographics of asthma.
1.3	Home postcode	Rationale
		Enables linkage and investigation into deprivation
	/	demographics.
1.4	Patient GP practice	Rationale
	(Drop down list)	Enables linkage
1.5	Gender (Select one only)	Select one only
	│	Rationale
	☐ Female	Enables investigation into asthma demographics.
	☐ Other	
1.6	Was the patient regularly exposed to smoke at home?	Helpnotes
	(Select one only)	Select one only.

	☐ Yes	This question aims to ascertain if the patient has prolonged or
	□ No	acute exposure to second hand smoke. These are associated
	☐ Not recorded	with an increased risk of developing asthma and exacerbations
		respectively. Please select yes if the patient is exposed to
		second hand smoke in the home at least weekly.
		If the exposure stopped at least 4 weeks prior to the admission,
		please enter 'No'.
		•
		Rationale
		NRAD
		A history of smoking and/or exposure to second-hand smoke
		should be documented in the medical records of all people
		with asthma. Current smokers should be offered referral to a
		smoking-cessation service.
		Showing cessation service.
2. Pre-	hospital care and arrival at hospital	
2.1	In the 12 hours prior to arrival at hospital, were systemic	Helpnotes
	steroids administered to the patient? This includes	Select one only.
	administration by primary care professionals,	
	ambulance or first responder crew. (Select one only)	Only select 'Yes' if systemic steroids were administered within
		12 hours prior to arrival at hospital for this exacerbation.
	☐ Yes	
	□ No	Administration since the patient became symptomatic for this
	☐ Not known	exacerbation of asthma. Should include administration by
		healthcare professionals including primary care, ambulance
		crew and first responder crew.
		Rationale
		See rationale given for systemic steroids in question 4.2.
2.2	In the 4 hours prior to arrival at hespital wars high dose	Helmotes
2.2	In the 4 hours prior to arrival at hospital, were high dose	Helpnotes Select one only.
	β2 agonist administered to the patient? This includes administration by primary care professionals,	Select one only.
	ambulance or first responder crew. (Select one only)	Only select 'Yes' if high dose \(\beta \) agonists were administered
	ambulance of mist responder crew. (Select one omy)	within 4 hours prior to arrival at hospital for this exacerbation.
	□ Yes	within 4 hours prior to arrival at hospital for this exactination.
	□ No	Administration since the patient became symptomatic for this
		exacerbation of asthma. Should include administration by
	☐ Not known	healthcare professionals including primary care, ambulance
		crew and first responder crew.
		crew and instresponder crew.
		BTS guidance on high dose β2 agonist:
		Children (>5 years):
		Moderate attack: β2 agonist 2–10 puffs via spacer and
		mouthpiece (given one puff at a time inhaled separately using
		tidal breathing)
		Acute severe: β2 agonist 10 puffs via spacer or nebulised
		salbutamol 5 mg
		Life threatening: Nebulised β2 agonist: salbutamol 5 mg plus

		ipratropium bromide 0.25 mg nebulised
		Rationale See rationale given for high dose β2 agonists n question 4.3.
2.3	Prior to arrival at hospital, was oxygen administered to the patient, for this exacerbation? This includes	Helpnotes Select one only.
	administration by primary care professionals, ambulance or first responder crew. (Select one only)	Only select 'Yes' if oxygen was administered prior to arrival at hospital for this exacerbation .
	☐ Yes ☐ No ☐ Not known	Administration since the patient became symptomatic for this exacerbation of asthma . Should include administration by healthcare professionals including primary care , ambulance crew and first responder crew .
		Rationale See rationale given for oxygen in question 4.1.
2.4	Did the patient arrive by ambulance? (Select one only) ☐ Yes ☐ No ☐ Not known	Helpnotes Select one only
2.5	What was the date and time of arrival at your hospital? Date of arrival / (dd/mm/yyyy) Time of arrival / (24hr clock 00:00)	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Arrival date/time refers to the patient's first contact with the hospital. The point of arrival is most often the ED or MAU, although patients do occasionally come straight from home/elsewhere into other wards, and you must include these cases also. The time is best determined from the ambulance transfer sheet, the ED record or the MAU, ward arrival record. Rationale To ensure that analysis can take place against individual time pressured metrics (administration of steroids, β2 agonists etc) and enables investigation into length of stay etc.
3. Acut	e observations	
3.1	What was the first recorded heart rate for the patient following arrival at hospital? (Whole number. Range 30-250 BPM)	Helpnotes Whole number. Range 30-250 BPM.
	beats/min (BPM)	Record the first measured heart rate upon arrival at hospital for this exacerbation.
		The number of times the heart beats per minute.

		Rationale NICE quality standard 25, statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. RCEM dataset Included in audit questions BTS guideline Includes pulse (heart) rate in diagrams for assessing and recording asthma severity. No NICE support. BTS Paediatric Asthma Audit Included in dataset
3.2	What was the first recorded respiratory rate for the patient following arrival at hospital? (Whole number. Range 0-80 BPM)	Helpnotes Whole number. Range 0-80 BPM. Record the first measured respiratory rate upon arrival at
	breaths/min	hospital for this exacerbation.
		The number of times the person breathes per minute.
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the
		time of presentation.
		RCEM dataset
		Included in audit questions
		BTS guideline
		Includes respiratory rate in diagrams for assessing and
		recording asthma severity.
		No NICE support. BTS Paediatric Asthma Audit
		Included in dataset
3.3	What was the first recorded oxygen saturation (SpO ₂)	Helpnotes
	measurement for the patient following arrival at	Whole number. Range 60 – 100%.
	hospital?	
	(Whole number. Range 60 – 100%)	Record the first measured SpO ₂ upon arrival at hospital for this exacerbation.
	%	
	☐ Not recorded	If 'Not recorded' go straight to Q.3.4
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of their
		symptoms receive an objective measurement of severity at the

		time of presentation RCEM dataset
		Included in audit questions
		BTS Paediatric Asthma Audit
		Included in dataset
		BTS guideline
		Children with life-threatening asthma or SpO ₂ <94% should
		receive high-flow oxygen via a tight fitting face mask or nasal
		cannula at sufficient flow rates to achieve normal saturations
		of 94-98%.
3.3.1	Was this measurement taken whilst the patient	Helpnotes
	was on supplementary oxygen?	Select one only.
	(Select one only.)	
	_	Do not answer if Q3.3 = 'Not recorded'.
	☐ No, on room air	
	☐ Yes	
	☐ Not recorded	
3.4	What was the first recorded peak flow measurement	Helpnotes
	(PEF) for this patient following arrival at hospital?	Whole number. Range 30-800.
	(Whole number. Range – 30-800)	
		The best pre-bronchodilator value should be recorded in L/min.
	L/min (on arrival)	Record the first measured peak flow (PEF) upon arrival at
	☐ Patient too unwell	hospital for this exacerbation.
	☐ Not recorded	
		PEF on arrival to hospital must be completed for all patients,
		unless they are too unwell.
		Rationale
		NICE quality standard 25, statement 7: Assessing severity
		People with asthma who present with an exacerbation of their
		symptoms receive an objective measurement of severity at the
		time of presentation.
		BTS guideline
		Throughout the guideline peak flow is the recommended
		measurement system reference to for assessing asthma
		severity.
		RCEM dataset
		Included in audit questions.
		BTS Adult Asthma Audit
		Included in dataset
3.4.1	What was the patient's previous best PEF?	Helpnotes
	(Whole number. If not recorded, enter predicted.	Whole number. If not recorded, enter predicted. Range 30-800.
	Range – 30-800)	
		Previous best according to Personalised Asthma Action Plan
	L/min	(PAAP), patient notes or the patient themselves is to be given
	☐ Not recorded	to accompany PEF on arrival. If previous best is not available,

		predicted should be entered.
	If previous best = 'Not recorded' please give	
	predicted PEF:	Rationale
	L/min	See rationale for question 3.4.
-		
	4: Acute treatment	I
4.1	Was oxygen prescribed for the patient at any point	Helpnotes
	during admission?	Select one for prescribed and one for administered only.
	(Select one for prescribed and one for administered only)	
		Patients likely to require oxygen at any point during their
	Prescribed 	admission should have it formally prescribed. This is to ensure
	Yes	that all patients admitted with an exacerbation of asthma have
	□ No	a prescription of oxygen available to them at all times during
		their in-patient stay and includes patients who do not use the
	Administered	prescription.
	☐ Yes	Rationale
	□ No	RCEM dataset
		O2 should be given on arrival to maintain sats of 94-98%
		O2 should be prescribed on arrival to maintain sats 94-98%
		BTS guideline
		Give supplementary oxygen to all hypoxaemic patients with
		acute severe asthma to maintain an SpO ₂ level of 94-98%.
		BTS/SIGN 2.6.1
		BTS Paediatric Asthma Audit Included in dataset
		Enables timings of administration to be measured.
4.2	When was the patient first administered systemic	Helpnotes
	steroids (including oral or IV) following arrival at	dd/mm/yyyy
	hospital?	24 hour clock 00:00
	(Enter date and time <u>OR</u> Not recorded <u>OR</u> Not	Enter ONE of: Date and time OR Not recorded OR Not
	administered alone).	administered alone.
	Date of steroids / / (dd/mm/yyyy)	Date and time should not be before date and time of arrival.
	Time of steroids / _ (24hr clock 00:00)	
	☐ Not recorded	Please record the date and time of the first administration of
	☐ Not administered	systemic steroids. i.e. prednisolone given orally, hydrocortisone
		IV or (rarely) IM steroids such as Depomedrone or Kenalog
		upon arrival at hospital for this exacerbation.
		Rationale
		NICE quality standard 25, statement 8: Treatment of acute
		asthma
		People aged 5 years or older presenting to a healthcare
		professional with severe or life threatening acute exacerbation
		of asthma receive oral or intravenous steroids within 1 hour of
		presentation.
		Also, BTS guideline.
		To enable analysis into time to administration of steroids
		To chasic analysis into time to duffillistration of sterolas

		RCEM dataset Included in dataset
		If not already given before arrival to the ED, steroids should be
		given as soon as possible (within 60 minutes of arrival for
		severe and 4 hours of arrival for moderate).
		BTS guideline
		People aged 5 years or older presenting to a healthcare
		professional with a severe or life-threatening acute
		exacerbation of asthma receive oral or intravenous steroids
		within 1 hour of presentation.
		Give steroids in adequate doses to all patients with an acute
		asthma attack. BTS/SIGN 2.6.1
		Give oral steroids early in the treatment of acute asthma
		attacks in children
		Use a dose of 10 mg prednisolone for children under 2 years of
		age, 20 mg for children aged 2–5 years and 30–40 mg for
		children older than 5 years. Those already receiving
		maintenance steroid tablets should receive 2 mg/kg
		prednisolone up to a maximum dose of 60 mg.
		Repeat the dose of prednisolone in children who vomit and
		consider intravenous steroids in those who are unable to retain
		orally ingested medication.
		BTS/SIGN 9.8.4
		BTS Paediatric Asthma Audit
		Included in dataset
		Enables timings of administration to the measured.
		Enables timings of administration to the measured.
4.3	When was the patient been first administered high dose	Enables timings of administration to the measured. Helpnotes
4.3	When was the patient been first administered high dose β2 agonist via nebuliser or inhaler + spacer following	
4.3		Helpnotes
4.3	$\beta 2$ agonist via nebuliser or inhaler + spacer following	Helpnotes dd/mm/yyyy
4.3	$\beta 2$ agonist via nebuliser or inhaler + spacer following arrival at hospital?	Helpnotes dd/mm/yyyy 24 hour clock 00:00
4.3	β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not
4.3	β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not
4.3	β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration//(dd/mm/yyyy)	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone.
4.3	β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00)	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone. Date and time should not be before date and time of arrival.
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes $dd/mm/yyyy$ 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose $\beta 2$ agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children.
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist:
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years):
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and mouthpiece (given one puff at a time inhaled separately using
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and mouthpiece (given one puff at a time inhaled separately using tidal breathing)
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and mouthpiece (given one puff at a time inhaled separately using tidal breathing) Acute severe: β2 agonist 10 puffs via spacer or nebulised
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and mouthpiece (given one puff at a time inhaled separately using tidal breathing) Acute severe: β2 agonist 10 puffs via spacer or nebulised salbutamol 5 mg
4.3	 β2 agonist via nebuliser or inhaler + spacer following arrival at hospital? (Enter date and time <u>OR</u> Not recorded <u>OR</u> Not administered alone). Date of administration/ (dd/mm/yyyy) Time of administration/_ (24hr clock 00:00) Not recorded 	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Enter ONE of: Date and time OR Not recorded OR Not administered alone. Date and time should not be before date and time of arrival. Please record the time and date of the first administration of high dose β2 agonist (eg salbutamol) upon arrival at hospital for this exacerbation. A pMDI + spacer is the preferred option for children. BTS guidance on high dose β2 agonist: Children (>5 years): Moderate attack: β2 agonist 2–10 puffs via spacer and mouthpiece (given one puff at a time inhaled separately using tidal breathing) Acute severe: β2 agonist 10 puffs via spacer or nebulised

		Rationale BTS guideline In hospital, ambulance and primary care, nebulisers for giving nebulised β2 agonist bronchodilators should preferably be driven by oxygen BTS/SIGN 9.3.2 In patients with severe asthma that is poorly responsive to an initial bolus dose of β2 agonist, consider continuous nebulisation with an appropriate nebuliser. BTS/SIGN 9.3.2 RCEM dataset Included in dataset High dose nebulised β2 agonist bronchodilator should be given within 10 minutes of arrival at ED. BTS Paediatric Asthma Audit Included in dataset Enables timings of administration to be measured.
5. Revi	ew and discharge	
5.1	What was the date of discharge from your hospital? Date of discharge / / (dd/mm/yyyy) Time of discharge / (24hr clock 00:00)	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date should not be before date and time entered into sections, 1, 2, 3 or 4. The date of discharge is usually found at the end of the admission record, or on the discharge summary. If the patient is discharged onto another hospital, an early discharge scheme, hospital at home or community scheme, please give the date of discharge from your hospital and not the scheme. If the patient discharged themselves or their parent/guardian discharged them, use date of self/parental-discharge. Rationale Enables length of stay to be measured. BTS Paediatric Asthma Audit Included in dataset RCEM dataset Included in dataset
5.2	What was the date of death, if patient died as an inpatient? Date of death// (dd/mm/yyyy) Time of death/ (24hr clock 00:00)	Helpnotes dd/mm/yyyy 24 hour clock 00:00 Date should not be before date and entered into sections 1, 2, 3 or 4. Rationale BTS Adult Asthma Audit

		Included in dataset
		RCEM dataset
		Included in dataset
		Enables investigation into mortality rates within hospital.
5.3	Was a discharge bundle completed for this admission?	<u>Helpnotes</u>
	(Select one only. If yes, select one only for BTS discharge	Select one only. If yes, select one only for BTS discharge
	bundle)	bundle.
	□ Yes	A structured way of improving discharge processes and care
	□ No	leading to improved patient outcomes. Based on evidence
	☐ Self/Parental-discharge	based clinical interventions or actions.
	,	
	If yes, was this a BTS discharge bundle?	BTS care bundle for asthma
	□ Yes	https://www.brit-thoracic.org.uk/standards-of-care/quality-
	□ No	improvement/care-bundle-for-asthma/
		If 'No' or 'Self/Parental-discharge' are selected please still
		complete what elements of a discharge bundle were complete
		for this patient in Q 5.3.1.
		Rationale
		NICE quality standard 25, statement 9: Specialist review
		People admitted to hospital with an acute exacerbation of
		asthma have a structured review by a member of the specialist
		respiratory team before discharge
		As below
5.3.1	Which of the following elements were undertaken as	Helpnotes
	part of the patients discharge? (Tick all that apply)	Tick all that apply.
	Inhaler technique	If any of the discharge elements have not been completed
	\square Inhaler technique checked	and/or are not applicable please do not select that the
		component has been completed.
	Assessment of medication and adherence	
	☐ Medication issued/classes reviewed	If 'No' or 'self/Parental-discharge' are selected (question 5.3)
	$\ \square$ Doses reviewed (increasing/decreasing as	please still complete which elements of a discharge bundle
	necessary)	were completed for this patient (if any).
	☐ Importance of adherence to preventer	
	medication discussed with patient/family	Smoking
		Parents with asthma should be advised about the dangers, to
	Personal Asthma Action Plan	themselves and to their children with asthma, of smoking, and
	☐ Reviewed/Modified existing	be offered appropriate support to stop smoking.
	<u>OR</u>	Advice can include schemes such as Step Right Out
	☐ Issued new	http://www.stopsmokingleic.co.uk/step-right-out-sign-up/.
		Follow up requests
	Triggers and exacerbating factors	
	Triggers and exacerbating factors ☐ Triggers and exacerbating factors documented	Communication directly with a named individual responsible
		Communication directly with a named individual responsible for asthma care within the practice, by means of fax or email
		Communication directly with a named individual responsible

about smoking	with the necessary information they need to make the follow
☐ Referral to stop smoking services	up appointment(s) themselves within the recommended
☐ If current smoker, smoking cessation	timeframe please select that the component was completed.
discussed/prescribed	
	TAPES =
Following up requests	Technique and Medication + Action Plan + Environment +
\square In the community within 2 working days	S ubsequent care
☐ Paediatric asthma clinic within 4 weeks	An acronym used to describe the elements of the British
☐ Paediatric respiratory specialist review if there	Thoracic Society (BTS) discharge bundle.
have been life-threatening features	https://www.brit-thoracic.org.uk/document-library/audit-and-
g .	quality-improvement/asthma-care-bundle/care-bundle-
	statement/
	Bundle Statement 1:
	All patients (or family members/carers administering
	medicines) should have their inhaler technique assessed prior
	to discharge.
	Bundle Statement 2:
	All patients should have their medications assessed. The
	importance of medication adherence to good asthma control
	should be reinforced to patients (and/or any family members
	or carers administering medicines) prior to discharge.
	Bundle Statement 3:
	A written asthma action plan for how to manage care should
	be provided to patients and families/carers.
	Bundle Statement 4:
	Triggering and exacerbating factors in the patient's overall
	environment should be considered.
	Bundle Statement 5:
	Subsequent care: follow-up in the community to be arranged within 2 working days plus specialist care according to criteria*
	within 2 weeks.
	WITHIN 2 WEEKS.
	* BTS/SIGN British guideline on the management of asthma,
	sections 8.6.3.
	'It is essential that the patient's primary care practice is
	informed within 24 hours of discharge from the emergency
	department or hospital following an asthma attack. Ideally this
	communication should be directly with a named individual
	responsible for asthma care within the practice, by means of
	fax or email.'
	Rationale
	NICE quality standard 25, statement 3: Written personalised
	action plans
	NICE quality standard 25, statement 4: Inhaler technique
	NICE quality standard 25, statement 6: Assessing asthma
	control
	NICE quality standard 25, statement 10: Follow up on Primary
	•

		Care
		RCEM dataset
		Evidence of assessment before discharge:
		The patient's inhaler technique
		The patient's inhaler type is satisfactory
		GP or clinical follow up arranged according to local policy for
		discharged patients within 2 working days.
		BTS guideline
		BTS/SIGN 2.4
		2.5 and 7.1
		BTS/SIGN 2.2
		8.6.3
		(Page 43)
		BTS Paediatric Asthma Audit
		Some elements included in dataset
		NRAD
		Please see NRAD recommendations in Why Asthma Still Kills
		on:
		Personal asthma action plan (PAAP) which details triggers and
		current treatment, and specifies how to prevent relapse and
		when to seek help in an emergency.
		Follow-up arrangements for patients who have attended ED or
		out of hours services.
		Factors that trigger or exacerbate asthma.
		An assessment of recent asthma control should be undertaken
		at every asthma review.
		Non-adherence to preventer inhaled corticosteroids.
		Health professionals' awareness of factors that increase risk of
		asthma attacks.
		Patient self-management.
		Parents and children, and those who care for or teach them,
		should be educated about managing asthma.
		BTS Paediatric Asthma Audit
		Included in dataset
5.4	Mas the patient prescribed produitaless at discharge	Helmotes
5.4	Was the patient prescribed prednisolone at discharge?	Helpnotes
	(Select one only. If yes, enter mgs per day and number of	Select one only. If yes, enter mgs per day AND number of days.
	days)	Range 5-60 (mgs) and 1-30 (days).
	□ Yes	Complete dose AND days OR No, completed course prior to
	□ No	discharge OR No OR maintenance dose.
	□ No, completed course prior to discharge	
	· · · · · · · · · · · · · · · · · · ·	Please record the daily dose and length of the course of
	☐ No, patient on maintenance steroids	prednisolone in days. Do not include dose and days prior to
	If yes	discharge.
	If yes,	
	mgs per day (Range – 5-60) days (Range – 1-30)	If self or parental discharge, select 'No'.
	days (name = 1 30)	_
		Rationale

RCEM dataset

Included in dataset

Discharged patients should have oral prednisolone prescribed.

BTS guideline

Give Steroids in adequate doses for all acute attacks. BTS/SIGN 2.6.1

Use a dose of 10 mg prednisolone for children under 2 years of age, 20 mg for children aged 2–5 years and 30–40 mg for children older than 5 years. Those already receiving maintenance steroid tablets should receive 2 mg/kg prednisolone up to a maximum dose of 60 mg.

Treatment for up to three days is usually sufficient, but the length of course should be tailored to the number of days necessary to bring about recovery. Tapering is unnecessary unless the course of steroids exceeds 14 days. BTS/SIGN 9.8.4

BTS Paediatric Asthma Audit

Included in dataset

Appendix 11: Secondary care audit: Paediatric organisational dataset

Secondary Care (Paediatric)

Organisational audit (Paediatric) data collection sheet

(Version – AADP Final 1.1: 31 January 2018)

All items are compulsory unless stated otherwise.

Section 1	Admissions – Number and beds		
	Admission numbers		
1.1	How many paediatric medical emergencies did your hospital admit in XXXX:	Whole numbers only.	
	Number	This refers to emergency admissions only.	
	Training of	Please include all paediatric medical emergency admissions for your hospital.	
		Please include admissions to a medical admissions ward via Accident and Emergency Department.	
		Adults (>18 years of age) are not included. Adolescents between 16 and 18 years old are included if they are admitted	
		to and treated on a paediatric ward. A stay in hospital of less than 4 hours would be classed as a	
		non-admission and would not be included. These patients may be identified from your hospital's discharge/death codes using the appropriate ICD10 codes.	
		Consider asking your IT or clinical coding department to help with this.	
		XXXX means from 1 April—31 March XXXX.	
1.2	How many paediatric respiratory coded emergencies did your hospital admit in XXXX	Whole numbers only.	
	Number	This refers to all primary emergency respiratory coded admissions only. This will include respiratory coded admissions not admitted to the respiratory ward, as well as those admitted to the respiratory ward. Codes J00-J99 should be included. Adults (>18 years of age) are not included.	
1.3	How many paediatric asthma emergency coded admissions did your hospital admit in XXXX?	Whole numbers only.	
	Number	This refers to all emergency paediatric asthma coded admissions only. A stay in hospital of less than 4 hours would be classed as a non-admission and would not be included. XXXX refers to 1 April—31 March XXXX inclusive. This may be identified from your hospital's discharge/death codes—ICD10 codes. Consider asking your IT or clinical coding department to help with this. We have already approached your audit department to facilitate the derivation of this metric. Asthma should be primary diagnosis. The main codes that will be relevant are:	

JAS.3- Mild persistent JAS.4- Moderate persistent JAS.5- Severe persistent JAS.5- Severe persistent JAS.5- Other and unspecified asthma JAG- Acute severe asthma This additionally includes all sub-component codes within these (e.g., JAS.20 uncomplicated, JAS.21 with facute) exacerbation), JAS.22 with status asthmaticus). See links below for full list: http://www.icd10dsta.com/iCD10CM/Codes/100-J99/JAD-JA7/JAS- AND RMS.2- Wheeze. For children between the ages of 1-5 only Advits (>15 years of age) who are treated on an odult ward are not included. Whole numbers only: admissions (response to 1.3) were discharged from your dedicated paediatric ward(s) in XXXX. Number Whole numbers only: This is the number of paediatric asthma coded admissions discharged from your dedicated paediatric/respiratory wards in XXXX. We have already approached your audit department to facilitate the derivation of this metric. Adults >15 years of age) who are treated on an adult ward are not included. Whole numbers only: This is the number of paediatric asthma patients? Number Whole numbers only:			J45.2- Mild intermittent
145.5 - Severe persistent 145.9 - Other and unspecified asthma 146 - Acute severe asthma 147 - Acute severe as the severe acute 147 - Acute severe 147			J45.3- Mild persistent
M5.9- Other and unspecified asthma A46 - Acute severe asthma This additionally includes all sub-component codes within these (e.g., I45.20 uncomplicated, I45.21 with (acute) exacerbation), I45.22 with status asthmaticus). See links below for full list: http://www.icd10data.com/ICD10CM/Codes/J00-J99/J40-J47/J45- AND R06.2 - Wheeze. For children between the ages of 1-5 only Adults (5-16 years of age) who are treated on an adult ward are not included. 1.4			J45.4- Moderate persistent
A6 - Acute severe asthma This additionally includes all sub-component codes within these (e.g., IAS. 20 uncomplicated, I45.21 with (acute) exacerbation), I45.22 with status asthmaticus). See links below for full list: http://www.icd10data.com/iCD10CM/Codes/J00-J99/J40-J47/J45-See links below are treated on an adult ward are not included. Abed See for admitted paediatric patients Beds for admitted paediatric patients Whole numbers only.			J45.5- Severe persistent
A6 - Acute severe asthma This additionally includes all sub-component codes within these (e.g., IAS. 20 uncomplicated, I45.21 with (acute) exacerbation), I45.22 with status asthmaticus). See links below for full list: http://www.icd10data.com/iCD10CM/Codes/J00-J99/J40-J47/J45-See links below are treated on an adult ward are not included. Abed See for admitted paediatric patients Beds for admitted paediatric patients Whole numbers only.			J45.9- Other and unspecified asthma
This additionally includes all sub-component codes within these (e.g., 145, 20 uncomplicated, J45, 21 with (acute) exacerbation), J45, 22 with status asthmaticus). See links below for full list: http://www.icd10data.com/iCD10CM/Codes/J00-J99/J40-J47/J45- AND R05, 2 - Wheeze. For children between the ages of 1-5 only Adults (>15 years of age) who are treated on an adult ward are not included. 1.4 How many emergency paediatric coded respiratory admissions (response to 1.3) were discharged from your dedicated paediatric ward(s) in XXXX. Number Number We don't have a dedicated paediatric ward Beds for admitted paediatric patients How many medical paediatric beds are there in your hospital which can be used for paediatric asthma patients? Number Whole numbers only. Whole numbers only. Whole numbers only. This is the number of paediatric osthma coded admissions discharged from your dedicated paediatric/respiratory wards in XXXX. Whole numbers only. Whole numbers only. This refers to beds designated to acute paediatric medicine at the time of the audit. This can include beds such as escalation ambulatory and day surgery beds as long as these are available to paediatric patients (including asthma) as and when required and the patients within them are under the care of the paediatric patients (including asthma) as and when required and the patients within them are under the care of the paediatric patients, please do not include them. Tick all that apply) Tick all that apply). Tick all that apply. Tick all that apply.			·
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 A mixed medical/surgical HDU A respiratory HDU 			
A respiratory HDU		A medical HDU	
		 A mixed medical/surgical HDU 	
Other, please specify			
		A respiratory HDU	

1.7	How many paediatric HDU beds are there in your	Select one only.
	hospital to which asthma patients can be admitted?	
	(Select one only)	
	• Zero	
	•	
	• Two	
	• Three	
	•	
	• Five	
	_	
	• Six	
	• Seven	
	•	
	•	
	• U Ten	
	• L Ten+	
1.8	How many general operational Paediatric Intensive Care	Select one only.
	Unit (PICU) beds, does your hospital have to which	
	asthma patients can be admitted? (Select one only)	In practice this will be PICU beds to which asthma patients are
	• Zero	admitted.
	•	
	• Two	
	• Three	
	• Four	
	• Five	
	• Six	
	• Seven	
	• 🔲 Eight	
	• Nine	
	• Ten	
	• Ten+	
Section 2	Staffing levels	
2.1	How many of the following staff posts are there in your	Up to 2 decimal places.
2.1	paediatric team?	op to 2 decimal places.
		Full time = 1 WTE (Whole Time Equivalent)
		Tull tille – 1 WTE (Whole Tille Equivalent)
	CT1/CT2ST3 and above	Please give the number of WTEs for each type of post (include
		both filled and vacant posts for each type of post (include
	Respiratory paediatric consultant	paediatric patients have access to adult or generic staff
	Paediatric consultant	routinely please include these in the WTE provided. For
	Associate specialist	example, if you have two respiratory physiologists who provide
	Staff grade	routine support to both adults and paediatric patients please
	Asthma nurse specialist	include the WTE for both of these in the box provided. If this is
	 Nurse consultant/other specialist nurse 	on a full time basis (paediatric asthma patients have access to
	 Respiratory/paediatric physiologist (lung function technician) 	them as and when required) enter 2 WTE. If they have time
	 Specialist respiratory/paediatric physiotherapist 	assigned specifically for treating paediatric patients, please only include the WTE assigned to this service. For example, if
	Other (please specify)	half of their time it dedicated to paediatric patients, enter 1 WTE, to represent 0.5 each.

2.2	How many unfilled paediatric team vacancies are there	Up to 2 decimal places.
	in your hospital at the time of completing this audit?	
	• FY1/FY2	Full time = 1 WTE (Whole Time Equivalent)
	• CT1/CT2	
	 ST3 and above 	Please give number of WTE vacant posts for types of post listed
	 Respiratory paediatric consultant 	at the time of completing this audit. If your paediatric patients
	 Paediatric consultant 	will have access to planned/unfilled posts for adult or generic
	Associate specialist	staff routinely please include these in the WTE provided. For
	Staff grade	example, there are two unfilled respiratory physiologists who
	Asthma nurse specialist	will provide routine support to both adults and paediatric
	Nurse consultant/other specialist nurse	patients as and when required on a full time basis please
	Respiratory/paediatric physiologist (lung function	include the 2 WTE for both of these in the box provided. If they
	technician)	will have time assigned specifically for treating paediatric
	 Specialist respiratory/paediatric physiotherapist 	patients, please only include the WTE to be assigned to this
	Specialist respiratory/paediatric physiotherapist	service. So, if only a quarter of their time will be dedicated to
	Other (places marify)	paediatric patients, enter 0.5 WTE to represent 0.25 each.
	 Other (please specify) 	
		The total WTE will be worked out by the audit team.
Section 3	Access to specialist staff and 7-day working	
	Access to specialist staff	
3.1	How many of the following staff members contribute to	Up to 2 decimal places.
	the acute paediatric intake rota?	
	Paediatric consultant	Full time = 1 WTE (Whole Time Equivalent)
	Respiratory paediatric consultant (separate or	
	on-call)	Please note the number of Whole Time Equivalents (WTEs).
	(Paediatric) Intensive care consultant	Acute paediatric consultant refers specifically to consultants in
	 Other specialist paediatric consultant 	acute paediatric medicine, not other physicians participating in
	Paediatric SpR/Specialty trainee and fellow	the on-call rota for medicine.
	. accuration opiny operatory training and renoting	
3.2	How frequently are paediatric patients on the	Select one for both weekdays and weekends.
	admissions ward reviewed by a senior decision maker	
	(SpR or above)?	
	(Select one only for both weekdays and weekends)	
	On weekdays?	
	Twice daily	
	o Daily	
	Other, please specify	
	• On weekends?	
	Twice daily	
	o Daily	
	Other, please specify	
3.3	Which asthma patients have access to a paediatric	Select one only.
3.3	respiratory nurse? (Select one only)	Science one only.
	• None	
	All asthma patients	
	Those under the care of a paediatric respiratory consultant	
	respiratory consultant	
	Other (please specify)	

3.4	Does your hospital have a designated, named clinical	Select one only.
	lead for asthma? (Select one only)	
	Paediatric lead only	NRAD
	Adult and paediatric lead	Every NHS hospital and general practice should have a
	No lead	designated, named clinical lead for asthma services,
	The read	responsible for formal training in the management of acute
		asthma.
3.4.1	If your hospital does have an asthma lead: Is this person	Select one only.
3.4.1	responsible for formal training in the management of	Select one only.
	acute paediatric asthma?	
	(Select one only)	
	Paediatric only	
	Adult and paediatric	
	• No	
	7 day working	
3.5	On which days does your hospital provide a PICU	Tick all that apply.
	outreach service for critically ill cases requiring PICU	
	management, and when is it available?	Outreach teams are usually PICU doctors or nurses who are
	(Tick all that apply)	called by ward staff to review deteriorating patients in order to
	• None	avoid or expedite transfer to the PICU.
	• Monday	
	• 🔲 Tuesday	This these days can vary, select the days this is most likely to or
	Wednesday	'usually' happen(s) on.
	• Thursday	
	• ☐ Friday	
	• Saturday	
	• Sunday	
	The outreach service runs overnight	
	Every night	
	Some nights	
3.6	On which days is there an on-call respiratory paediatric	Tick all that apply.
3.0	consultant available? (Tick all that apply)	тек ин тийт ирргу.
	• None	This is asking whether there is a formal on-call respiratory
		paediatric consultant available even if there isn't a respiratory
	•	1.
	• Tuesday	take as such.
	•	If the sea days are seen some select the description of the second like to
	• Thursday	If these days can vary, select the days this is most likely to or
	• Friday	'usually' happen(s) on.
	Saturday	
	• Sunday	
:		
3.6.1	When is the on-call respiratory paediatric consultant	Tick all that apply.
	available on these days? (Tick all that apply)	
	• 🔲 In-hours	It is understood that in-hours and out-of-hours differs from
	Out of hours	Trust to Trust and is dependent on the structure of contracts

		and handovers.
		For consistency within this audit, in-hours is considered 9am –
		5pm, Monday – Friday.
		Out of hours is considered 24/7 on-call availability (including
		telephone) of a respiratory paediatric consultant.
3.7	On which days does a senior decision maker from the	Tick all that apply.
	paediatric team (ST3 or above) undertake a ward round	,
	of new paediatric respiratory patients on the Paediatric	This question refers to whether formal ward rounds are
	Admissions Unit (PAU)? (Tick all that apply)	undertaken by these staff, as opposed to 'Board Rounds'.
	• None	
	Monday	If these days can vary, select the days this is most likely to or
	• Tuesday	'usually' happen(s) on.
	Wednesday	
	•	
	• Friday	
	• Saturday	
	• Sunday	
	Sullday	
3.8	On which days does a senior decision maker from the	Tick all that apply.
3.0	paediatric team (ST3 or above) undertake a ward round	rick all that apply.
	of new paediatric respiratory patients on the paediatric	This question refers to whether formal ward rounds are
	ward(s)? (Tick all that apply)	undertaken by these staff, as opposed to 'Board Rounds'.
	• None	undertaken by these stayy, as opposed to board nounds.
	• Monday	If these days can vary, select the days this is most likely to or
	Tuesday	'usually' happen(s) on.
		usuany happen(s) on.
	• Wednesday	
	• Thursday	
	•	
	• Saturday	
	• Sunday	
2.0		The state of the s
3.9	On which days is a respiratory nurse(s) available to	Tick all that apply.
	review asthma paediatric asthma patients?	This could be a respiratory nurse specialist, Asthma nurse or
	(Tick all that apply)	clinical nurse specialist with an interest in asthma.
	• None	chinical nurse specialist with an interest in astrima.
	•	If these days can your select the days this is most likely to an
	• Tuesday	If these days can vary, select the days this is most likely to or 'usually' happen(s) on.
	• Wednesday	usually happen(s) on.
	•	
	•	
	• Saturday	
	• Sunday	

Smoking cessation Select one only.			
hospital, and if so how many WTEs are provided to run the service? (Select one only) No service available			
the service? (Select one only) No service available delivered in the hospital either by your staff or a visiting smoking cessation programme, delivered in the hospital either by your staff or a visiting smoking cessation practitioner. 1			
 No service available Less than 0.5 0.5 1 2 3 4+ Other, please specify Are smoking-cessation pharmacotherapies prescribed to paediatric patients in your hospital? (Select one only) Yes, to patients over 12 years old No, to adults only 4.2.1 If yes, what type of pharmacotherapy is prescribed for paediatric patients (over 12 or 16 years old) (Tick all that apply) Nicotine replacement therapy Other, please specify Varenicline and Bupropion are not licensed for use in people under the age of 18.			
Less than 0.5 Less than 0.5 D.5 D.5 Less than 0.5 D.5 Less than 0.5 L			
1			
http://www.nhs.uk/Conditions/Smoking- 2			
http://www.nhs.uk/Conditions/Smoking- 2			
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Other, please specify 4.2 Are smoking-cessation pharmacotherapies prescribed to paediatric patients in your hospital? (Select one only)			
4.2 Are smoking-cessation pharmacotherapies prescribed to paediatric patients in your hospital? (Select one only)			
paediatric patients in your hospital? (Select one only) Yes, to patients over 12 years old Yes, to patients over 16 years old No, to adults only			
Yes, to patients over 12 years old Yes, to patients over 16 years old No, to adults only 4.2.1 If yes, what type of pharmacotherapy is prescribed for paediatric patients (over 12 or 16 years old) (Tick all that apply) Nicotine replacement therapy Other, please specify Tick all that apply. Varenicline and Bupropion are not licensed for use in people under the age of 18.			
Yes, to patients over 16 years old No, to adults only 4.2.1 If yes, what type of pharmacotherapy is prescribed for paediatric patients (over 12 or 16 years old) (Tick all that apply) Nicotine replacement therapy Other, please specify Tick all that apply. Varenicline and Bupropion are not licensed for use in people under the age of 18.			
No, to adults only 4.2.1 If yes, what type of pharmacotherapy is prescribed for paediatric patients (over 12 or 16 years old) (Tick all that apply) Nicotine replacement therapy Other, please specify Tick all that apply. Varenicline and Bupropion are not licensed for use in people under the age of 18.			
4.2.1 If yes, what type of pharmacotherapy is prescribed for paediatric patients (over 12 or 16 years old) (Tick all that apply) • Nicotine replacement therapy • Other, please specify Tick all that apply. Varenicline and Bupropion are not licensed for use in people under the age of 18.			
paediatric patients (over 12 or 16 years old) (Tick all that apply) Nicotine replacement therapy Other, please specify Varenicline and Bupropion are not licensed for use in people under the age of 18.			
 (Tick all that apply) Nicotine replacement therapy Other, please specify Varenicline and Bupropion are not licensed for use in people under the age of 18.			
 Nicotine replacement therapy Other, please specify under the age of 18.			
Other, please specify			
Prescriptions Prescriptions			
Prescriptions			
4.3 Does your hospital have paper prescriptions or Select one only.			
electronic prescribing? (Select one only)			
Paper prescriptions only			
Electronic prescribing partially implemented			
Electronic prescribing fully implemented			
throughout the organisation			
Managing respiratory failure – emergency oxygen therapy			
4.4 Does your hospital have a paediatric oxygen policy in Select one only.			
place? (Select one only) This can often be found on the trust intranet or via the trust			
Yes, paediatric specific clinical governance department.			
Yes, combined adult and paediatrics			
• No RCEM dataset			
 Not known O2 should be given on arrival to maintain sats 94-98% 			
O2 should be prescribed on arrival to maintain sats 94-98%			
BTS guideline			
Give supplementary oxygen to all hypoxaemic patients with			
acute severe asthma to maintain an SpO ₂ level of 94-98%.			
BTS/SIGN 2.6.1			
Questions around oxygen are included in the BTS paediatric			
clinical audit.			

4.5	Does the ward based paediatric medication chart/record	Select one only.	
	have a designated place in which to prescribe oxygen?		
	(Select one only)	RCEM dataset	
	• Yes	O₂ should be given on arrival to maintain sats 94-98%	
	• No	O ₂ should be prescribed on arrival to maintain sats 94-98%	
	-		
		BTS guideline	
		Give supplementary oxygen to all hypoxaemic patients with	
		acute severe asthma to maintain an SpO_2 level of 94-98%.	
		BTS/SIGN 2.6.1	
4.6	Does your hospital use a system of paediatric early	Select one only.	
	warning detection? (Select one only)		
	• Yes	This would include PEWS scoring as an alert to patient	
	• No	deterioration and need for clinical and/or Outreach Review.	
4.6.1	If yes, does your PEWS chart enable the recording of the	Tick all that apply.	
	following: (Tick all that apply)		
	• None		
	Target saturation		
	Actual saturation		
	 Amount of oxygen administered 		
4.6.2	If none, or no to any of the above, (PEWS and oxygen	Tick all that apply.	
	recording for PEWS), is there another monitoring chart		
	that allows the following to be recorded:	Validation required: Question and options available based on	
	(Tick all that apply)	what has been answered previously. Eg if PEWS chart captures	
	• None	target saturation, that is greyed out for this question.	
	Target saturation		
	Actual saturation		
	 Amount of oxygen administered 		
4.7	Does the hospital have an oxygen training programme in	Select one only.	
	place? (Select one only)		
	• Yes	Examples would include an online e-learning module, or formal	
	• No	teaching sessions around this topic	
	Not known		
	_		
Section 5	Integrated care		
5.1	Does your hospital have a paediatric specialist asthma	Select one only.	
	service? (Select one only)		
	• Yes	NRAD	
	• No	Patients with asthma must be referred to a specialist asthma	
	Not known	service if they have required more than two courses of systemic	
		corticosteroids, oral or injected, in the previous 12 months or	
		require management using British Thoracic Society (BTS)	
		stepwise treatment 4 or 5 to achieve goal.	

Section 6	Patient and carer engagement			
6.1	How often is there a formal survey seeking patient/carer	Select one only.		
	views on paediatric services? (Select one only)			
	 Continuous (every patient) 	Excludes the Friends and Family Test.		
	 More than 4 times a year 			
	3-4 times a year			
	• 1-2 times a year			
	Less than once a year			
	• Never			
6.2	Do you have a strategic group for paediatric services?	Select one only.		
	(Select one only)	,		
	• Tyes	This group is defined as consisting of senior clinical and		
	• No	management representatives, who meet regularly, set and		
	Not known	review targets, implement the stroke strategy and make plans		
	- Not known	for the future of the service.		
6.2.1	Does this group have a patient representative?	Select one only.		
	(Select one only)	,		
	• Tyes	A patient or carer who attends to provide a user perspective of		
	• No	care.		
	Not known			
6.3	Are patients able to access their own electronic records	Select one only.		
	at your hospital, as part of a collaborative self-	,		
	management scheme? (Select one only)	This question aims to establish if patients have access to their		
	• Tyes	own electronic records routinely, outside of the FOI request		
	• No	process, as part of a self-management scheme.		
	Not known			
6.4	Is your paediatric service linked with a specific patient	Select one only.		
	support or engagement group? (Select one only)	,		
	• Tyes			
	• No			
	Not known			
6.4.1	If yes, what is this group's name?	Free text.		
	7,			
	Free text			
Section 7	Transitional care			
7.1	Do your processes for transitioning young people from	Tick all that apply.		
	paediatric to adult services include ensuring that:			
	(Tick all that apply)	NICE guideline: Transition from children's to adults' services		
	 we do not have any formal transition 	for young people using health or social care services		
	arrangements	https://www.nice.org.uk/guidance/ng43/chapter/Implementa		
	 the young person has a full record of their 	tion-getting-started		
	condition			
	•	Ready Steady Go programme		
	 the young person has a transition plan and 	A suite of resources designed to deliver high-quality transition		
	that they have worked on this with both	for young people (YP) across all subspecialties.		
	paediatric and adult clinicians	https://www.nice.org.uk/sharedlearning/implementing-		
	 the young person has a named case worker 	transition-care-locally-and-nationally-using-the-ready-steady-		
	to assist in signposting for them and their family	go-programme		

BTS/SIGN guideline

TRANSITION TO ADULT-BASED HEALTHCARE

Transition to adult services is important for all adolescents with asthma, irrespective of the asthma severity. No studies on transition of adolescents with asthma to adult services were identified although there are many studies looking at transition of adolescents with chronic illness. Few studies compare different approaches and many recommendations come from consensus statements rather than randomised controlled trials.

It is important that the process of transition is co-ordinated and it is recommended that a healthcare professional be identified to oversee transition and either link with a counterpart in adult services or remain involved until the young person is settled within adult services.

In the initial period after transition to adult services in secondary care, adolescents are best seen by one consultant to build their confidence and encourage attendance.

PREPARATION FOR TRANSITION

Transition should be seen as a process and not just the event of transfer to adult services.

It should begin early, be planned, involve the young person, and be both age and developmentally appropriate (see Table 15).

British guideline on the management of asthma Table 15:

Recommendations for organising transition services (Table 15)

- Young people should be given the opportunity to be seen without their parents/ carers.
- Transition services must address the needs of parents/carers whose role in their child's life is evolving at this time.
- Transition services must be multidisciplinary and multiagency. Optimal care requires a co-operative working relationship between adult and paediatric services, particularly where the young person has complex needs with multiple specialty involvement.
- Co-ordination of transitional care is critical. There should be an identified coordinator who supports the young person until he or she is settled within the adult system.
- Young people should be encouraged to take part in transition/support programmes and/or put in contact with other appropriate youth support groups. The

		involvement of adult physicians prior to transfer supports
		attendance and adherence to treatment.
	•	Transition services must undergo continued evaluation.



Generic Questions

Pre-hospital and arrival

Acute observation and treatment

Review and discharge

All patients (Adults and children)

Case definition: See below for all.

- NHS number
- Date of birth
- Home postcode
- **Patient GP practice**
- Gender
- Exposure to second hand smoke at home

- Systemic steroids within 12 hours prior to arrival
- High dose B2 agonist within 4 hours prior to arrival
- Oxygen prior to arrival
- Arrival by ambulance
- Arrival at hospital (1)

- First recorded heart rate
- First recorded respiratory rate
- First recorded Sp02
- Supplementary oxygen
- Oxygen prescribed
- First administration of β2 agonists ##

Discharge from hospital



- If died as an inpatient
- Discharge bundle for admission
- Discharge bundle elements:
- \Rightarrow Inhaler technique
- ⇒ Assessment of medication
- $\Rightarrow PAAP$
- \Rightarrow Triggers
- \Rightarrow Follow up requests

Adults (16+)

Case definition: Admitted to hospital <u>adult</u> services for an acute exacerbation of asthma (primary diagnosis).

Smoking status

First recorded PEF

- Previous best PEF (if not recorded, predicted)

- First administration of system steroids ## (2)

- Discharge bundle elements:
- \Rightarrow Smoking cessation
- \Rightarrow (6 18 only) Smoking advice to parents/guardian
- Prescribed prednisolone (Mgs and days)

Children (6—18 years)

Case definition: Admitted to hospital paediatric services for an acute exacerbation of asthma (primary diagnosis).

Children (1—5 years)

Case definition: Admitted to hospital paediatric services for an acute exacerbation of asthma OR wheeze which is responsive to salbutamol (ICD-10 R06.2) (primary diagnosis).

 Systemic steroid administered

- Discharge bundle elements:
 - ⇒Smoking advice to parents/ quardian
- Prescribed prednisolone

Appendix 13: Removed questions: Secondary care audit - adult clinical dataset

Question Response Helpnotes		Helpnotes	Rationale
			REASON FOR REMOVAL
1. Generic questions			
Date and time of admission at your hospital	Date of admission to unit// (dd/mm/yyyy) Time of admission to unit/ (24hr clock 00:00)	Cannot be before date and time of arrival.	MOSCOW rating = M Consistency with COPD audit dataset. To ensure that analysis can take place against individual time pressured metrics and enables investigation into length of stay etc. REMOVE – Time of arrival adequate for analysis of time pressured metrics (steroids).
Route of referral to your hospital for this admission?	□ Ambulance □ Another hospital □ Advised by GP □ Advised by NHS 111 □ Self- referral □ Minor Injury Unit □ Walk-in Centre □ Other Please specify □ Not recorded	Select one only.	MOSCOW rating = C No NICE or BTS to support. Not in RCEM dataset. REMOVE – No evidence and not collected by other asthma audits.
Comorbidities recorded for this patient	☐ Diabetes ☐ Obesity ☐ Hypertension ☐ Coronary heart disease ☐ Stroke ☐ Heart failure ☐ Painful conditions (determined by repeat prescribed analgesics) ☐ Lung cancer ☐ COPD	Select all that apply	MOSCOW rating = C NICE Quality statement 5: Review Text re asthma reviews for both children and adults include reference to assessment of comorbidities. Statement = 'A structured review can improve clinical outcomes for people with asthma. Benefits associated with structured review may include reduced absence from school or work, reduced exacerbation rate, improved symptom control and reduced attendance in accident and emergency departments'

	 □ Bronchiectasis □ Osteoporosis □ Other Please specify □ None □ Not recorded 		Does having this information enable clinicians to assess and change medication and treatment in a more informed way? REMOVE – No evidence and not collected in other asthma audits and time consuming to complete. Could consider Charlson Index as a measure or derive from HES?
Did this patient have a recorded mental health diagnosis?	☐ Yes ☐ No	Select one only. The record of the patientspatient's mental health diagnosis should be as documented within the patient's notes and identified at the point of admission to hospital. Do and not include a diagnosis made during this hospital stay.	Included in dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 BTS Adult Asthma Audit Included in dataset. NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. REMOVE — Following investigation into other data sources, this question will be removed as it can be accessed via HES diagnostic fields (ICD-10 codes) to identify records with a mental health diagnosis.

Psychosocial or social factors recorded for this patient	 □ Depression □ Anxiety □ Psychiatric treatment (in the last 12 months?) □ Drug or alcohol abuse □ Deliberate self-harm □ Learning disability □ Social isolation/Lived alone □ Other Please specify □ None □ Not recorded 	Select all that apply	MOSCOW rating = S RCEM dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 BTS Adult Asthma Audit Included in dataset. NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. REMOVE – Following investigation into other data sources, this question will be removed as it can be accessed via HES diagnostic fields (ICD-10 codes) to identify records with a mental health diagnosis.
Did this patient have a recorded learning disability?	☐ Yes ☐ No	The record of learning disability diagnosis should be the point of admission to hospital and not include diagnosis during this hospital stay.	RCEM dataset Included in dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including

			the significance of concurrent psychological and mental health issues. REMOVE – Following investigation into other data sources, this question will be removed as it can be accessed via HES diagnostic fields (ICD-10 codes F70-F79) to identify records with recorded learning difficulties.
2. Asthma history			
Had asthma been diagnosed prior to this admission?	☐ Yes ☐ No ☐ Not recorded If no or Not recorded proceed to section 3.	Select one only	MOSCOW rating = S BTS Adult Asthma Audit Included in dataset REMOVE – Felt too time consuming and onerous for a continuous audit and not necessarily something that improves the care a patient receives.
Type of asthma at admission to your hospital	☐ Allergic asthma ☐ Late Onset asthma ☐ Occupational asthma ☐ Seasonal asthma ☐ Aspirin sensitive asthma ☐ Infective asthma ☐ Eosinophilic asthma ☐ Fungal asthma ☐ Other Please specify ☐ Not recorded	Select all that apply Allergic asthma: where there is specific allergic triggers for the patients asthma Late onset asthma: adult onset asthma with no previous history Occupational asthma: where substances found in the workplace cause asthma symptoms, attacks and exacerbations Seasonal asthma: asthma only experienced at certain times of year Aspirin sensitive asthma: medications such as non-steroidal anti-inflammatory drugs (NSAIDS) or aspirin cause asthma symptoms, attacks and exacerbations Infective asthma: asthma attacks triggered by bacterial infection Eosinophilic asthma: eosinophilia usually>0.3cell/µl often associated with nasal polyps and elevated FeNO Fungal asthma: IgE sensitisation to a	MOSCOW rating = S NICE quality statement 2: Diagnosing occupational asthma Adults with new onset asthma are assessed for occupational cause No BTS or NICE support. Enables us to track if change in diagnosis/type of asthma was made during admission. REMOVE - No evidence for inclusion and not asked in other audits. Having at discharge shows that review and triggers have been carried out appropriately and used to make an informed and accurate (re) diagnosis.

Did the patient have a Personalised	☐ Yes ☐ No ☐ Not recorded	fungal spore such as Aspergillus usually with high total IgE and eosinophilia Select one only	MOSCOW rating = S
Asthma Action Plan (PAAP) in place at the time of this admission?		A formal record and effective self-management document which contains information and advice on a specific patient's asthma. Components of the PAAP are: • Medication and inhaler techniques • Advice on medication and when this should be increased • Triggers and exacerbating factors • Advice on peak flow levels (best and worst) • Advice on how to recognise loss of control and when help should be sought • Actions to take if asthma deteriorates This document should be reviewed and updated at each annual asthma review and following each acute asthma episode.	NICE quality standard 3: Written personalised action plans People with asthma receive a written personalised action plan. BTS guideline All people with asthma (and/or their parents or carers) should be offered self-management education which should include a written personalised asthma action plan and be supported by regular professional review. 5.2.2 (A) NRAD All people with asthma should be provided with written guidance in the form of a personal asthma action plan (PAAP) which details their own triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency. REMOVE – Feedback received regarding difficulty accessing PAAP. To retain PAAP element at discharge, the focal point of the audit according to the HQIP NCAP spec.
If yes, what information was available within the existing PAAP?	 ☐ Current medication and adherence (go to Q2.4) ☐ Triggers and exacerbating factors (go to Q2.5) ☐ Advice on taking current medication ☐ Advice on increasing medication (action points) 	Select all that apply	MOSCOW rating = C NICE quality standard above. Not in BTS or RCEM datasets. REMOVE - No evidence for asking about content of PAAP. Onerous and time consuming for audit participants and not asked by other asthma audits.

	 □ Advice on calling for help □ Peak flow levels at which action should be taken □ A record of best personal peak expiratory flow (PEF) □ A record of best predicted peak expiratory flow (PEF) □ Not recorded 		
Asthma medication on admission	 □ Short acting reliever inhaler (SABA) □ Inhaled steroid inhalers (ICS) □ Inhaled steroid as a combined ICS/LABA □ Long acting bronchodilators (LABA) □ Xolair □ Methotrexate □ Spacer inhaler device □ Leukotriene Receptor Antagonist □ Home nebuliser □ Not recorded 	Select all that apply	MOSCOW rating = C BTS dataset asks about being on inhaled ICS before admission. Enables us to track if a change in medication was necessary as a result of attack/exacerbation and hospital admission. REMOVE - No evidence for collecting in this detail and also not asked by other asthma audits. BTS dataset asks about ICS before admission – we could include this alone.
Triggers and exacerbating factors identified for this patient prior to admission	 ☐ Food allergy ☐ Animal allergy ☐ Hay fever/allergic rhinitis ☐ Virus infection/URTIs ☐ Drugs ☐ Exercise ☐ Other Please specify ☐ Not recorded 	Select all that apply	MOSCOW rating = C Not in BTS or RCEM datasets, only that should form part of PAAP – not that it should be asked on admission. NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. Not in BTS or RCEM datasets, only that should form part of PAAP – not that it should be asked on admission. Enables us to track if a change in triggers was identified during hospital admission. REMOVE - No evidence specific to including these and not included in other asthma datasets.

3. Acute presentation and admission				
Level of consciousness on admission	GCS Scale (1-15) Not recorded	Glasgow Come Scale (GCS) is a neurological scale which gives a reliable and objective record of conscious state of a person for initial as well as subsequent assessment. A patient is assessed against the criteria of the scale and a score out of 15 is given. Elements of the scale include eyes, verbal and motor responses.	MOSCOW rating = S RCEM dataset Included in audit questions (considered vital sign – I think?) No BTS or NICE support. REMOVE- Little/no evidence for inclusion. Severity of asthma/attack can be obtained from Peak Flow and Sp02.	
Date and time PEF measurement taken	Date of PEF// (dd/mm/yyyy) Time of PEF/ (24hr clock 00:00) □ Not recorded	Enter a range of 1-15 only. Date and time should not be before date and time of arrival or admission.	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation Peak flow and Sp02 are recommended for doing this. BTS guideline Throughout the guideline peak flow is the recommended measurement system reference to for assessing asthma severity. RCEM dataset Included in audit questions BTS Adult Asthma Audit Included in dataset REMOVE - Record of measurement considered enough information to obtain necessary information.	
Date and time Sp02 was taken	Date of Sp02// (dd/mm/yyyy) Time of Sp02/_ (24hr clock 00:00) □ Not recorded	Date and time should not be before date and time of arrival or admission.	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation Peak flow and Sp02 are recommended for doing this. RCEM dataset	

			Included in audit questions BTS Adult Asthma Audit Included in dataset No BTS guideline or NICE support. REMOVE - Record of measurement considered enough
Was an objective measurement of asthma severity made on admission?	☐ Yes ☐ No ☐ Not recorded	Select one only	information to obtain necessary information. MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. Peak flow and Sp02 are recommended for doing this. BTS guideline Throughout the guideline peak flow is the recommended measurement system reference to for assessing asthma severity. RCEM dataset Included in audit questions. REMOVE – Sufficient to have Peak flow and Sp02
			recorded.
If yes, severity of asthma measured	☐ Mild ☐ Moderate ☐ Severe ☐ Life threatening	Select one only Children (older than 5) Moderate: Sp02 ≥92%, PEF ≥50% best or predicted, able to talk, heart rate ≤125/minute, respiratory rate ≤30/minute Severe: Sp02 <92%, PEF 33-50% best or predicted, too breathless to talk, heart rate >125/minute, respiratory rate >30/minute, use of accessory neck muscles Life threatening: Sp02 <92% plus any of: PEF 33% best or predicted, silent chest, poor respiratory effort too, agitation,	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. BTS guideline Diagrams on management of acute asthma in adults in hospital and ED include assessing asthma severity. REMOVE – Sufficient to have Peak flow and SpO2 recorded.

		confusion, cyanosis	
		Adults Moderate: Sp02 ≥92%, PEF >50-75% best or predicted, no features of acute severe asthma Severe: Sp02 ≥92%, PEF <50% best or predicted, heart rate ≥110/minute, respiratory rate ≥25/minute, cannot complete sentence in 1 breath Life threatening: Sp02 <92% plus any of: silent chest, cyanosis, poor respiratory effort, arrhythmia, hypotension, exhaustion, altered consciousness	
Patient admitted to an Intensive Care Unit (ICU)	☐ Yes ☐ No ☐ Not recorded	Select one only.	MOSCOW rating = C No BTS guideline or NICE support. REMOVE - No evidence to include and not used in
If yes, patient was ventilated in ICU	☐ Yes ☐ No ☐ Not recorded	Select one only.	other asthma audits. MOSCOW rating = C No BTS guideline or NICE support.
			REMOVE - No evidence to include and not used in other asthma audits.
4. Management in hospital			
Was the patient reviewed by a member of respiratory medical team within 24 hours of admission?	☐ Yes ☐ No ☐ Not recorded	Select one only.	MOSCOW rating = S/C Could compare this with COPD? REMOVE - No evidence/not applicable to asthma.
			COPD only.
If yes, date and team of review	Date of assessment// (dd/mm/yyyy) Time of assessment/ (24hr clock 00:00)	Date and time should not be before date and time of arrival or admission.	MOSCOW rating = S/C REMOVE – As above
l			

Assessment by a multidisciplinary	Date of assessment//		MOSCOW rating = S
severe asthma service offered	(dd/mm/yyyy)		NICE quality statement 11: Difficult asthma
Severe astrilla service offered	Time of assessment/ (24hr clock		People with difficult asthma are offered an assessment
	00:00)		by a multidisciplinary difficult asthma service.
	□ Not recorded		
			NRAD
	☐ Not offered		Patients with asthma must be referred to a specialist
			asthma service if they have required more than two
			courses of systemic corticosteroids, oral or injected, in
			the previous 12 months or require management using
			British Thoracic Society (BTS) stepwise treatment 4 or 5
			to achieve goal.
			BTS guideline
			Patients with difficult asthma should be systematically
			evaluated, including:
			- confirmation of the diagnosis of asthma, and
			- identification of the mechanism of persisting
			symptoms and assessment of adherence to therapy.
			This assessment should be facilitated through a
			dedicated multidisciplinary difficult asthma service, by a
			team experienced in the assessment and management
			of difficult asthma.
			REMOVE: Although would be interesting, not a priority
			for audit due to requirement of refined continuous
			dataset.
Best PEF recorded in 24 hours pre-	L/min	Range 1-100%	MOSCOW rating = S
discharge	% best		BTS guideline
	☐ Not recorded		Measure and record PEF 15–30 minutes after starting
			treatment, and thereafter according to the response.
			Measure and record PEF before and after nebulised or
			inhaled β2 agonist.
			BTS/SIGN 8.4
			BTS Adult Asthma Audit
			Included in dataset
			No NICE support.
			INO MICE Support.

Lowest PEF recorded in 24 hours pre- discharge	L/min % best □ Not recorded	Range 1-100%	REMOVE – The BTS guidelines are that PEF should be >75%predicted prior to discharge and using %best will not allow the audit to capture this data. To remove to ensure reduced dataset as only purpose is to ensure patient is discharged correctly. MOSCOW rating = M BTS guideline Measure and record PEF 15–30 minutes after starting treatment, and thereafter according to the response. Measure and record PEF before and after nebulised or inhaled β2 agonist. BTS/SIGN 8.4 BTS Adult Asthma Audit Included in dataset No NICE support.
			REMOVE – Does not seem to adhere to standard.
5. Review and discharge			
Type of asthma at discharge from your hospital	 □ Allergic asthma □ Late Onset asthma □ Occupational asthma □ Seasonal asthma □ Aspirin sensitive asthma □ Infective asthma □ Eosinophilic asthma □ Fungal asthma □ Other Please specify □ Not recorded 	Allergic asthma: where there is specific allergic triggers for the patients asthma Late onset asthma: adult onset asthma with no previous history Brittle asthma: Type 1: wide PEF variability (>40% diurnal variation for >50% of the time over a period of >150 days) despite intense therapy. Type 2: sudden severe attacks on the background of apparently well controlled asthma) Occupational asthma: where substances found in the workplace cause asthma symptoms, attacks and exacerbations Seasonal asthma: asthma only experiences at certain times of year	MOSCOW rating = S NICE quality statement 2: Diagnosing occupational asthma Adults with new onset asthma are assessed for occupational cause No BTS or NICE support. Enables us to track if change in diagnosis/type of asthma was made during admission. REMOVE - Although would be interesting, not a priority for audit due to requirement of refined continuous dataset.

Structured review by a member of the specialist respiratory team prior to discharge?	☐ Yes ☐ No ☐ Self-discharge	Aspirin sensitive asthma: medications such as non-steroidal anti-inflammatory drugs (NSAIDS) or aspirin cause asthma symptoms, attacks and exacerbations Infective asthma: asthma caused by an infection Eosinophilic asthma: Fungal asthma: Select one only. A team in which the clinical lead is a respiratory consultant (adult or paediatric) or specialist with an interest in respiratory disease (adult or paediatric) or a trained specialist nurse with expertise in managing asthma. The review should include: • An assessment of events leading up to the attack (including exposure to triggers, adherence and inhaler technique • review of written personalised asthma action plan (PAAP)	MOSCOW rating = M NICE quality statement 9: Specialist review People admitted to hospital with an acute exacerbation of asthma have a structured review by a member of the specialist respiratory team before discharge BTS Adult Asthma Audit Included in dataset REMOVE - Considered the same as a discharge bundle being carried out.
		review of regular treatment including considering whether this needs to be changed.	
If yes, date and time of review	Date of review// (dd/mm/yyyy) Time of review/ (24hr clock 00:00)	Date and time should not be before date and time of dates and times entered into sections, 1, 2, 3 or 4.	MOSCOW rating = M Enables investigation into how long reviews take to happen and if there is a delay between this and discharge of the patient (potential issues with social care etc)
			REMOVE - Considered the same as a discharge bundle being carried out.
Asthma medication at discharge from your hospital	 ☐ Short acting reliever inhaler (SABA) ☐ Inhaled corticosteroid inhalers (ICS) ☐ Inhaled corticosteroid as a combined 	Select all that apply.	MOSCOW rating = C Enables us to track if change in medication was required

	ICS/LABA Long acting bronchodilators (LABA) Xolair Methotrexate Spacer inhaler device Leukotriene Receptor Antagonist Home nebuliser Other Please specify Not recorded		following admission. REMOVE: Inclusion of 5.10 enough. Potentially only collected ICS on arrival instead of what was 2.4 so 5.10 would link to this?
Regular inhaled corticosteroid step-up treatment prescribed at discharge	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = S
from your hospital?			REMOVE - Considered part of medication review within discharge bundle.
Has a review of inhaled corticosteroid treatment before discharge for this patient been documented?	☐ Yes ☐ No	Select one only Evidence within the notes that the patients inhaled corticosteroid treatment was reviewed and/or changed prior to discharge for this admission.	BTS Adult Asthma Audit Included in dataset BTS guideline Clinicians should be aware that higher doses of inhaled corticosteroids may be needed in patients who are smokers or ex-smokers. Prevention of readmissions = decreases the risk of future exacerbations. REMOVE - Considered part of medication review within discharge bundle.
Triggers and exacerbating factors identified upon discharge for this admission	 ☐ Food allergy ☐ Animal allergy ☐ Hay fever/allergic rhinitis ☐ Virus infection/URTIs ☐ Drugs ☐ Exercise 	Select all that apply	MOSCOW rating = C NICE quality standard 2: Diagnosing occupational asthma Adults with new onset asthma are assessed for occupational cause

	☐ Other Please specify		Enables comparisons with triggers identified prior to attack/exacerbation and acute admission. REMOVE –Triggers asked within discharge bundle question.
If current smoker, was the patient prescribed smoking-cessation pharmacotherapy during the current admission? (e.g. NRT, varenicline)	☐ Not recorded ☐ Offered by declined ☐ No ☐ Yes	Select one only	MOSCOW rating = S Consistency with COPD dataset BTS guideline Clinicians should be aware that higher doses of inhaled corticosteroids may be needed in patients who are smokers or ex-smokers BTS/SIGN 6.2.4 REMOVE - Smoking cessation already covered within discharge bundle question.
6. Transition of care			within discharge bundle question.
Has the patient been recently referred to adult care?	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = S REMOVE – Deemed unnecessary for clinical dataset. Questions have been incorporated into organisational dataset.
If yes, is there	 □ evidence that the young person has a full record of their condition? □ evidence that the GP had the same record? □ evidence that the young person has a transition plan that he/she has worked on with Paed and adult clinicians □ evidence that young person has a named case worker to assist in signposting for them and their family. □ Not recorded 	Select all that apply	MOSCOW rating = S BTS guideline In the initial period after transition to adult services in secondary care, adolescents are best seen by one consultant to build their confidence and encourage attendance. (Page 122) REMOVE – Deemed unnecessary for clinical dataset. Questions have been incorporated into organisational dataset.
Was a referral letter drafted and sent to community service/specialist performing follow up?	☐ Yes ☐ No		REMOVE – Deemed unnecessary for clinical dataset. Questions have been incorporated into organisational dataset.

If yes, what elements were included in	☐ Medications (current and recent	
the referral letter?	changes)	REMOVE – Deemed unnecessary for clinical dataset.
	☐ Compliance (asthma control?)	Questions have been incorporated into organisational
	☐ Best PEF	dataset.
	prescriptions of steroids	
	☐ Previous admissions and ED visits	
	☐ Annual review information	
	☐ Smoking history	
	☐ Evidence of variability (how	
	diagnosis was made)	
	☐ Triggers and exacerbating factors	
	identified	
	☐ Associated atopy	

Appendix 14: Removed questions: Secondary care audit - paediatric clinical dataset

Question	Response	Helpnotes	Rationale
			REASON FOR REMOVAL
1. Generic questions			
Height			MOSCOW rating = M Ensures that the correct doses of medication are being given and prescribed for the children height and weight. REMOVE - Dosages not being asked and only minimal questions on medication. No evidence for inclusion and not asked in
			other asthma audits.
Weight			MOSCOW rating = M Ensures that the correct doses of medication are being given and prescribed for the children height and weight.
			REMOVE - Dosages not being asked and only minimal questions on medication. No evidence for inclusion and not asked in other asthma audits.
Date and time of admission at your hospital	Date of admission to unit// (dd/mm/yyyy) Time of admission to unit/ (24hr clock 00:00)	Cannot be before date and time of arrival.	MOSCOW rating = M Consistency with COPD audit dataset. To ensure that analysis can take place against individual time pressured metrics and enables investigation into length of stay etc.
			REMOVE - Time of arrival adequate for analysis of time pressured metrics (steroids).

Route of referral to your hospital for this admission?	☐ Ambulance ☐ Another hospital ☐ Advised by GP ☐ Advised by NHS 111 ☐ Self/Parental referral ☐ Minor Injury Unit ☐ Walk-in Centre ☐ Other Please specify ☐ Not recorded	Select one only.	MOSCOW rating = C No NICE or BTS to support. REMOVE - No evidence and not collected by other asthma audits.
Did this patient have a record of being under Child and Adolescent Mental Health Services (CAMHS)?	Yes No Not recorded	Select one only.	MOSCOW rating = S RCEM dataset Included in dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 BTS Adult Asthma Audit Included in dataset. NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. NOTE: Would using a screening tool such as the PHQ-2 which is a two item question which enquires about the degree to which

			an individual has experienced depressed mood and anhedonia over the past two weeks be more appropriate? Would obtaining this information be more time consuming/onerous than the current question? Is this type of tool something that is consistently and reliably used? REMOVE - Can capture via outcomes data (HES APC – Diagnosis).
Did this patient have a recorded learning disability?	☐ Yes ☐ No ☐ Not recorded		RCEM dataset Included in dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. REMOVE - Can capture via outcomes data (HES APC – Diagnosis).
Was this child known to social services?	⊠ Yes □ No □ Not recorded	Paediatric only	MOSCOW rating = C 14% of children in NRAD were known to social services. No NICE or BTS to support. Not in RCEM dataset.

Was this child subject to an existing child protection plan?	☐ Yes ☐ No ☐ Not recorded	Paediatric only	REMOVE - No evidence and not collected by other asthma audits. 14% of children in NRAD known to social services as only relatively small number. MOSCOW rating = C No NICE or BTS to support. Not in RCEM dataset.
Type of asthma at the time of this	☐ Allergic asthma	Select all that apply.	REMOVE – No evidence and not collected by other asthma audits. MOSCOW rating = S
admission	 □ Brittle asthma □ Occupational asthma □ Seasonal asthma □ Aspirin sensitive asthma □ Infective asthma 	,	This will enable us to track if a change in diagnosis was made during the admission. No NICE or BTS to support asking this. Not in RCEM dataset.
	☐ Eosinophilic asthma ☐ Fungal asthma ☐ Other Please specify ☐ Not recorded		REMOVE - No evidence and not asked in other audits. Kept in discharge as shows that review and triggers collected and used to make informed (re)diagnosis.
Had asthma been diagnosed prior to this admission?	☐ Yes ☐ No ☐ Not recorded If no or Not recorded proceed to section 3.	Select one only	MOSCOW rating = S BTS Paediatric Asthma Audit Included in dataset REMOVE - Although interesting not a priority for refined continuous dataset.
Is there a history of atopy?	☐ Yes ☐ No ☐ Not recorded	Tendency to present with allergic diseases such as allergic rhinitis (hay fever), food allergy and atopic dermatitis (eczema). Atopy is typically associated with heightened immune responses to common allergens, especially inhaled allergens and food allergens.	No NICE or BTS to support asking this. Not in RCEM dataset. REMOVE - A lot of mention of atopy throughout the guideline but no specific recommendations, guidance or standards.

Is there a family history of any of the	☐ Diabetes		No NICE or BTS to support asking this.
following?	☐ Obesity		Not in RCEM dataset.
	☐ Hypertension		
	☐ Coronary heart disease		REMOVE – No evidence and not collected in
	☐ Stroke		other asthma audits and time consuming to
	☐ Heart failure		complete.
	☐ Painful conditions (determined		
	by repeat prescribed analgesics)		
	☐ Lung cancer		
	□ COPD		
	☐ Asthma		
	☐ Bronchiectasis		
	☐ Osteoporosis		
	☐ Other		
	Please specify		
	□ None		
	☐ Not recorded		
Last known ACT score	(5-25)	A patient self-administered tool for	No NICE or BTS to support asking this.
		identifying those with poorly	Not in RCEM dataset.
		controlled asthma . The scores range from	
		5 (poor control of asthma) to 25	Again, a lot of mention of 'symptom scores'
		(complete control of asthma), with	throughout the guideline but no specific
		higher scores reflecting	recommendations.
		greater asthma control. An ACT score >19 indicates well-controlled asthma .	DEMONIE No suidence for inclusion and not
		indicates well-controlled astnma .	REMOVE- No evidence for inclusion and not used in any other asthma audits. Does not
			appear to be paediatric specific and cannot
			be used for under 12s? Also, seems to be
			GSK test and need to be careful with
			pharma endorsement.
Did the patient have a Personalised	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = S
Asthma Action Plan (PAAP) in place at		,	NICE quality standard 3: Written
the time of this admission?		A formal record and effective self-	personalised action plans
		management document which contains	People with asthma receive a written
		information and advice on a specific	personalised action plan.

		patient's asthma. Components of the PAAP are: • Medication and inhaler techniques • Advice on medication and when this should be increased • Triggers and exacerbating factors • Advice on peak flow levels (best and worst) • Advice on how to recognise loss of control and when help should be sought • Actions to take if asthma deteriorates This document should be reviewed and updated at each annual asthma review and following each acute asthma episode.	BTS guideline All people with asthma (and/or their parents or carers) should be offered self-management education which should include a written personalised asthma action plan and be supported by regular professional review. 5.2.2 (A) NRAD All people with asthma should be provided with written guidance in the form of a personal asthma action plan (PAAP) which details their own triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency. REMOVE - Although interesting not a priority for refined continuous dataset.
If yes, what information was available within the existing PAAP?	□ Current medication (go to Q2.4) □ Triggers and exacerbating factors (go to Q2.5) □ Advice on taking current medication □ Advice on increasing medication (action points) □ Advice on calling for help □ Peak flow levels at which action should be taken □ A record of best personal peak expiratory flow (PEF) □ A record of best predicted peak expiratory flow (PEF) □ Not recorded	Select all that apply	MOSCOW rating = C NICE quality standard above. Not in BTS or RCEM datasets. REMOVE - No evidence for asking for content of PAAP, onerous and time consuming for audit participants and not asked by other asthma audits.

Asthma medication on admission	☐ Short acting reliever inhaler (SABA) ☐ Inhaled corticosteroid inhalers (ICS) ☐ Inhaled corticosteroid as a combined ICS/LABA ☐ Long acting bronchodilators (LABA) ☐ Xolair ☐ Methotrexate ☐ Spacer inhaler device ☐ Leukotriene Receptor Antagonist ☐ Home nebuliser ☐ Not recorded	Select all that apply	MOSCOW rating = C BTS dataset asks about being on inhaled ICS before admission. Enables us to track if a change in medication was necessary as a result of attack/exacerbation and hospital admission. REMOVE - No evidence for collecting in this detail and not asked by other asthma audits. BTS dataset asks about ICS before admission – we could include this alone.
How many of the following have been prescribed in the last 12 months?	Courses of steroids Inhaled corticosteroid inhalers (ICS) β2 agonist		All asthma patients who have been prescribed more than 12 short-acting reliever inhalers in the previous 12 months should be invited for urgent review of their asthma control, with the aim of improving their asthma through education and change of treatment if required. REMOVE - Little or no evidence for inclusion. Not included in any other asthma audits. Very time consuming and onerous to collect.
Triggers and exacerbating factors identified for this patient prior to admission	☐ Food allergy ☐ Animal allergy ☐ Hay fever/allergic rhinitis ☐ Virus infection/URTIs ☐ Drugs ☐ Exercise ☐ Other Please specify	Select all that apply	MOSCOW rating = C NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. Not in BTS or RCEM datasets, only that

Number of A&E visits and hospital admissions in the last 12 months	□ Not recorded A&E visits hospital admissions		should form part of PAAP – not that it should be asked on admission. Enables us to track if a change in triggers was identified during hospital admission. REMOVE- No evidence specific to including these and not included in other asthma datasets. BTS Paediatric Asthma Audit Included in dataset
			REMOVE - As continuous audit, hospital admissions will be captured through previous submissions. Can A&E visits be captured via linkage instead?
Smoking status of patient, as document for the current admission	□ Not recorded □ Never smoked □ Ex-smoker □ Current smoker	Select one only	MOSCOW rating = S Consistency with COPD dataset BTS Paediatric Asthma Audit Included in dataset BTS guideline Parents with asthma should be advised about the dangers, to themselves and to their children with asthma, of smoking, and be offered appropriate support to stop smoking. NRAD A history of smoking and/or exposure to second-hand smoke should be documented in the medical records of all people with asthma. Current smokers should be offered referral to a smoking-cessation service. REMOVE - Not deemed appropriate to ask of paediatric patients, but smoking

			cessation section will remain in discharge bundle for 6-18 year olds. If prescribed, this would provide enough information to suggest a child smoked at time of admission.
If current smoker, was the patient prescribed smoking-cessation pharmacotherapy during the current admission? (e.g. NRT, varenicline)	☐ Not recorded ☐ Offered by declined ☐ No ☐ Yes	Select one only	MOSCOW rating = S Consistency with COPD dataset BTS guideline Clinicians should be aware that higher doses of inhaled corticosteroids may be needed in patients who are smokers or ex-smokers BTS/SIGN 6.2.4 REMOVE - Smoking cessation already covered within discharge bundle question.
3. Acute presentation and admission			question
Level of consciousness on admission	GCS Scale (1-15) Not recorded	Glasgow Come Scale (GCS) is a neurological scale which gives a reliable and objective record of conscious state of a person for initial as well as subsequent assessment. A patient is assessed against the criteria of the scale and a score out of 15 is given. Elements of the scale include eyes, verbal and motor responses. Enter a range of 1-15 only.	MOSCOW rating = S RCEM dataset Included in audit questions (considered vital sign – I think?) No BTS or NICE support. REMOVE - Little/no evidence for inclusion. Severity of asthma/attack can be obtained from Peak Flow and Sp02.
PEWS(R)/Asthma Severity Score	(Range?)		REMOVE - Little/no evidence for inclusion and not included in any other asthma audits. Equivalent (NEWS/MEWS) not being added to adult form.
Date and time PEF measurement taken (children 5 and over only)	Date of PEF// (dd/mm/yyyy) Time of PEF/_ (24hr clock 00:00)	Date and time should not be before date and time of arrival or admission.	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an

	□ Not recorded		objective measurement of severity at the time of presentation Peak flow and Sp02 are recommended for doing this. BTS guideline Throughout the guideline peak flow is the recommended measurement system reference to for assessing asthma severity. RCEM dataset Included in audit questions BTS Paediatric Asthma Audit Included in dataset REMOVE - Record of measurement considered enough to obtain necessary information.
Date and time Sp02 was taken	Date of Sp02// (dd/mm/yyyy) Time of Sp02/ _ (24hr clock 00:00) □ Not recorded	Date and time should not be before date and time of arrival or admission.	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation Peak flow and Sp02 are recommended for doing this. RCEM dataset Included in audit questions BTS Paediatric Asthma Audit Included in dataset REMOVE - Record of measurement considered enough information to obtain necessary information.
Was an objective measurement of asthma severity made on admission?	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an

			objective measurement of severity at the time of presentation. Peak flow and Sp02 are recommended for doing this. BTS guideline Throughout the guideline peak flow is the recommended measurement system reference to for assessing asthma severity. RCEM dataset Included in audit questions. REMOVE – Although interesting, sufficient to have Peak flow and Sp02 recorded.
If yes, severity of asthma measured	☐ Mild ☐ Moderate ☐ Severe ☐ Life threatening	Select one only Children 2-5 years Moderate: Sp02 ≥92%, able to talk, heart rate ≤140/minute, respiratory rate ≤40 minute Severe: Sp02 <92%, too breathless to talk, heart rate >140/minute, respiratory rate >40 minute, use of accessory neck muscles Life threatening: Sp02 <92%, silent chest, poor respiratory effort, agitation, confusion, cyanosis Children (older than 5) Moderate: Sp02 ≥92%, PEF ≥50% best or predicted, able to talk, heart rate ≤125/minute, respiratory rate ≤30/minute Severe: Sp02 <92%, PEF 33-50% best or predicted, too breathless to talk, heart rate >125/minute, respiratory rate >30/minute, use of accessory neck muscles	MOSCOW rating = M NICE quality statement 7: Assessing severity People with asthma who present with an exacerbation of their symptoms receive an objective measurement of severity at the time of presentation. BTS guideline Diagrams on management of acute asthma in children in hospital and ED include assessing asthma severity. REMOVE – Although interesting, sufficient to have Peak flow and SpO2 recorded.

		Life threatening: Sp02 <92% plus any of: PEF 33% best or predicted, silent chest, poor respiratory effort too, agitation, confusion, cyanosis	
High dose β2 agonist bronchodilator administered	Date of β2 agonist// (dd/mm/yyyy) Time of β2 agonist/ (24hr clock 00:00) □ Not recorded □ Not administered	Date and time should not be before date and time of arrival or admission.	MOSCOW rating = M BTS guidelines 2.6.1 (A) Use high-dose inhaled B2 agonists as first-line agents in patients with acute asthma and administered as early as possible. 2.6.2 (A) Inhaled B2 agonists are the first line treatment for acute asthma in children. RCEM dataset (within 10 minutes) High dose nebulised β2 agonists bronchodilator should be given within 10 minutes of arrival at ED BTS Paediatric Asthma Audit Included in dataset Enables timings of administration to be measured. REMOVE - Combine with nebuliser
Patient admitted to an Paediatric Intensive Care Unit (PICU)	☐ Yes ☐ No ☐ Not recorded	Select one only.	question. MOSCOW rating = C No BTS guideline or NICE support. BTS Paediatric Asthma Audit Included in dataset
If yes, patient was ventilated in PICU	☐ Yes ☐ No ☐ Not recorded	Select one only.	REMOVE- No evidence to include, and not used in other asthma audits. MOSCOW rating = C No BTS guideline or NICE support. BTS Paediatric Asthma Audit Included in dataset

			REMOVE - No evidence to include, and not used in other asthma audits.
4. Management in hospital			
Assessment by a multidisciplinary severe asthma service offered	Date of assessment		NICE quality statement 11: Difficult asthma People with difficult asthma are offered an assessment by a multidisciplinary difficult asthma service. NRAD Patients with asthma must be referred to a specialist asthma service if they have required more than two courses of systemic corticosteroids, oral or injected, in the previous 12 months or require management using British Thoracic Society (BTS) stepwise treatment 4 or 5 to achieve goal. BTS guideline Patients with difficult asthma should be systematically evaluated, including: - confirmation of the diagnosis of asthma, and - identification of the mechanism of persisting symptoms and assessment of adherence to therapy. This assessment should be facilitated through a dedicated multidisciplinary difficult asthma service, by a team experienced in the assessment and management of difficult asthma. REMOVE - Not applicable for paediatrics.
Best PEF recorded in 24 hours pre-	L/min		MOSCOW rating = S
discharge (children 5 and over only)	% best	Range 1-100%	BTS guideline
	☐ Not recorded	Shuaib – can you add more clinical detail please.	Measure and record PEF 15–30 minutes after starting treatment, and thereafter according to the response. Measure and

Lowest PEF recorded in 24 hours pre- discharge (children 5 and over only)	L/min % best □ Not recorded	Range 1-100% Shuaib – can you add more clinical detail please.	record PEF before and after nebulised or inhaled β2 agonist. BTS/SIGN 8.4 No NICE support. REMOVE – The BTS guidelines are that PEF should be >75%predicted prior to discharge and using %best will not allow the audit to capture this data. To remove to ensure reduced dataset as only purpose is to ensure patient is discharged correctly. MOSCOW rating = M BTS guideline Measure and record PEF 15–30 minutes after starting treatment, and thereafter according to the response. Measure and record PEF before and after nebulised or inhaled β2 agonist. BTS/SIGN 8.4 No NICE support.
			seem to adhere to standard.
5. Review and discharge			
Was a referral to a specialist made following this admission?	☐ Yes ☐ No ☐ Not recorded		REMOVE - Unsure how this question is different to the element included within the discharge bundle question.
If yes, what of the following criteria were fulfilled in order to make this referral?	□ Increased ICS dosage □ Decreased number of ICS inhalers □ Increased β2 agonist □ Increased number of courses of oral steroids □ Increased number of A&E visits		NRAD Patients with asthma must be referred to a specialist asthma service if they have required more than two courses of systemic corticosteroids, oral or injected, in the previous 12 months. The use of combination inhalers should be encouraged. Where long-acting beta agonist (LABA) bronchodilators are prescribed for

			people with asthma, they should be prescribed with an inhaled corticosteroid in a single combination inhaler. REMOVE - All patients are given appointment post discharge. No/little evidence for inclusion. Not asked by other
			asthma audits.
Type of asthma at discharge from your hospital	☐ Allergic asthma ☐ Brittle asthma ☐ Occupational asthma ☐ Seasonal asthma ☐ Aspirin sensitive asthma ☐ Infective asthma ☐ Eosinophilic asthma ☐ Fungal asthma ☐ Other Please specify ☐ Not recorded	Select all that apply Allergic asthma: where there is specific allergic triggers for the patients asthma Brittle asthma: Type 1: wide PEF variability (>40% diurnal variation for >50% of the time over a period of >150 days) despite intense therapy. Type 2: sudden severe attacks on the background of apparently well controlled asthma) Occupational asthma: where substances found in the workplace cause asthma symptoms, attacks and exacerbations Seasonal asthma: asthma only experiences at certain times of year Aspirin sensitive asthma: medications such as non-steroidal anti-inflammatory drugs (NSAIDS) or aspirin cause asthma symptoms, attacks and exacerbations Infective asthma: asthma caused by an infection Eosinophilic asthma: Fungal asthma:	MOSCOW rating = S NICE quality standard 2: Diagnosing occupational asthma Adults with new onset asthma are assessed for occupational cause No BTS or NICE support. Enables us to track if change in diagnosis/type of asthma was made during admission. REMOVE - Although would be interesting, not a priority for audit due to requirement of refined continuous dataset.
Structured review by a member of the	☐ Yes ☐ No ☐ Self-discharge	Select one only.	MOSCOW rating = M
specialist respiratory team prior to	Les Livo Lisellaige	A team in which the clinical lead is a	NICE quality statement 9: Specialist review
discharge?		respiratory consultant (adult or	People admitted to hospital with an acute
alsonarge:		paediatric) or specialist with an interest in	exacerbation of asthma have a structured
1	1	pacada io, or openiant with an intelest in	, chassing of activity have a structured

		respiratory disease (adult or paediatric) or a trained specialist nurse with expertise in managing asthma. The review should include: • An assessment of events leading up to the attack (including exposure to triggers, adherence and inhaler technique • review of written personalised asthma action plan (PAAP) review of regular treatment including considering whether this needs to be changed.	review by a member of the specialist respiratory team before discharge REMOVE - Considered the same as a discharge bundle being carried out.
If yes, date and time of review	Date of review// (dd/mm/yyyy) Time of review/ (24hr clock 00:00)	Date and time should not be before date and time of dates and times entered into sections, 1, 2, 3 or 4.	MOSCOW rating = M Enables investigation into how long reviews take to happen and if there is a delay between this and discharge of the patient (potential issues with social care etc) REMOVE - Considered the same as a discharge bundle being carried out.
Was the patient referred for assessment by a multidisciplinary severe asthma service?	Date of assessment/ (dd/mm/yyyy) Time of assessment/ (24hr clock 00:00) Not recorded Not offered Patient was already under the care of severe asthma service		MOSCOW rating = S NICE quality statement 11: Difficult asthma People with difficult asthma are offered an assessment by a multidisciplinary difficult asthma service. NRAD Patients with asthma must be referred to a specialist asthma service if they have required more than two courses of systemic corticosteroids, oral or injected, in the previous 12 months or require management using British Thoracic Society (BTS) stepwise treatment 4 or 5 to achieve goal. BTS guideline Patients with difficult asthma should be

Asthma medication at discharge from your hospital	☐ Short acting reliever inhaler (SABA) ☐ Inhaled steroid inhalers (ICS) ☐ Inhaled steroid as a combined ICS/LABA ☐ Long acting bronchodilators (LABA) ☐ Xolair ☐ Methotrexate ☐ Spacer inhaler device ☐ Leukotriene Receptor Antagonist ☐ Home nebuliser	Select all that apply.	and - identification of the mechanism of persisting symptoms and assessment of adherence to therapy. This assessment should be facilitated through a dedicated multidisciplinary difficult asthma service, by a team experienced in the assessment and management of difficult asthma. REMOVE - Not applicable for paediatrics. MOSCOW rating = C Enables us to track if change in medication was required following admission. REMOVE - Inclusion of 5.10 enough. Potentially only collected ICS on arrival instead of what were 2.4 so 5.10 would link to this?
	☐ Other Please specify ☐ Not recorded		
Regular inhaled corticosteroid step-up treatment prescribed at discharge from your hospital?	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = S ? REMOVE - Considered part of medication review within discharge bundle.

Triggers and exacerbating factor identified upon discharge for thi admission	0,	Select all that apply	MOSCOW rating = C NICE quality standard 2: Diagnosing occupational asthma Adults with new onset asthma are assessed for occupational cause Enables comparisons with triggers identified prior to attack/exacerbation and acute admission. REMOVE - Triggers included within
			discharge bundle question.
6. Transition of care Has the patient been recently referred to adult care?	☐ Yes ☐ No ☐ Not recorded	Select one only	MOSCOW rating = S REMOVE - Not necessary for continuous
			clinical audit.
If yes, is there	 □ evidence that the young person has a full record of their condition? □ evidence that the GP had the same record? □ evidence that the young person has a transition plan that he/she has worked on with Paed and adult clinicians □ evidence that young person has a named case worker to assist in signposting for them and their family. □ Not recorded 		MOSCOW rating = S BTS guideline In the initial period after transition to adult services in secondary care, adolescents are best seen by one consultant to build their confidence and encourage attendance. (Page 122) REMOVE - Not relevant for clinical dataset and to be included in organisational dataset.
Was a referral letter drafted and sent to community service/specialist performing follow up?			REMOVE: No referral required as mist will be followed up at the same hospital, but should assess these components in a GP referral letter.
			REMOVE - No referral required as most will be followed up at same hospital but should

		assess these components in GP referral letter.
If yes, what elements were included in the referral letter?	 ☐ Medications (current and recent changes) ☐ Compliance (asthma control?) ☐ Best PEF ☐ Exacerbation history i.e. dates and prescriptions of steroids ☐ Previous admissions and ED visits ☐ Annual review information ☐ Smoking history ☐ Evidence of variability (how diagnosis was made) ☐ Triggers and exacerbating factors identified ☐ Associated atopy 	REMOVE - No referral required as most will be followed up at same hospital but should assess these components in GP referral letter.

Appendix 15: Removed questions: Secondary care audit - adult org. dataset

Question	Category	Help note
Section 1: Admissions, staffing levels, general organisation	of care	
Do any of the consultant acute physicians have an	COPD and Asthma	
interest in respiratory medicine?		
• Yes		
• No		NOTE - Question will potentially be removed from the next audit due to lack of clarity
 Not know 		around what defines interest and what if only 1/20 etc.
Section 5: Integrating care across primary and secondary s	ectors	
Are COPD discharge bundles used for COPD patients	Asthma and	An example would include the BTS discharge bundle or similar.
discharged from your hospital following exacerbation,	COPD/COPD only	For asthma
and if so which elements of care do they cover (tick all		NICE quality statement 3: Written personalised action plans
that apply)?		NICE quality statement 4: Inhaler technique
• No		NICE quality statement 6: Assessing asthma control
 Smoking cessation advice 		NICE quality statement 10: Follow up on Primary Care
 Assessment of suitability for pulmonary 		BTS guideline
rehabilitation		Before initiating a new drug therapy practitioners should check adherence with existing
 Written COPD patient information 		therapies, inhaler technique and eliminate trigger factors. Inhaled corticosteroids are the
 Satisfactory use of inhalers demonstrated and understood 		recommended preventer drug for adults and children for achieving overall treatment goals. BTS/SIGN 2.4
		Prescribe inhalers only after patients have received training in the use of the device and have
 Confirmation of follow-up arrangements after discharge 		demonstrated satisfactory technique. BTS/SIGN 2.5 and 7.1
Other (please specify)		Adherence to long-term asthma treatment should be routinely and regularly addressed by all
()		healthcare professionals within the context of a comprehensive programme of accessible
OPTION 1: AMENDED QUESTION		proactive asthma care. BTS/SIGN 2.2
Are discharge bundles used for COPD and asthma		Before initiating a new drug therapy practitioners should check adherence with existing
patients discharged from your hospital following		therapies, inhaler technique and eliminate trigger factors. BTS/SIGN 2.4
exacerbation?		Prior to discharge, inpatients should receive written personalised asthma action plans, given
COPD patients		by healthcare professionals with expertise in providing asthma education. BTS/SIGN 2.2
·		Follow up should be arranged prior to discharge with the patient's general practitioner or

 Asthma patients No If yes, which elements of care do they cover (tick all that apply)? [Following list to appear if COPD patients selected] Smoking cessation advice Assessment of suitability for pulmonary rehabilitation Written COPD patient information Satisfactory use of inhalers demonstrated and understood Confirmation of follow-up arrangements after discharge Other (please specify) [Following list to appear if asthma patients selected] Review of triggers and exacerbating factors Medication assessment Review of inhaler technique Personal (Written) Asthma Action Plan (PAAP) Smoking cessation advice Follow up arrangements (with Primary care and specialist if necessary) 		asthma nurse within two working days and with a hospital specialist asthma nurse or respiratory physician at about one month after admission. BTS/SIGN 8.6.3 A hospital admission represents a window of opportunity to review self-management skills. No patient should leave hospital without a written personalised asthma action plan. (Page 43) NRAD – see recommendations on: Personal asthma action plans (PAAP) which details their own triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency. Follow-up arrangements. Factors that trigger or exacerbate asthma must be elicited routinely and documented. An assessment of recent asthma control should be undertaken at every asthma review. Non-adherence to preventer inhaled corticosteroids is associated with increased risk of poor asthma control and should be continually monitored. Patient self-management. Parents and children, and those who care for or teach them, should be educated about managing asthma Health professionals must be aware of the factors that increase the risk of asthma attacks and death. BTS Adult Asthma Audit: Included in dataset
specialist if necessary)		REMOVED - As questions around using care bundles are asked in both the COPD and asthma clinical datasets this information can be obtained from there.
OPTION 2: NEW/SEPARATE QUESTION Are asthma discharge bundles used for asthma patients discharged from your hospital following exacerbation and if so which elements of care do they cover (tick all that apply)? NO Review of triggers and exacerbating factors	Asthma only	

• 1	Medication assessment		
• F	Review of inhaler technique		
	Personal (Written) Asthma Action Plan		
	Smoking cessation advice		
	Follow up arrangements (with Primary care and		REMOVED - As questions around using care bundles are asked in both the COPD and
	specialist if necessary)		asthma clinical datasets this information can be obtained from there.
NEW QUE		Asthma only	BTS guideline for PR includes asthma patients (https://www.brit-
-	pulmonary rehabilitation service available to	,	thoracic.org.uk/document-library/clinical-information/pulmonary-rehabilitation/bts-
	atients discharged following exacerbations?		guideline-for-pulmonary-rehabilitation/) although routine referral is not recommended
·	ğ ğ		
			as it is for COPD and some information on the BLF website suggests that asthma patients can benefit from this.
			benefit from this.
			REMOVED - Not appropriate for asthma patients.
NEW QUE	STION	Asthma only	New VED Not appropriate for assuma patients.
-	5.5 is it available within X weeks of hospital	7.Stillia Silly	
discharge	-		REMOVED - Not appropriate for asthma patients.
	TION/QUESTION	Asthma and COPD	Based on BTS Adult Asthma Audit
	r hospital provide dedicated education for		NRAD
-	d asthma patients?		Parents and children, and those who care for or teach them, should be educated about
•	No		managing asthma. This should include emphasis on 'how', 'why' and 'when' they should use
•	Printed information		their asthma medications, recognising when asthma is not controlled and knowing when and
•	Face to face sessions		how to seek emergency advice.
•	Groups sessions		
•	Website information		Patient self-management should be encouraged to reflect their known triggers, eg increasing
•	Other (please specify)		medication before the start of the hay-fever season, avoiding non-steroidal anti-
	-		inflammatory drugs or by the early use of oral corticosteroids with viral- or allergic-induced
			exacerbations.
			BTS guideline
			Computerised decision support systems for patient use can be considered as an approach to
			supporting self-management.

	DEMOVED. Decreed many immediately desired and advantage of the
Is there a regular MDT meeting for patients with severe asthma, and if so which staff attend? (Tick all that apply) No Respiratory consultant Specialist trainee Other doctor (including GP) Asthma nurse specialist Nurse consultant/other specialist nurse Community based matron/nurse Respiratory physiotherapist Dietician Palliative care specialist Thoracic surgeon Psychologist Radiologist with interest in respiratory Respiratory pharmacist Other (please specify)	REMOVED - Deemed more important to include a patient and carer engagement section. All hospitals should have access to a severe asthma centre via a regional network and have the ability to discuss severe asthma cases. Please indicate if your centre is the regional NHSE commissioned centre for Severe Asthma, a mini-hub with ability to deliver some aspects of severe asthma care, or a networked centre engaging in MDTs with the Severe Asthma Centre, Some hospitals and/or integrated care services run MDT meetings to discuss complex cases in order to improve communication and care integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. Consistency with COPD audit. Included based on BTS Audit Asthma Audit: RCEM dataset Evidence to consideration given to psychosocial factors in adults prior to discharge. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 BTS Adult Asthma Audit Included in dataset. NRAD
	Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues.
	REMOVE: Only applicable to severe asthma services and as part of the requirements for a severe asthma service was not felt useful to include.
If yes to 5.5: How frequently does the meeting occur?	REMOVE: Only applicable to severe asthma services and as part of the requirements for a
(Select one only)	severe asthma service was not felt useful to include.

•	Weekly		
•	Fortnightly		
•	Monthly		
•	☐ Quarterly		
•	Other (please specify)		

Appendix 16: Removed questions: Secondary care audit - paediatric org. dataset

All items are compulsory.

Question	Help note
Section 1: Admissions, staffing levels, general organisation of care	
How many dedicated respiratory beds are there in your hospital?	Please enter the total number of respiratory beds. REMOVE - Not applicable for paediatrics as beds not specified as specifically for paediatrics.
How many designated Level 2 beds are there on your dedicated respiratory ward(s)?	Level 2 beds are those where the sickest cases are nursed outside of HDU/ICU, ie those requiring CPAP, NIV, step-down from HDU/ICU or requiring heightened observation and support:
 Zero One Two Three Four Five Five + 	Link to the Intensive Care Society Guidelines: https://www.ics.ac.uk/ICS/quidelines-and-standards.aspx
	REMOVE - Not applicable to paediatrics.
Is there an inpatient dietetic service in the hospital which paediatric asthma patients can be referred to, and if so how many WTEs are allocated to the paediatric medical wards to run the service?	If applicable, please note the number of WTE dieticians who staff the dietetic service.
 No service available Less than 0.5 0.5 1 2 3 	

 4+ Other Does your hospital run its own Pulmonary Rehabilitation	REMOVE - Not applicable to paediatrics.
Does your hospital run its own Pulmonary Rehabilitation	REMOVE - Not applicable to paediatrics.
	REMOVE - Not applicable to paediatrics.
	REMOVE - Not applicable to paediatrics.
Service?	
• Yes	
• No	REMOVE – Not applicable to paediatrics.
Section 2: Organisation of acute respiratory care	
Does your hospital operate a respiratory take separate from the rest of acute medicine?	Answer yes to this question if respiratory physicians have a dedicated on-call rota.
• Yes	
• No	REMOVE - Not applicable to paediatrics.
On which days does your hospital operate a system of specialty triage of cases to respiratory medicine? (tick all that apply)	I.e. are patients directed to particular wards dependent upon their diagnosis?
• None	
• Monday	
• Tuesday	
• Wednesday	
• Thursday	
• Friday	
• Saturday	
• Sunday	REMOVE – Not applicable to paediatrics.
On which days is there an on-call respiratory	This is asking about the availability of respiratory SpR opinion.

SpR/specialty trainee available? (tick all that apply)	
SpR/specialty trainee available? (tick all that apply) No on-call respiratory SpR/specialty trainee available Monday Tuesday Wednesday Thursday Friday Saturday Sunday On which days is a physiotherapist(s) available to review COPD patients where necessary? No physiotherapist available to review patients Monday Tuesday Wednesday Thursday Friday Saturday Saturday Sunday	REMOVE – Not applicable to paediatrics. Many hospitals use rotational physios (usually Band 5 or 6) to provide on-call respiratory services. Please answer according to the most appropriate for your hospital.
Which patients have access to a respiratory physiotherapist? None	REMOVE – Not applicable to paediatrics.
 All asthma patients Only those admitted under a respiratory physician Only those under the 'severe asthma service' Other (please specify) 	REMOVE – Not applicable to paediatrics.
On which days does a senior decision maker from the	This question refers to whether formal ward rounds are undertaken by these staff, as opposed to 'Board
paediatric respiratory team (SpR or above) undertake a	Rounds'.

ward round of new respiratory patients on the other	If your hospital only has a paediatric ward and paediatric patients would not be admitted to any wards by
ward(s)? (tick all that apply)	this please select 'Paediatric patients would only be admitted to a paediatric ward'.
• None	
•	This these days can vary, select the days this is most likely to or 'usually' happen(s) on.
•	
Wednesday	REMOVE: Now redundant due to changes made to question 2.3 (On which days does a senior decision maker
• Thursday	from the paediatric team (SpR or above) undertake a ward round of new paediatric respiratory patients on
• 🔲 Friday	the paediatric ward(s)?)
Saturday	
• Sunday	
Section 3: Managing respiratory failure – emergency oxyge	en therapy
Is there a named lead clinician responsible for the NIV service?	Please answer 'yes' if there is a specific named lead for the NIV service. This person may also lead the COPD or asthma service, and if that is the case, please also answer 'yes.'
• Yes	
• No	REMOVE - Different service and different context in children as no COPD. Consider removing unless discussing LTV service – optiflow is delivered by generalists on most wards. Not applicable to paediatrics.
In which setting(s) is non-invasive ventilation (NIV) given?	Please tick all sites where NIV is given
NIV not available	
 MAU/admissions ward 	
A&E/ED	
Respiratory ward	
General wardHDU	
• ICU	
Other	
	REMOVE - Different service and different context in children as no COPD. Consider removing unless discussing LTV service – optiflow is delivered by generalists on most wards. Not applicable to paediatrics.

Is there an NIV monitoring chart?	If so select all that it records.
 No NIV monitoring chart 	
 Initial blood gases 	
 Subsequent blood gases 	
 Time of application 	
Initial NIV pressures	
Subsequent NIV pressures	REMOVE – Different service and different context in children as no COPD. Consider removing unless discussing LTV service – optiflow is delivered by generalists on most wards. Not applicable to paediatrics.
Is there a training programme for staff providing NIV, if so what does it cover?	Examples would include an online e-learning module, or formal teaching sessions around this topic
 No training programme for staff providing NIV 	
Not known	
Indications for NIV	
 Initiation of NIV 	
 Nursing patients on NIV 	REMOVE – Not applicable to paediatrics
Has your hospital participated in the BTS NIV audit?	
Please select all the options applicable.	
 Not participated 	
• 2013	
• 2012	
• 2011	DENACY/E. Not applicable to prodictive
• 2010	REMOVE –Not applicable to paediatrics
Sub-question to 'Does your hospital have an oxygen training programme in place?	REMOVE –Not applicable to paediatric services
and the second s	
If yes, what does it cover? (Tick all that apply)	
 Prescription of emergency oxygen for doctors 	
Monitoring of emergency oxygen for nurses	
and other health professionals	
Not known	
Section 4: Integrating care across primary and secondary se	ectors
Are asthma discharge bundles used for paediatric asthma patients discharged from your hospital following	An example would include the BTS discharge bundle or similar. BTS discharge bundle should apply to all patients over the age of 2.

exacerbation and if so which elements of care do they cover (tick all that apply)?

- No
- Review of triggers and exacerbating factors
- Medication assessment
- Review of inhaler technique
- Personal (Written) Asthma Action Plan
- Smoking cessation advice
- Follow up arrangements (with Primary care and specialist if necessary)

NICE quality statements

NICE quality statement 3: Written personalised action plans

NICE quality statement 4: Inhaler technique

NICE quality statement 6: Assessing asthma control

NICE quality statement 10: Follow up on Primary Care

BTS guideline

Before initiating a new drug therapy practitioners should check adherence with existing therapies, inhaler technique and eliminate trigger factors. Inhaled corticosteroids are the recommended preventer drug for adults and children for achieving overall treatment goals. BTS/SIGN 2.4

Prescribe inhalers only after patients have received training in the use of the device and have demonstrated satisfactory technique. BTS/SIGN 2.5 and 7.1

Adherence to long-term asthma treatment should be routinely and regularly addressed by all healthcare professionals within the context of a comprehensive programme of accessible proactive asthma care. BTS/SIGN 2.2

Before initiating a new drug therapy practitioners should check adherence with existing therapies, inhaler technique and eliminate trigger factors. BTS/SIGN 2.4

Prior to discharge, inpatients should receive written personalised asthma action plans, given by healthcare professionals with expertise in providing asthma education. BTS/SIGN 2.2

Follow up should be arranged prior to discharge with the patient's general practitioner or asthma nurse within two working days and with a hospital specialist asthma nurse or respiratory physician at about one month after admission. BTS/SIGN 8.6.3

A hospital admission represents a window of opportunity to review self-management skills. No patient should leave hospital without a written personalised asthma action plan. (Page 43)

NRAD – see recommendations on:

Personal asthma action plans (PAAP) which details their own triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency.

Follow-up arrangements.

Factors that trigger or exacerbate asthma must be elicited routinely and documented.

An assessment of recent asthma control should be undertaken at every asthma review.

Non-adherence to preventer inhaled corticosteroids is associated with increased risk of poor asthma control and should be continually monitored.

Patient self-management .

	Parents and children, and those who care for or teach them, should be educated about managing asthma. Health professionals must be aware of the factors that increase the risk of asthma attacks and death.
	REMOVE - Clinical datasets asks this for every paediatric patient – will be captured there and therefore not needed here.
Does your hospital have access to a team that undertakes integrated care of patients with COPD (tick all that apply)?	Examples would include an early/assisted discharge service, supported discharge service, admissions avoidance service, hospital at home service and so on
 No – no access to team undertaking integrated care of patients with COPD Yes - hospital based team Yes – community based team Yes – single team that works across the primary/secondary care interface Not known 	We are asking these questions to understand the extent of integrated care that is now taking place across the NHS in England and Wales
	REMOVE – Not applicable to paediatrics.
How many days a week does it operate (tick all that apply)?	
 Monday Tuesday Wednesday Thursday Friday Saturday Sunday	
	REMOVE – Not applicable to paediatrics.
What services are provided and by whom (tick all that apply)?	I think we need to decide about the severe asthma service vs integrated care team above before I can say if COPD only or both.

- Not known
- Other (please specify)
- Out-reach Early/Supported discharge
 - Service not provided
 - Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- In-reach Early/Supported discharge,
 - Service not provided
 - Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- Admissions avoidance,
 - Service not provided
 - Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- Pulmonary rehabilitation,
 - Service not provided
 - Hospital based team
 - Community based team

- If yes, what is the name of the team?
- Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- Oxygen assessment service,
 - Service not provided
 - o Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- Medicines/chronic disease management,
 - o Service not provided
 - Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?
- Nebuliser service,
 - Service not provided
 - o Hospital based team
 - Community based team
 - If yes, what is the name of the team?
 - Single team that works across the primary/secondary care interface
 - If yes, what is the name of the team?

Constitute acception advise	
Smoking cessation advice,	
 Service not provided 	
 Hospital based team 	
 Community based team 	
If yes, what is the name of the	
team?	
 Single team that works across the 	
primary/secondary care interface	
If yes, what is the name of the team?	
if yes, what is the hame of the team:	

	T
	REMOVE – Not applicable to paediatrics
Is there a pulmonary rehabilitation service available to COPD patients discharged following exacerbation? • Yes • No • Not known	This refers to any pulmonary rehabilitation service that is available to patients discharged following an admission. REMOVE – Not applicable for paediatrics.
If yes to 5.4: is it available within 4 weeks of hospital discharge? • Yes • No • Not known	REMOVE – Not applicable for paediatrics.
Please select the option that provides the patient with their NIV, even if they receive their NIV treatment at home: No Not known Provided via our own hospital Provided via the regional centre only Provided via other provider	If the patient is provided with NIV by the hospital, regional centre or other provider, but actually receives the NIV treatment at home, then please select the option that has provided the patient with NIV. REMOVE – Not applicable to paediatrics. Consider something similar for LTV?
Is there specifically funded sessional time devoted to developing integrated respiratory services in your area, and if so what is the designation of the individual(s) responsible for developing these services? (tick all that apply)	

 No Integrated care respiratory physician Respiratory physician Respiratory nurse specialist Nurse consultant Respiratory physiotherapist 	
Other (please specify)	
	REMOVE – Not applicable to paediatrics.
Is there an integrated care pathway for managing COPD locally?	
• Yes	
• No	
Not known	
	REMOVE - Not applicable to paediatrics.
	NEMOTE Not applicable to paculation
Is there a regular MDT meeting for paediatric patients	Some hospitals run MDT meetings to discuss complex cases in order to improve communication and care
with asthma, and if so which staff attend?	Some hospitals run MDT meetings to discuss complex cases in order to improve communication and care integration. Please tick yes if your hospital runs such a meeting.
with asthma, and if so which staff attend?	
with asthma, and if so which staff attend? (Tick all that apply)	integration. Please tick yes if your hospital runs such a meeting.
with asthma, and if so which staff attend? (Tick all that apply) No	integration. Please tick yes if your hospital runs such a meeting.
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade.
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee Other doctor (including GP)	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee Other doctor (including GP) Asthma nurse specialist	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee Other doctor (including GP) Asthma nurse specialist Nurse consultant/other specialist nurse	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues.
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee Other doctor (including GP) Asthma nurse specialist Nurse consultant/other specialist nurse Community based matron/nurse	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death,
with asthma, and if so which staff attend? (Tick all that apply) No Respiratory paediatric consultant Paediatric consultant Specialist trainee Other doctor (including GP) Asthma nurse specialist Nurse consultant/other specialist nurse Community based matron/nurse Respiratory/paediatric physiotherapist	integration. Please tick yes if your hospital runs such a meeting. Other doctor includes FY, CT, associate specialist and staff grade. BTS guideline Healthcare professionals must be aware that patients with severe asthma and one or more adverse psychosocial factors are at risk of death. BTS/SIGN 8.1.3 NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues. REMOVE: Applicable to severe asthma services only and as part of their requirements not considered

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 Radiologist with interest in respiratory Respiratory pharmacist 		
Other (please specify)		
If yes to 4.2: How frequently does the meeting occur? (Select one only) Weekly Fortnightly Monthly Quarterly Other (please specify)	REMOVE: Applicable to severe asthma services only and as part of their requirements not considered useful to include.	
Section 5: Patient and carer engagement		
NEW SECTION/QUESTION	Based on BTS Adult Asthma Audit	
Does your hospital provide dedicated education for asthma patients? No Printed information Face to face sessions Groups sessions Website information Other (please specify)	Parents and children, and those who care for or teach them, should be educated about managing asthma. This should include emphasis on 'how', 'why' and 'when' they should use their asthma medications, recognising when asthma is not controlled and knowing when and how to seek emergency advice. Patient self-management should be encouraged to reflect their known triggers, eg increasing medication before the start of the hay-fever season, avoiding non-steroidal anti-inflammatory drugs or by the early use of oral corticosteroids with viral- or allergic-induced exacerbations. BTS guideline Computerised decision support systems for patient use can be considered as an approach to supporting self-management. REMOVE - Additional question in BTS audit but felt that falls under discharge bundle umbrella.	

Appendix 17: Case study 1 – Queen Elizabeth Hospital, Birmingham

Hospital details

Pilot hospital/site name:	Queen Elizabeth Hospital, Birmingham	
Service case study from :	Adult	
Region :	West Midlands	
Hospital type:	Acute	
Clinical and audit lead names:	Clinical lead: Col Duncan Wilson	
	Author: Sharon Rees	
	Audit contact: Stephen Rosser	
Number of asthma patients	397	
admitted a year (2016):		

Case study

How the AADP pilot process and data collection system were introduced and maintained at your hospital:

For example:

- How were patients identified
- How was the organisational information obtained
- Who collected the data
- Who entered the data
- Were cases entered prospectively (i.e. as the patients were admitted) or retrospectively, in batches)
- What communication pathways were set up between staff, departments, units etc.

All sixteen patients where identified via an automated alert system, which enters patients details and location into the Respiratory Support Teams email inbox. Alerts are generated by patients being admitted into medicine and Prednisolone >30mg OD along with nebulised bronchodilators being prescribed. This level of IT support reduces respiratory clinician time to patient's bed side, enhancing respiratory care.

Organisational data was collated by the trust informatics department having been requested by audit lead, Col Wilson.

AADP paper data sheets were completed by asthma clinical nurse specialists (CNS) in addition to completing electronic Asthma Care Bundle by the bedside as part of admission review. Some data sets being completed retrospectively, depending on service demands for that day. 10/16 data sets partially completed, pending discharge date, whilst the remainder sets completed retrospectively in full, following discharge.

Having fully completed 16 paper data sets the nursing team uploaded onto electronic audit spreadsheet. With audit lead submitting both organisational and clinical data once completed.

A pre-audit asthma service meeting was held. Prior to this meeting a CNS had highlight where information could be obtained from, possible difficulties and IT support required. Audit lead liaised with critical care colleagues asking them to inform the asthma team if any patients were admitted to level 3 care straight from the A + E department.

A laminated AADP patient information poster was displayed in our Difficult to Treat Asthma clinic.

What worked well:

For example:

- Patient identification
- Organisational data retrieval
- Acquisition of patient notes
- Data collection and entry
- Communication pathways
- Service processes
- Patient pathway

As a trust we are very fortunate to have IT systems which are labour saving by reducing manual information finding via paper notes. Although, some elements of the data set relies on effective documentation and uploading front door contacts on to electronic systems.

Having had an electronic Asthma Care Bundle in place for some time, a large amount of audit data is already being collected, reducing additional workloads.

Revision of the service Asthma Action Plan (AAP) coincided prior to audit launch. Although smoking status, occupational triggers and inhaler technique review date was included in the electronic Asthma Care Bundle, we did not display any evidence on the AAP that these elements had been discussed/reviewed. All point included now. As a service we were keen to ensure both the AAP and Asthma Care Bundle are fit for purpose, and as much data can be either 'pulled' or 'push' onto patients discharge letter, improving transition of care. As a trust and respiratory team we have already achieved this with our COPD service.

Prior to audit launch communications between Clinical, Managerial, Compliance and Informatics' teams was initiated to ensure IT systems/pathways would be in place for audit launch. Developing long-term strategies allowing future sustainability of data collection demonstrating evidence based care. Currently achieving Best Practice Tariffs within COPD care, all involved are keen for this to be achievable in Asthma care; if/when tariffs are introduced.

Audit experience demonstrated effective team working, especially between medical and nursing components.

What issues were encountered and how these were overcome (or will be in the future):

For example:

As above

As a team we are reliant on A + E administration staff to upload front door documentation, not all 16 patients had theirs available electronically. Which did resort in having to access paper notes.

Data point 1.3, 1.4 and 1.7 were not always documented. The need to refresh colleagues knowledge of the significance of these elements

is something as a team we need to take forward.

Since audit completion A + E prescribing is now carried out on an electronic system known as Patient Information Communication System (PICS). Prior to this paper scripts had

been used until the decision to admit was made, transferring prescriptions to PICS. Revision of the Asthma Bundle is required; hopefully once pilot audit data is published we can use this to escalate those discussions as a priority. Adapting and including additional points to AAP not only **Examples of learning, service** adaptation and improvements provides evidence of discussions had to other colleagues, it will made as a result of participating hopefully empower our patients to adjust their own plans, plus in the AADP pilot: request health professionals review their inhaler technique for intense, if not observed recently, in turn giving our patients insight into what they should expect from an asthma review in For example: acute or stable state. Asthma care Further education to colleagues on not just acute presentations pathway/treatment of asthma but to think about, once discharged, are our patients Service processes linked into the right service for follow up? And to consider the Communication pathways impact of psychosocial elements. Patient identification and/or audit data acquisition Data collection and entry Quality Improvement project/s initiated Other recommendations on how Having IT systems to support data collection is a huge other respiratory/paediatric advantage. How viable this is for other trusts to develop and services can incorporate the new sustain obviously comes with a cost attached. Although if asthma audit into their day to targets are attached to evidence based care these costs need to day working life): be factored in, and in the long term hopefully become cost neutral. Effective team working with early communication and planning is key to developing services long term ensuring consistent care and information gathering.

Appendix 18: Case study 2 – Royal Derby Hospital

Hospital details

Pilot hospital/site name:	Royal Derby Hospital	
Service case study from :	Paediatric	
Region:	East Midlands	
Hospital type:	District general	
Clinical and audit lead names:	Clinical leads: Dr Simon Chitnis, Dr Nigel Ruggins, Dr Helen Riches	
	Audit contacts: Sarah Stevenson, June Bettson- Green, Stuart	
	Benney	
Number of asthma patients	Information not available.	
admitted a year (2016):		

Case study

How the AADP pilot process and
data collection system were
introduced and maintained at
your hospital:

For example:

- How were patients identified
- How was the organisational information obtained
- Who collected the data
- Who entered the data
- Were cases entered prospectively (i.e. as the patients were admitted) or retrospectively, in batches)
- What communication pathways were set up between staff, departments, units etc.

Patients were identified by keeping an eye on the patients admitted with wheeze/asthma by the juniors on the ward, or asking other doctors to alert the juniors involved in the audit if they were not on the ward when a patient with wheeze or asthma was admitted. Info was obtained from patient notes, usually after completed discharge. Ideally patient notes were collected prior to being sent for coding, but often had to be found after coding.

Junior doctors collected the data and entered it. Info was entered retrospectively.

Due to only a small number of patients required and a small team in the hospital, patients were either identified by those doing the audit when on the ward, or asking colleagues to let them know if a suitable patient had been admitted. If more patients were needed, a suitable option would be to alert all staff of the need for wheeze or asthma patients and to have a central document accessible by all staff to upload patient details. Then once patient details were uploaded, those involved in the audit would be able to collect data at a convenient time.

What worked well:

For example:

- Patient identification
- Organisational data retrieval
- Acquisition of patient notes
- Data collection and entry
- Communication pathways
- Service processes
- Patient pathway

Patient identification was not very efficient, however could be streamlined so that suitable patients were not missed.

Data retrieval was simple. Notes contained all relevant information and information was easily found.

Patient notes acquisition was the most time-consuming part of

Patient notes acquisition was the most time-consuming part of the process. Junior doctors were left to retrieve notes if they were not caught before being sent to coding. This involved several phone calls and trips to the coding department to retrieve them which would have been problematic had the wards been busier.

Because of the need for patient data to be in as soon as the 4 weeks ended, it was not possible to keep track of patient details, and then source all notes for the patients in one session and then input the data. The simplest way was to keep on top of it as you went along, however with busier wards and if one was not able to find the notes before they were sent for coding then this would have been tricky.

Perhaps a week window at the end of the 4 week period would allow all notes to be sourced and entered in one afternoon to ensure no patients were missed.

What issues were encountered and how these were overcome (or will be in the future):

The main issues were tracking patient notes and getting to them before they were sent for coding, as once they were sent, it became more time consuming to gather the information afterwards.

For example:

Identifying patients was not particularly efficient as mentioned but can be improved on for next time.

As above

Examples of learning, service adaptation and improvements made as a result of participating in the AADP pilot:

For example:

- Asthma care pathway/treatment
- Service processes
- Communication pathways
- Patient identification and/or audit data acquisition
- Data collection and entry
- Quality Improvement project/s initiated

Need for more use of BTS bundles.

Need for improved documentation in all aspects of patient care. Setting up the audit again for more patients would be much easier next time having learnt how to do it for this pilot.

If a departmental audit was being carried out in the future, it would be easier to fit the audit into daily working life having carried out the pilot.

Achieved a better understanding of patient notes acquisition.

Other recommendations on how other respiratory/paediatric services can incorporate the new asthma audit into their day to day working life):

care.

FY1's generally discharge most of the patients/organise discharges. If they filled in the data sheet with patient information after discharging a patient, no patients would be missed and it would save a lot of time as they would already know most of the information and be able to find it easily. If the person discharging the patient filled in the information, then that would save lots of time. If it was also only one patient at a time, then the workload would seem reduced.

Plus lessons learnt from doing the audit (eg. Noticing that bundles were not being used regularly) may improve patient

Appendix 19: Case study 3 – Royal United Hospital, Bath

Hospital details

Pilot hospital/site name:	Royal United Hospitals Bath NHS Foundation Trust (Royal United	
	Hospital)	
Service case study from :	Adult	
Region:	South West	
Hospital type:	District general	
AADP clinical and audit lead	Clinical lead: Dr Sharon Sturney	
names:	Audit contact: Mr Robert Eliot	
Number of asthma patients	234	
admitted a year:		

Case study

How the AADP pilot process and data collection system were introduced and maintained at your hospital:

For example:

- How were patients identified
- How was the organisational information obtained
- Who collected the data
- Who entered the data
- Were cases entered prospectively (i.e. as the patients were admitted) or retrospectively, in batches)
- What communication pathways were set up between staff, departments, units etc.

The Royal United Hospital Bath is a District General Hospital with a relatively low number of acute asthma admissions. All adults admitted with an acute exacerbation of asthma, whether referred by their GP or from the Emergency Department (ED), should be seen on the Medical Assessment Unit (MAU) and then either discharged or moved to an inpatient ward. Fewer than 20% of patients with a coded emergency admission due to asthma are cared for on the respiratory ward, therefore it was important to devise several mechanisms by which we were alerted when they presented to hospital.

- A GP specialist trainee is currently responsible for reviewing the electronic admission record each weekday morning to locate patients admitted with COPD exacerbations for the on-going national COPD audit, and her role was expanded to include highlighting patients with acute exacerbations of asthma.
- Respiratory consultants provide a weekday morning review of a number of patients with respiratory conditions on MAU, thereby assessing a number of patients with asthma.
- 3) Once asthmatic patients were admitted to a medical ward, we relied on our colleagues to contact the RNSs to ask for them to be reviewed.

We prepared for the introduction of the Asthma Audit Development Project (AADP) by updating our acute asthma discharge care bundle and temporary asthma action plan, then used several different modalities of communication (face-to-face, email, electronic staff bulletin, presentation at the weekly respiratory department teaching meeting) to make staff on the

medical wards aware that the audit was taking place. Ward teams were asked to contact our Respiratory Nurse Specialists (RNSs) with any appropriate patients so that they could review them on the ward prior to discharge, perform inhaler checks as necessary, fill in the discharge bundle and collect the necessary data for the audit. The asthma discharge bundle was relatively recently introduced within the hospital, so we used the audit as a driver to increase its profile and improve patient care.

Once datasets were completed on paper, information was inputted onto the spreadsheet by the asthma lead consultant. Some organisational information was obtained by the asthma lead consultant from the business intelligence unit/coding department.

What worked well:

For example:

- Patient identification
- Organisational data retrieval
- Acquisition of patient notes
- Data collection and entry
- Communication pathways
- Service processes
- Patient pathway

Organisational data was relatively easy to obtain, although we needed to be careful not to include elective admissions for omalizumab treatment or acute admissions for patients with asthma where an exacerbation was not the main diagnosis.

Data collection worked well for patients who spent more than 24 hours in the hospital, with the RNS team retrieving important information when completing discharge bundles. Once the data was collected in paper format it was then inputted onto the spreadsheet held on a departmental drive. This was efficient for the purposes of a four week audit, however in future this task could be performed by another member of the respiratory team; in our case an RNS, admin support worker or COPD audit nurse in a combined role. From next month the number of ward computers will increase, which would enable the direct inputting of data onto the spreadsheet at the time the discharge bundle is completed, saving time.

What issues were encountered and how these were overcome (or will be in the future):

For example:

As above

We encountered some difficulty in identifying suitable patients for the audit prior to their discharge, and this also had implications for patient review by a member of the respiratory team and completion of a discharge bundle. Firstly, a GP specialist trainee reviewed the electronic admission record each weekday morning to locate patients admitted with asthma exacerbations, but often the diagnosis was inaccurate and this caused extra work in looking through patient notes. We are examining ways in which the accuracy of clinical information, including working diagnoses, on that database can be improved to make the process more efficient.

Secondly, once asthmatic patients were admitted to a medical ward, we relied on our colleagues to contact the RNSs to ask for

them to be reviewed, which was a slight change to previous practice and meant that at times they heard about patients when it was difficult for them to be reviewed before discharge. Prior to the AADP launch, the RNS team had been educating ward nurses on MAU and the respiratory ward about the discharge bundle and this information was disseminated further to include all of the medical wards at morning safety briefings and via the electronic staff bulletin. Ward managers, medical nurse practitioners, and doctors working in acute medicine were contacted via personal email. This education will need to continue.

During the AADP, several patients were admitted over a weekend when specialist respiratory staff were not present to identify patients or collect data, and others were discharged from the post-take ward round meaning we were unaware of their admission until after they had gone home. In those cases, our secretaries were able to obtain the notes so that data could be submitted for the audit, but because the RNSs had not seen the patients no discharge bundle was completed and, therefore, patient care was not optimal. The roll out of an on-going national audit means that we need to address this issue, and one solution is to provide a lot of the information that our RNSs deliver face-to-face in an information leaflet which we are currently developing. This could be done in conjunction with educating ward nurses to complete the discharge bundle. This would be particularly useful on the respiratory ward, MAU and medical short stay ward where the majority of patients are discharged.

Next month, a new ED electronic patient record will be launched and we will investigate whether this could be used to flag asthma admissions automatically.

The audit data for patients still in the hospital was collected at same time as the discharge bundle was completed. One problem we encountered was that information was stored in three folders located in different places and within several documents: medical clerking/notes, ED nursing notes, observation chart, peak flow chart, drug chart, discharge bundle paperwork and electronic patient record. The introduction of a new ED electronic patient record and e-prescribing next month, and e-observation charts later in the year, as well as scanning the discharge bundle into the electronic patient record, should lead to increased efficiency in data collection when the on-going

Examples of learning, service adaptation and improvements made as a result of participating in the AADP pilot:

For example:

- Asthma care pathway/treatment
- Service processes
- Communication pathways
- Patient identification and/or audit data acquisition
- Data collection and entry
- Quality Improvement project/s initiated

Other recommendations on how other respiratory/paediatric services can incorporate the new asthma audit into their day to day working life:

audit commences. It may mean that a non-clinical member of the team can collect and input data remotely, particularly in cases where the RNSs did not meet the patient.

Our involvement in the AADP has had a positive impact on the awareness of acute asthma within our department and the wider medical division and has enabled us to make improvements in patient management, most notably the use of the discharge care bundle. Education and awareness of staff are key to identifying the correct patients, as well as utilising information technology (IT) to make the process more efficient. In our case, planned changes to IT should lead to improved incorporation of the national on-going audit process into daily clinical practice when it is launched in the New Year. The audit process will increase our workload and this will need to be shared by several members of the respiratory clinical and administrative team so that absences etc. do not affect our ability to fulfil requirements. Some of this work may be taken on by members of the team responsible for the national COPD audit. Our focus now needs to be on patients who present and are then discharged out-of-hours and how we can improve their care. Initially we will try to do this by producing an information leaflet, including signposts to useful online material.

One of the most important things when incorporating the audit into day to day working was to involve several members of the team, as this will enable the data to be collected when individuals are away from work for whatever reason. Depending on your set-up for the ongoing national COPD audit, it may be possible to use the same staff members to identify patients, collect and input the data.

Appendix 20: Case study 4 – Watford General Hospital

Hospital details

Pilot hospital/site name:	West Hertfordshire NHS Trust Watford General Hospital
Service case study from:	Adult
Region:	East of England
Hospital type:	District General Hospital
Clinical and audit lead names:	Clinical lead: Dr Kay Roy
	Audit contact: Henah Tabassam, Nariman Feituri
Number of asthma patients admitted a	345
year (2016):	

Case study

How the AADP pilot process and data collection system were introduced and maintained at your hospital:

For example:

- How were patients
- identified
- How was the organisational
- information obtained
- Who collected the data
- Who entered the data
- Were cases entered
- prospectively (i.e. as the
- patients were admitted) or
- retrospectively, in batches)
- What communication
- pathways were set up
- between staff,
- departments, units etc.

An audit proforma was created which also served as an admissions tool for acute asthma in adults. This simple single page document covered all parameters required for the audit as well as others required by the acute admitting teams. It also covered an algorithm for management of asthma as per BTS guidelines as well as the asthma discharge bundle with all the essential components.

Patients admitted with acute asthma were identified by the respiratory team (consultant covering acute respiratory admissions) and the community respiratory nurses who provide a daily in-reach service to the hospital and review patients at all stages of their pathway from ED to the acute admissions unit for medicine, the specialised respiratory ward as well as other hospital wards. The community nurses were also largely involved in collecting the data and completing the proforma before data was entered by our audit department although overseen and checked by the secondary care respiratory consultant. Data was collected and entered prospectively. The organisational information was obtained from the coding department.

Clear communication pathways have been set up and established between the acute services (AAU and ED), who can use the asthma proforma as an admissions tool before referring to the specialist respiratory services compromising of both the hospital and community teams. This interface between 3 teams and staff members ensures a streamlined patient journey from acute admission to discharge and very importantly continued care and follow up thereafter once back in the community.

What worked well:

For example:

- Patient identification
- Organisational data retrieval
- Acquisition of patient notes
- Data collection and entry
- Communication pathways
- Service processes
- Patient pathway

This audit demonstrated that clear working relationships between departments and staff involved in different aspects of patient journey as described above can be effectively achieved through good communication.

We developed a proforma led pathway for acute adult asthma which not only incorporated all the essential elements of the audit but also assessment metrics which the ED consultant lead perceived as important to the initial evaluation of the patient. The pathway was hence conceived with input from the acute services which helped maintain engagement with the audit process and ensured early referral to the respiratory services and allows improved integration of all services as well.

Data was always collected prospectively by the same respiratory nurse from start to end of the patients admission and at the same time for all 100% of patients, the discharge bundle was provided with in depth counselling around careful management of their asthma as well as follow up arrangements. A copy of the proforma was kept in the patient notes in both secondary care and entered onto the community respiratory service database.

What issues were encountered and how these were overcome (or will be in the future):

For example: As above Continuing education is an essential part of the asthma service and through talks and discussions with the acute services we aim to ensure a smooth patient journey is achieved. Due to staff turnover amongst doctors especially in all departments we must ensure that a regular rolling teaching programme is embedded in the service programme to familiarise all involved in the care of adult asthma with the pathway, management and referral mechanisms to both the respiratory team in secondary care as well as the community respiratory team. There are contact numbers (fax and telephone) at the end of the asthma proforma to ensure that anyone discharged outside normal working hours by ED or AAU has their discharge summary faxed straight away to ensure the patient is then followed up by phone call or home visit the next working day by the community respiratory team.

Examples of learning, service adaptation and improvements made

This audit has allowed us to look closely at the asthma pathway and its effective uptake and implementation in

as a result of participating in the AADP pilot:

For example:

- Asthma care pathway/ treatment
- Service processes
- Communication pathways
- Patient identification and/or audit data acquisition
- Data collection and entry
- Quality Improvement project/s initiated

all hospital departments and settings through better communication links and education and training. There is now improved scope to refer to the specialist respiratory services. We have worked closely with ED to develop this pathway and ensure services are more integrated and also have improved our management of patients who have not been seen by a respiratory specialist but discharged by acute medicine. The discharge summary can be faxed to the community respiratory team as described above and we have also now ensured through close working with the pharmacy team in the hospital, that a steroid inhaler can be issued and dispensed from ED to any adult asthmatic (Step 1 BTS not on any steroid inhaler previously) discharged without the patient needing to pick the prescription up from pharmacy. As the audit was completed prospectively and continuity of care maintained by the same community nurse following up the patient, we ensured that 100% patients had a complete discharge bundle assessed and delivered as well as follow up organised and this was all stated on the proforma in detail.

We work closely with our audit department who provide regular staffing to maintain data entry from the completed proformas for all our ongoing respiratory audits.

Aside from the quality improvement projects initiated above, we will be looking at re auditing in 6 months to review in more detail the impact that the service improvements above have had on asthma care in hospital and admission and readmission rates. In particular we will also be looking at smart health initiatives for personalised and self-directed asthma care and how this influences upon adherence and compliance with personal asthma action plans as well as exacerbation rates.

Other recommendations on how other respiratory/ paediatric services can incorporate the new asthma audit into their day to day working life):

Our service described above is unique in some ways but essentially delivers an integrated respiratory service which includes secondary care and the community working together to provide the respiratory care pathways including asthma. The strength of this model is that the community respiratory nurse specialists in reach into the acute hospital and are then able to follow patients in the

	community and the Respiratory Consultant provide the
	community specialist clinics. The shared clinical
	governance structure allows for integration of clinical
	audit across organisations and breaks down barriers for
	patients. The asthma audit is an example where there has
	been partnership working so that the development of the
	asthma proforma enhances the care as well as allows
	audit to occur prospectively.
,	

Appendix 21: Primary care audit: Primary care queries

Primary Care: Asthma Audit Development Project

Audit Queries

(Version – 1.1 Final 31 January 2018

No Question	Rationale
Section 1: Demographics and Co-morbidities	
Number of patients with asthma	
NOTE: Date of diagnosis to be included.	
The following patient demographics will be extracted	For analysis of equity of access.
and reported on:	
• Ethnicity	
Gender	
• Age	
Deprivation score: WIMD/IMD	
The presence of the following co-morbidities will be	To allow assessment of the percentage of asthma patients with co-
assessed:	morbidities (to better categorise the audited cohort).
 Diabetes 	
 Hypertension 	BTS/SIGN guideline
• COPD	Healthcare professionals must be aware that patients with severe
 Bronchiectasis 	asthma and one or more adverse psychosocial factors are at risk of
 Depression 	death. BTS/SIGN 8.1.3
Anxiety	
Schizophrenia, Bipolar and other psychotic	<u>NRAD</u>
illness	Health professionals must be aware of the factors that increase the
CAMHS referral	risk of asthma attacks and death, including the significance of
Learning disability	concurrent psychological and mental health issues.
 Osteoporosis 	
Eczema	NICE quality statements (QS25)
Atopy	Quality Statement 5: Review
Nasal polyps	Assessment of comorbidities is included in the recommended
Reflux	content for annual reviews for both adults and children.
Hay fever	
Family history of asthma	
Allergic rhinitis	
Obesity (BMI)	
, , ,	
NOTES:	
SMI codes will also be used for analysis of parity of	
esteem.	
The first round of audit should be used to explore if this	

list can be amended and/or reduce as there are currently a lot of data point.

Section 2: Asthma exacerbations

The proportion of patients who have a record of an asthma exacerbation treated within primary care in the last 15 months

NOTES:

Coded exacerbations will be compared with the number of exacerbations calculated from use of validated combinations of the following codes:

 ≤300 mg oral corticosteroids (OCS) (not prescribed during an annual review)

A cut-off of <150 mg will be used for children </p>

A cut-off of ≤150 mg will be used for children < 5 years old

 Lower respiratory tract infections treated with same day prescription of appropriate antibiotics (Ax-LRTI)

There must be at least 14 days between each event for them to be considered separate events.

Obtain date of exacerbation.

Enables exploration of exacerbations per demographic, if frequent exacerbations are more likely in certain patient groups.

To investigate if issues around adherence and control are being addressed. Where patients are continuing to have repeated asthma attacks may show that assessments and reviews may not be taking place appropriately.

NICE Quality Statements (QS25)

NICE quality statement 6: Assessing asthma control

People with asthma who present with respiratory symptoms receive an assessment of their asthma control.

The proportion of patients with asthma who have had ≥ 3 courses of prednisolone for asthma exacerbations AND referred for specialist care in the last 15 months

NRAD

Patients with asthma must be referred to a specialist asthma service if they have required more than two courses of systemic corticosteroids, oral or injected, in the previous 12 months or require management using British Thoracic Society (BTS) stepwise treatment 4 or 5 to achieve control.

Section 3: Getting the diagnosis right

Spirometry

1b

2a **Spirometry**

The proportion of patients with asthma who have a record (ever recorded) of spirometry (+ reversibility) test

NOTE:

Result of test to be included in this query. Last recorded measurement with result to be used. Definitions of reversibility testing include:

 Spirometry evidence or trial of treatment with OCS,

OR

 ICS with medcodes for "positive" reversibility testing

BTS/SIGN guideline

Undertake a structured clinical assessment to assess the initial probability of asthma. This should be based on:

- a history of recurrent episodes (attacks) of symptoms, ideally corroborated by variable peak flow when symptomatic and asymptomatic
- symptoms of wheeze, cough, breathlessness and chest tightness that vary over time
- recorded observation of wheeze heard by a healthcare professional
- personal/family history of other atopic conditions (in particular, atopic eczema/dermatitis, allergic rhinitis)
- no symptoms/signs to suggest alternative diagnoses.

OR

	 evidence of variable PEFR Will include all relevant spirometry codes. FEV¹, FVC etc Children under 6 to be excluded from denominator as not commonly used in this cohort. 	Compare the results of diagnostic tests undertaken whilst a patient is asymptomatic with those undertaken when a patient is symptomatic to detect variation over time. Carry out quality-assured spirometry using the lower limit of normal to demonstrate airway obstruction, provide a baseline for assessing response to initiation of treatment and exclude alternative diagnoses. • Obstructive spirometry with positive bronchodilator reversibility increases the probability of asthma. Normal spirometry in an asymptomatic patient does not rule out the diagnosis of asthma.
2b	The proportion of patients with asthma who have had a spirometry (+ reversibility) test in the last 15 months	As above
	NOTE: Result of test to be included in this query. Last recorded measurement and result to be used. Definitions of reversibility testing include: • Spirometry evidence or trial of treatment with OCS, OR • ICS with medcodes for "positive" reversibility testing OR • evidence of variable PEFR Will include all relevant spirometry codes. FEV ¹ , FVC etc	
Peak fl	ow	
3a	Peak flow The proportion of patients with asthma who have a record (ever recorded) of a peak flow test (reading and/or diary records)	BTS/SIGN guideline In adults with no evidence of airflow obstruction on initial assessment, and in whom other objective tests are inconclusive but asthma remains a possibility, consider referral for challenge tests.
	NOTE: Result of test to be included in this query if available (coding of test does not always have result). Last recorded measurement with result to be used.	
3b	The proportion of patients with asthma who have a record of a peak flow test (reading and/or diary records) in the last 15 months	As above
	NOTE: Result of test to be included in this query if available (coding of test does not always have result). Last recorded measurement with result to be used.	

Fractio	nal exhaled nitric oxide (FeNO)	
4a	The proportion of patients with asthma who	BTS/SIGN Guideline
	have a record (ever recorded) of a fractional	Fractional exhaled nitric oxide (FeNO)
	exhaled nitric oxide FeNO test	Use measurement of FeNO (if available) to find evidence of
		eosinophilic inflammation. A positive test increases the probability
	NOTE: Result of test to be included in this	of asthma but a negative test does not exclude asthma.
	query.	
	44-57	
4b	The proportion of patients with asthma who	As above
	have a record of a fractional exhaled nitric oxide	
	FeNO test in the last 15 months	
	NOTE: Result of test to be included in this	
	query.	
	' '	
Record	of ANY objective measurement	
5	The proportion of patients with asthma who	As above for spirometry, peak flow and FeNO test.
	have a (ever recorded) record of <u>any</u> objective	
	measurement	
	Spirometry (+reversibility)	
	Peak Flow	
	Fractional exhaled nitric oxide FeNO	
Section	1 4: Personalised Asthma Action Plans	
6	The proportion of people with asthma who	NICE Quality Statements (QS25)
	have had a Personalised Asthma Action Plan	NICE quality standard 3: Written personalised action plans
	(PAAP) anytime in the last 15 months.	People with asthma receive a written personalised action plan.
	NOTE: Look at results for 1-5 and 6-18 year olds.	BTS/SIGN guideline
	NOTE. LOOK at results for 1-3 and 0-18 year olds.	All people with asthma (and/or their parents or carers) should be
		offered self-management education which should include a written
		personalised asthma action plan and be supported by regular
		professional review.
		In adults, written personalised asthma action plans may be based on
		symptoms and/or peak flows: symptom-based plans are generally
		preferable for children.
		Written PAAPs (for example, those for adults and children from
		Asthma UK, available at www.asthma.org.uk/resources/#action
		plans) are crucial components of effective self-management
		education. One systematic review identified the features of PAAPs
		associated with beneficial outcomes. These include:
		 specific advice about recognising loss of asthma control,
		assessed by symptoms or peak flows or both.91,150,151 In
		children, symptom-based written plans are effective in
		reducing emergency consultations for asthma, although (in
		older children) peak flow-based plans may be as effective for
		other outcomes.
		actions, summarised as two or three action points, to take if
<u> </u>	<u> </u>	· · · · · · · · · · · · · · · · · · ·

asthma deteriorates, including seeking emergency help, starting oral steroids (which may include provision of an emergency course of steroid tablets), restarting or temporarily increasing (as opposed to just doubling) ICS, as appropriate to clinical severity.

Summary of table 8 from guideline

- Current medication and adherence
- Triggers and exacerbating factors
- Advice on taking current medication
- Advice on increasing medication (action points)
- Advice on calling for help
- Peak flow levels at which action should be taken
- A record of best personal peak expiratory flow (PEF)
- A record of best predicted peak expiratory flow (PEF)

NRAD

All people with asthma should be provided with written guidance in the form of a personal asthma action plan (PAAP) which details their own triggers and current treatment, and specifies how to prevent relapse and when to seek help in an emergency.

Section 5: Providing the right care

7 The proportion of people with asthma who have had an annual review in the last 15 months.

NOTE: This query will just establish the number of patients who have had an annual appointment to review their asthma. Due to the fact that this does not necessarily mean that all annual review elements will be reviewed together (at this appointment), but still reviewed annually this query has been separated from the next.

NICE Quality Statements (QS25)

NICE quality statement 4: Inhaler technique

People with asthma are given specific training and assessment in inhaler technique before starting any new inhaler treatment

NICE quality statement 5: Review

People with asthma receive a structured review at least annually.

NRAD

People with asthma should have a structured review by a healthcare professional with specialist training in asthma, at least annually. People at high risk of severe asthma attacks should be monitored more closely, ensuring that their personal asthma action plan (PAAPs) are reviewed and updated at each review.

An assessment of inhaler technique to ensure effectiveness should be routinely undertaken and formally documented at annual review, and also checked by the pharmacist when a new device is dispensed. An assessment of recent asthma control should be undertaken at every asthma review. Where loss of control is identified, immediate action is required, including escalation of responsibility, treatment change and arrangements for follow-up.

BTS/SIGN guideline

In primary care, people with asthma should be reviewed regularly by a nurse or doctor with appropriate training in asthma management. Review should incorporate a written action plan.

Before initiating a new drug therapy practitioners should check adherence with existing therapies, inhaler technique and eliminate

trigger factors. BTS/SIGN 2.4

Monitoring children in primary care

Asthma is best monitored in primary care by routine clinical review on at least an annual basis (see section 14.3).

The factors that should be monitored and recorded include:

- symptom score, eg Children's Asthma Control Test, Asthma Control Questionnaire
- asthma attacks, oral corticosteroid use and time off school/nursery due to asthma since last assessment
- inhaler technique
- adherence, which can be assessed by reviewing prescription refill frequency
- possession of and use of a self-management plan/written personalised asthma action plan
- exposure to tobacco smoke
- growth (height and weight centile).

Monitoring adults in primary care

In adults the following factors should be monitored and recorded in primary care:

- symptomatic asthma control
- lung function assessed by spirometry or by PEF
- asthma attacks,
- oral corticosteroid use and time off work since last assessment
- inhaler technique
- adherence
- bronchodilator reliance
- possession of and use of a self-management plan/personal action plan

QOF and NICE Quality Statements (QS25)

Paediatrics

- Assessment of symptomatic asthma control using recognised tool (RCP 3 questions, asthma control questionnaire, children's asthma control test, paediatric asthma quality of life questionnaire)
- Review of exacerbations, oral corticosteroid use and time off school or nursery since last assessment
- Inhaler technique
- Assessing adherence (review of prescriptions)
- Adjustment of treatment
- Possession and review of PAAP
- Exposure to tobacco smoke
- Measurement of growth centile (height and weight)
- Assessment of comorbidities
- Review of diagnosis

Adults

• Assessment of symptomatic asthma control using recognised tool

		(RCP 3 questions, asthma control questionnaire, asthma control
		test, asthma quality of life questionnaire)
		Measurement of lung function, assessment by spirometry or by
		peak expiratory flow
		 Review of exacerbations, oral corticosteroid use and time off work
		or study since last assessment
		Checking inhaler technique
		Assessing adherence (review of prescriptions)
		Adjustment of treatment
		Bronchodilator reliance (review of prescriptions)
		Possession and review of PAAP
		• Smoking status
		Assessment of comorbidities
		Review of diagnosis
		• Review of diagnosis
8	The proportion of patients with asthma who	BTS/SIGN guideline
	have a record of the RCP 3 questions being	In adults the following factors should be monitored and recorded in
	asked in the last 15 months	primary care:
		symptomatic asthma control
	NOTE: Only if the RCP 3 questions were asked	(Symptomatic asthma control is best assessed using directive
	can be captured, not the responses to them.	questions such as the Royal College of Physicians' '3 questions',129 or the Asthma Control Questionnaire or Asthma Control Test (see
		Table 7), since broad non-specific questions may underestimate
		symptoms)
		, ,
		Also see annual review rationale.
9	The proportion of people with asthma who	NICE Quality Statements (QS25)
	have received an inhaler check within the last	NICE quality statement 4: Inhaler technique
	15 months.	People with asthma are given specific training and assessment in
		inhaler technique before starting any new inhaler treatment
		BTS/SIGN guideline
		Before initiating a new drug therapy practitioners should check
		adherence with existing therapies, inhaler technique and eliminate
		trigger factors. BTS/SIGN 2.4
		Chapter ractors projective
		NRAD
		An assessment of inhaler technique to ensure effectiveness should
		be routinely undertaken and formally documented at annual review,
		and also checked by the pharmacist when a new device is dispensed.
		Monitoring children in primary care
		Asthma is best monitored in primary care by routine clinical review
		on at least an annual basis (see section 14.3).
		The factors that should be monitored and recorded include:
		symptom score, eg Children's Asthma Control Test, Asthma
		Control Questionnaire
		asthma attacks, oral corticosteroid use and time off
		school/nursery due to asthma since last assessment
		inhaler technique
	l .	

	T	
10a	The proportion of people with asthma who were recorded as being exposed to second hand smoke in the last 15 months.	 adherence, which can be assessed by reviewing prescription refill frequency possession of and use of a self-management plan/written personalised asthma action plan exposure to tobacco smoke growth (height and weight centile). Monitoring adults in primary care In adults the following factors should be monitored and recorded in primary care: symptomatic asthma control lung function assessed by spirometry or by PEF asthma attacks, oral corticosteroid use and time off work since last assessment inhaler technique adherence bronchodilator reliance possession of and use of a self-management plan/personal action plan BTS/SIGN guideline Parents with asthma should be advised about the dangers, to themselves and to their children with asthma, of smoking, and be offered appropriate support to stop smoking. Clinicians should be aware that higher doses of inhaled corticosteroids may be needed in patients who are smokers or exsmokers. NRAD A history of smoking and/or exposure to second-hand smoke should
		Current smokers should be offered referral to a smoking-cessation
		service.
10b	The proportion of people with asthma who	As above
	were recorded as a current smoker at any time	
	in the last 15 months and have received or been	
	'sign-posted to' to a behavioral change	
	intervention.	
10c	The proportion of people with asthma who	As above
	were recorded as a current smoker at any time	
	in the last 15 months and had a stop-smoking	
	drug prescribed.	
11	The proportion of people with asthma who	BTS/SIGN guideline
	have been prescribed more than 12 short-acting	Anyone prescribed more than one short-acting bronchodilator
	reliever inhalers in the last 12 months	inhaler device a month should be identified and have their asthma
		assessed urgently and measures taken to improve asthma control if
	[Time between the 12 SABA alert and the next	this is poor.
	asthma review code (query 12) will be	NRAD
	calculated]	NRAD

All asthma patients who have been prescribed more than 12 shortacting reliever inhalers in the previous 12 months should be invited for urgent review of their asthma control, with the aim of improving their asthma through education and change of treatment. 12 The proportion of people with asthma who BTS/SIGN guideline have been prescribed the following in the last 6 Long-acting inhaled β2 agonists should only be started in patients months: who are already on inhaled corticosteroids, and the inhaled corticosteroid should be continued. · a single component long-acting beta Combination inhalers are recommended to: agonists (LABA) guarantee that the long-acting $\beta 2$ agonist is not taken without • a single component inhaled corticosteroid (ICS) inhaled corticosteroid • improve inhaler adherence a single component LTRA • a combined LABA and ICS or MART inhaler a combined ICS and LTRA NRAD The use of combined inhalers should be encouraged. Where longacting beta agonist (LABA) bronchodilators are prescribed for people with asthma, they should be prescribed with an inhaled corticosteroid in a single combination inhaler. 13 The proportion of patients with asthma who NRAD have been prescribed fewer than 4 ICS devices Non-adherence to preventer inhaled corticosteroids is associated with increased risk of poor asthma control and should be continually in the last year. monitored. **BTS/SIGN Guideline** Inhaled corticosteroids are the recommended preventer drug for adults and children for achieving overall treatment goals. Inhaled corticosteroids should be considered for patients with any of the following asthma-related features: asthma attack in the last two years using inhaled $\beta 2$ agonists three times a week or more symptomatic three times a week or more waking one night a week Give inhaled corticosteroids initially twice daily (except ciclesonide which is given once daily). Once a day inhaled corticosteroids at the same total daily dose can be considered if good control is established. Before initiating a new drug therapy practitioners should check adherence with existing therapies, check inhaler technique, and eliminate trigger factors. Adherence to long-term asthma treatment should be routinely and regularly addressed by all healthcare professionals within the context of a comprehensive programme of accessible proactive asthma care. BTS/SIGN 2.2

Appendix 22: Removed queries: Primary care audit: Primary care queries

Query	Original rationale
	REASON FOR REMOVAL
Asthma exacerbations	
The proportion of patients who had an asthma exacerbation in the last 15 months, who were treat with any of the following for their exacerbation	Enables exploration of how exacerbations of asthma are being treatment and which, if any, are more effective in preventing future attacks.
 Oral steroids Antibiotics Spacers (+ nebuliser) Inhaler 	RCEM dataset Included in dataset Discharged patients should have oral prednisolone prescribed.
	BTS/SIGN guideline Give Steroids in adequate doses for all acute attacks. BTS/SIGN 2.6.1 Continue prednisolone 40–50mg daily for at least 5 days or until recovery. BTS/SIGN 8.3.3
	REASON FOR REMOVAL: This information would be very poorly coded.
The proportion of patients with asthma who have attended an emergency department with an exacerbation of asthma in the last 15 months.	Enables exploration of exacerbations per demographic, if frequent exacerbations are more likely in certain patient groups. Also investigation into if issues around adherence and control are being addressed. Where patients are continuing to have repeated asthma attacks may show that assessments and follow ups are not taking place.
	NICE Quality Statements (QS25) NICE quality statement 6: Assessing asthma control People with asthma who present with respiratory symptoms receive an assessment of their asthma control. NICE quality statement 10: Follow up in Primary Care People who received treatment in hospital or through out-of-hours services for an acute exacerbation of asthma are followed up by their own GP practice within 2 working days of treatment.
	BTS/SIGN guideline Follow up should be arranged prior to discharge with the patient's general practitioner or asthma nurse within two working days and with a hospital specialist asthma nurse or respiratory physician at about one month after admission. BTS/SIGN 8.6.3

NRAD Follow-up arrangements for patients who have attended ED or out of hours services. **REASON FOR REMOVAL: This is only possible with linked** data to establish that the coded exacerbation was actually treated in ED, rather than primary care. The proportion of patients with asthma who have been Enables exploration of exacerbations per demographic, if admitted to hospital with an exacerbation of asthma in the frequent exacerbations are more likely in certain patient last 15 months. groups. Also investigation into if issues around adherence and control are being addressed. Where patients are continuing to have repeated asthma attacks may show that assessments and follow ups are not taking place. **NICE Quality Statements (QS25)** NICE quality statement 6: Assessing asthma control People with asthma who present with respiratory symptoms receive an assessment of their asthma control. NICE quality statement 10: Follow up in Primary Care People who received treatment in hospital or through out-ofhours services for an acute exacerbation of asthma are followed up by their own GP practice within 2 working days of treatment. **BTS/SIGN** guideline Follow up should be arranged prior to discharge with the patient's general practitioner or asthma nurse within two working days and with a hospital specialist asthma nurse or respiratory physician at about one month after admission. BTS/SIGN 8.6.3 NRAD Follow-up arrangements for patients who have attended ED or out of hours services. **REASON FOR REMOVAL: This is only possible with linked** data to establish that the coded exacerbation was actually treated in ED, rather than primary care. The proportion of patients with asthma who have attended As for hospital/emergency department rationale. out of hours (OOH)/walk-in/urgent care centres with an exacerbation of asthma in the last <time period> **REASON FOR REMOVAL: Aspirational at the moment, as not** coded and would need to approach/recruit individual OOH/walk-in services for information.

Option: Aspirational, consider again at later date when coding

has improved?

The proportion of patients with asthma who received a review within 2 days of discharge from hospital or emergency department for an exacerbation of asthma

Enables exploration of follow up per demographic, if less likely in certain patient groups. Also investigation into if issues around adherence and control are being addressed. Where patients are continuing to have repeated asthma attacks may show that assessments and follow ups are not taking place.

NICE Quality Statements (QS25)

NICE quality statement 6: Assessing asthma control

People with asthma who present with respiratory symptoms receive an assessment of their asthma control.

NICE quality statement 10: Follow up on Primary Care
People who receive treatment in hospital or through out-ofhours services for an acute exacerbation of asthma are
followed up by their own GP practice within 2 working dates
of treatment.

BTS/SIGN guideline

Follow up should be arranged prior to discharge with the patient's general practitioner or asthma nurse within two working days and with a hospital specialist asthma nurse or respiratory physician at about one month after admission. BTS/SIGN 8.6.3

NRAD

Follow-up arrangements must be made after every attendance at an emergency department or out-of-hours service for an asthma attack. Secondary care follow-up should be arranged after every hospital admission for asthma, and for patients who have attended the emergency department two or more times with an asthma attack in the previous 12 months.

REASON FOR REMOVAL: This is not possible as exacerbations treated in emergency departments and hospitals cannot be captured without linked data.

Getting the diagnosis right

The proportion of patients with asthma who have a (ever recorded) record of <u>any</u> objective measurement

- Objective response recorded on a symptom score (following a trial of treatment)
- RCP 3 questions

BTS/SIGN guideline

Adults and children with airways obstruction Asthma is the by far the commonest cause of airways obstruction identified through spirometry in children. Obstruction due to other disorders is much more common in adults than in children. Patients may have more than one cause of airflow obstruction, which complicates the interpretation of any test. In particular, asthma and COPD commonly coexist in adults. A bronchodilator reversibility test and/or a monitored initiation of treatment (typically six weeks of inhaled corticosteroids (ICS) can establish whether or not the airflow obstruction reverses to normal with treatment. Evidence of a symptomatic response, ideally using objective measures of

asthma control and lung function, should be sought at a follow-up visit. If there is significant reversibility or improvement in symptom scores, confirm the diagnosis of asthma and record the basis on which the diagnosis was made. Continue to treat as asthma, but aim to find the minimum effective dose of therapy. If the patient remains asymptomatic consider a trial of reduction or withdrawal of treatment. This is particularly important in children in whom natural resolution of symptoms is more common than in adults.

In adults and children with an intermediate probability of asthma and airways obstruction identified through spirometry, undertake reversibility tests and/ or a monitored initiation of treatment assessing the response to treatment by repeating lung function tests and objective measures of asthma control.

REASON FOR REMOVAL: Symptom score and RCP 3 not considered objective measurements.

The proportion of patients with asthma who have been diagnosed with asthma as per NICE guidelines (2017)

Rationale note included as not published at point of removal.

REASON FOR REMOVAL: Will be captured via other diagnosis queries.

Addition of 'consistent with a diagnosis of asthma' was considered for diagnosis queries.

REASON FOR NON-INCLUSION: No diagnostic test is 100% for asthma. A patient could have a normal spirometry but diagnosis is made via another test.

Personalised Asthma Action Plans (PAAPs)

The proportion of people with asthma who have had triggers and exacerbating factors identified and recorded within the last 15 months

Triggers and exacerbating factors identified and recorded:

- Food allergy
- Animal allergy
- Virus infection/URTIs
- Drugs (e.g. NSAIDs)
- Exercise
- Tobacco
- Occupational factor

NRAD

Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the significance of concurrent psychological and mental health issues.

Patient self-management should be encouraged to reflect their known triggers, eg increasing medication before the start of the hay-fever season, avoiding non-steroidal anti-inflammatory drugs or by the early use of oral corticosteroids with viral- or allergic-induced exacerbations.

BTS/SIGN guideline

Every asthma consultation is an opportunity to review, reinforce and extend both the patient's knowledge and skills. This is true whether the patient is seen in primary care, the ED or the outpatient clinic. It is important to recognise that education is a process and not a single event. A hospital admission represents a window of opportunity to review self-management skills. No patient should leave hospital without a written personalised asthma action plan.

An acute consultation offers the opportunity to determine

what action the patient has already taken to deal with the asthma attack. Their self-management strategy may be reinforced or refined and the need for consolidation at a routine follow up considered.

A consultation for an upper respiratory tract infection or other known trigger is an opportunity to rehearse with the patient their self- management in the event of their asthma deteriorating.

Education should include personalised discussion of issues such as trigger avoidance and achieving a smoke-free environment to support people and their families living with asthma.

Brief simple education linked to patient goals is most likely to be acceptable to patients.

REASON FOR REMOVAL: This will be under estimated as existence of triggers is only noted initially, with individual triggers then mainly captured in free text (which is not analysable). Those which can be captured as co-morbidities have be moved to the co-morbidities query.

Annual review

The proportion of people with asthma who have received an annual review of each of the following in the last 15 months:

Paediatrics

- Assessment of symptomatic asthma control using recognised tool (RCP 3 questions, asthma control questionnaire, children's asthma control test, paediatric asthma quality of life questionnaire)
- Review of exacerbations, oral corticosteroid use and time off school or nursery since last assessment
- Inhaler technique
- Assessing adherence (review of prescriptions)
- Adjustment of treatment
- Possession and review of PAAP
- Exposure to tobacco smoke
- Measurement of growth centile (height and weight)
- Assessment of comorbidities
- Review of diagnosis

Adults

- Assessment of symptomatic asthma control using recognised tool (RCP 3 questions, asthma control questionnaire, asthma control test, asthma quality of life questionnaire)
- Measurement of lung function, assessment by spirometry or by peak expiratory flow

NICE Quality Statements (QS25)

NICE quality statement 4: Inhaler technique

People with asthma are given specific training and assessment in inhaler technique before starting any new inhaler treatment

NICE quality statement 5: Review

People with asthma receive a structured review at least annually.

NRAD

People with asthma should have a structured review by a healthcare professional with specialist training in asthma, at least annually. People at high risk of severe asthma attacks should be monitored more closely, ensuring that their personal asthma action plan (PAAPs) are reviewed and updated at each review.

An assessment of inhaler technique to ensure effectiveness should be routinely undertaken and formally documented at annual review, and also checked by the pharmacist when a new device is dispensed.

BTS/SIGN guideline

In primary care, people with asthma should be reviewed regularly by a nurse or doctor with appropriate training in asthma management. Review should incorporate a written action plan.

Before initiating a new drug therapy practitioners should check adherence with existing therapies, inhaler technique

- Review of exacerbations, oral corticosteroid use and time off work or study since last assessment
- Checking inhaler technique
- Assessing adherence (review of prescriptions)
- Adjustment of treatment
- Bronchodilator reliance (review of prescriptions)
- Possession and review of PAAP
- Smoking status
- Assessment of comorbidities

Review of diagnosis

and eliminate trigger factors. BTS/SIGN 2.4

Monitoring children in primary care

Asthma is best monitored in primary care by routine clinical review on at least an annual basis (see section 14.3).

The factors that should be monitored and recorded include:

- symptom score, eg Children's Asthma Control Test, Asthma Control Questionnaire
- asthma attacks, oral corticosteroid use and time off school/nursery due to asthma since last assessment
- inhaler technique
- adherence, which can be assessed by reviewing prescription refill frequency
- possession of and use of a self-management plan/written personalised asthma action plan
- exposure to tobacco smoke
- growth (height and weight centile).

Monitoring adults in primary care

In adults the following factors should be monitored and recorded in primary care:

- symptomatic asthma control
- lung function assessed by spirometry or by PEF
- asthma attacks,
- oral corticosteroid use and time off work since last assessment
- inhaler technique
- adherence
- bronchodilator reliance
- possession of and use of a self-management plan/personal action plan

QOF and NICE Quality Statements (QS25)

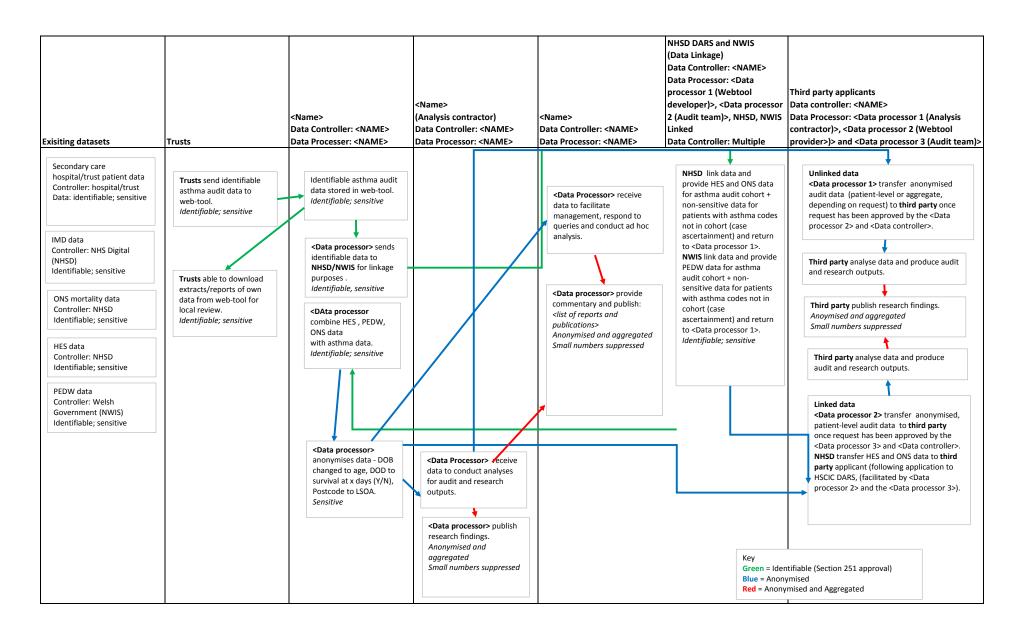
Paediatrics

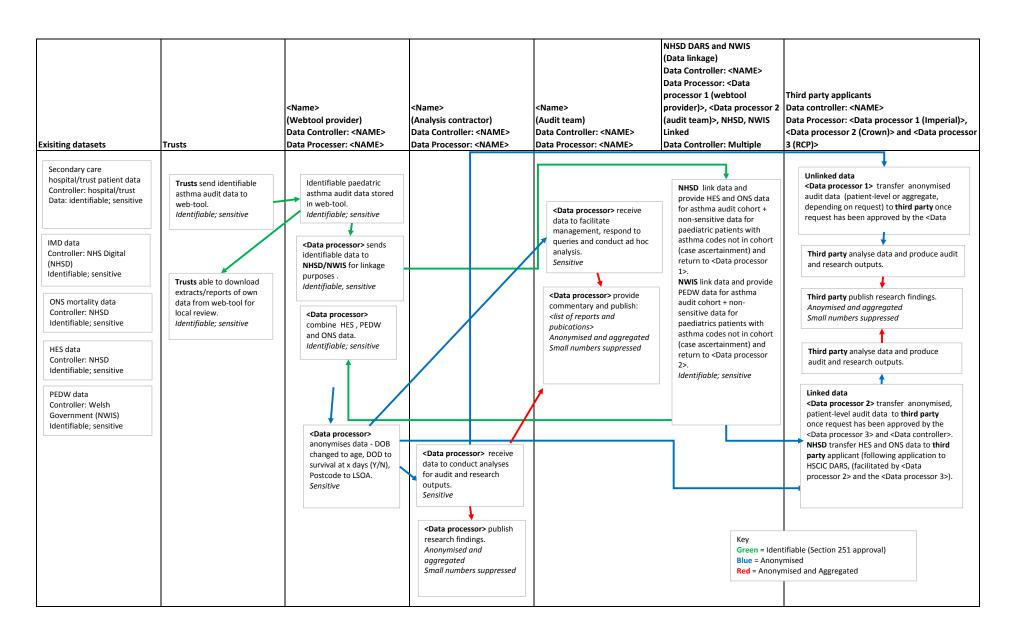
- Assessment of symptomatic asthma control using recognised tool (RCP 3 questions, asthma control questionnaire, children's asthma control test, paediatric asthma quality of life questionnaire)
- Review of exacerbations, oral corticosteroid use and time off school or nursery since last assessment
- Inhaler technique
- Assessing adherence (review of prescriptions)
- Adjustment of treatment
- Possession and review of PAAP
- Exposure to tobacco smoke
- Measurement of growth centile (height and weight)
- Assessment of comorbidities
- Review of diagnosis

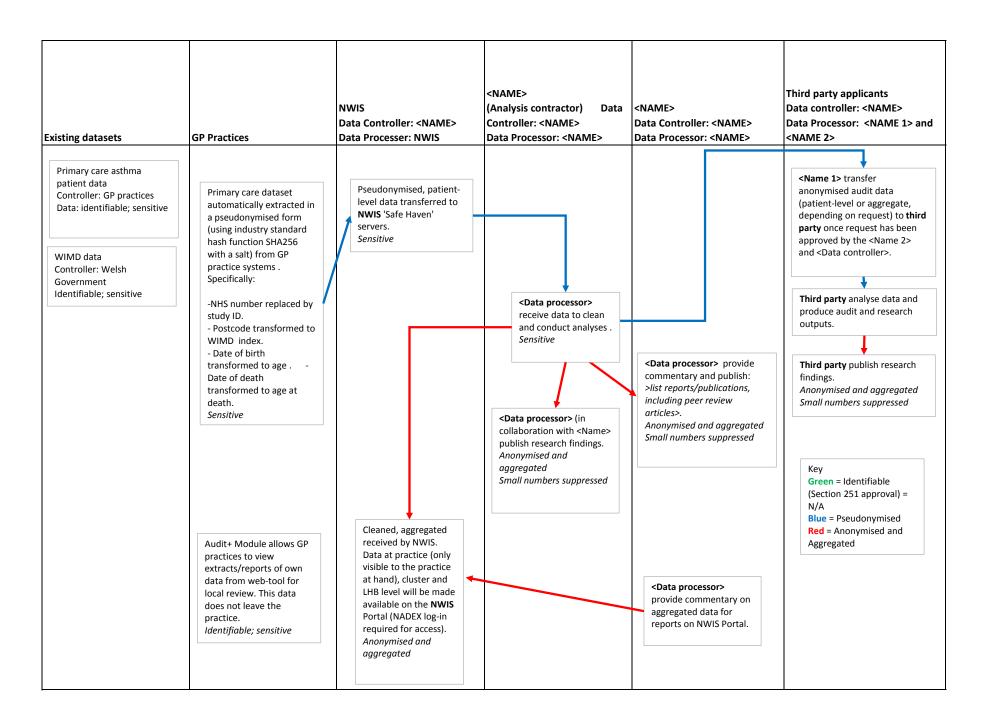
Adults • Assessment of symptomatic asthma control using recognised tool (RCP 3 questions, asthma control questionnaire, asthma control test, asthma quality of life questionnaire) • Measurement of lung function, assessment by spirometry or by peak expiratory flow • Review of exacerbations, oral corticosteroid use and time off work or study since last assessment • Checking inhaler technique • Assessing adherence (review of prescriptions) Adjustment of treatment • Bronchodilator reliance (review of prescriptions) • Possession and review of PAAP Smoking status Assessment of comorbidities • Review of diagnosis **REASON FOR REMOVAL: Individual annual review elements** are not coded, only the fact that an annual review has taken place. The proportion of people who have had their current See rationale under annual review. medication and adherence assessed at their year review **REASON FOR REMOVAL: As per reason for removal of** individual annual review elements, this is not coded. Only that an annual review as taken place. Providing the right care The proportion of people with asthma who have been NRAD referred to secondary/tertiary care. Patients with asthma must be referred to a specialist asthma service if they have required more than two courses of systemic corticosteroids, oral or injected, in the previous 12 months or require management using British Thoracic Society (BTS) stepwise treatment 4 or 5 to achieve control. **REASON FOR REMOVAL: There will be some practices who** use generic referral codes and some who don't use any. Data would therefore be too unreliable. Option: 1. Aspirational, consider again at later date once coding has improved? 2. Pull in initial round and report confirmation that reporting is too bad to use. The proportion of patients with asthma who have had the BTS/SIGN guideline influenza immunisation in the preceding <DATE>. Immunisations should be administered independent of any considerations related to asthma. Responses to vaccines may be attenuated by high-dose inhaled corticosteroids.

	People with chronic respiratory illness who are infected with
	the influenza virus have more serious illness and are at higher
	risk of mortality. The vaccine has variable effectiveness
	according to season and current health status when given. It
	is safe and the highest value intervention for the treatment of
	COPD but is used less than some other COPD interventions
	that have less value.
	that have less value.
	REASON FOR REMOVAL: Evidence not strong for asthma as a
	heterogenous condition with varying severity.
Proportion of patients with asthma who have been	BTS/SIGN guideline
screened for depression and anxiety or who have been	Healthcare professionals must be aware that patients with
diagnosed with these conditions within the last 2 years.	severe asthma and one or more adverse psychosocial factors
	are at risk of death. BTS/SIGN 8.1.3
	NRAD
	Health professionals must be aware of the factors that
	increase the risk of asthma attacks and death, including the
	significance of concurrent psychological and mental health
	issues.
	REASON FOR REMOVAL: Being captured within co-
	morbidities section
Proportion of people with asthma who have been	REASON FOR REMOVAL: Majority of query keep but
prescribed more than 12 short acting reliever inhalers in	'invitation to urgent review' removed as this is not coded.
the last 12 months and been invited to an urgent review of	
their asthma control.	
Proportion of patients with asthma who have received >2	REASON FOR REMOVAL: No evidence for inclusion
courses of oral corticosteroids in the last year.	
Proportion of children with asthma who have a record of	BTS/SIGN guideline
mental health service access, self-harm or overdose.	Healthcare professionals must be aware that patients with
mental health service access, sen-harm of overdose.	severe asthma and one or more adverse psychosocial factors
	are at risk of death. BTS/SIGN 8.1.3
	NRAD
	Health professionals must be aware of the factors that
	increase the risk of asthma attacks and death, including the
	significance of concurrent psychological and mental health
	issues.
	REASON FOR REMOVAL: Captured within co-morbidities
	query. If reinstated, must align with wording used in
	secondary care datasets.
The proportion of patients with asthma who have a	BTS/SIGN guideline
recorded learning disability.	Healthcare professionals must be aware that patients with
	severe asthma and one or more adverse psychosocial factors
	are at risk of death. BTS/SIGN 8.1.3
	·

	NRAD Health professionals must be aware of the factors that increase the risk of asthma attacks and death, including the
	significance of concurrent psychological and mental health issues.
	REASON FOR REMOVAL: Added to co-morbidities
Proportion of children with asthma whose height has been	REASON FOR REMOVAL: Component of annual review. No
recorded in the last year.	evidence for including individually.
Proportion of children with asthma how are exposed to	REASON FOR REMOVAL: Will be captured via
smoke at home	smoking/exposure to smoke query, not need individually.
Transitional care (additional analysis)	REASON FOR REMOVAL: Research, rather than audit.
Look at frequency of exacerbations in teenage to 25 year	
age group compared to rest of population	
Look at smoking status and advice given to teenage to 25 year age group compared with rest of population (already measured in QOF)	
Organisational information	
Staffing levels	REASON FOR REMOVAL: Not coded, would need to
Asthma diplomas	approach each individual GP practice for this information.
Frequency of asthma clinics	
Average time of asthma reviews	
Content of referral letters	











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1. Background to Research

The Royal College of Physicians (RCP) have been commissioned by the Healthcare Quality Improvement Partnership (HQIP) to deliver the Asthma Audit Development Project (AADP). The AADP will inform the specification of, and carry out some of the work required for, a National Asthma Audit which will start in 2018. The National Asthma Audit will collect information on every asthma patient who is admitted to hospital with an asthma attack in England and Wales, and ensure that their care matches existing asthma standards and guidelines. It has recently been confirmed that the National Asthma Audit will also cover Scotland. The National Asthma Audit will also develop a set of primary care queries for the purposes of direct data extraction from primary care practices in Wales, to provide insight into primary care practice and treatment of asthma patients. Exploration on how this can also be done in England and Scotland will be carried out. To ensure that asthma patients were able to contribute their views on how a National Asthma Audit should be conducted, RCP commissioned Asthma UK to undertake a focus group with a group of their Research and Policy volunteers.

2. Research Aims

To explore and understand:

- What people with asthma think is important to them about their asthma care
- What people with asthma think is important to record in an asthma audit
- Why a national asthma audit is important to people with asthma
- How people with asthma would like the draft patient level report and infographic to be

3. Method: Focus group

A focus group was carried out because it offers an interactive method to explore the views of people with asthma to meet the above research aims. The participants were given a brief written overview of the AADP prior to attending, and the example of the primary care section of the patient level report and COPD infographic were shared with the group when this point of the discussion was reached on the day. Although it is difficult to make firm conclusions based on just one focus group, it is sufficient to provide indicative answers to the aims and explore the thoughts and ideas of people with asthma in-depth.

3.1. Recruitment

- The focus group had 9 participants recruited using Asthma UK's network of Research and Policy volunteers. A recruitment specification was used to ensure a suitable mix of participants. The group contained representation from:
 - o Both men and women
 - o A range of ages from teens to 70's
 - A range of geographical locations across England and Wales
 - White British and Non-White British ethnicities
 - o A carer of someone with asthma
 - o A range of contact time with emergency hospital care
 - People diagnosed with asthma as both a child and adult

3.2. Facilitation

- The focus group took place on Saturday 21st October 2017 at Asthma UK's office in London.
- RCP's AADP Project Manager attended the focus group to observe the feedback first hand, and had the chance to ask questions at the end.

3.3. Analysis

- The focus group discussions were audio-recorded and transcribed professionally. Thematic analysis of the transcript was conducted in relation to the research aims.
- The report contains quotes to help bring life and evidence to the findings. The quotes have been chosen to best demonstrate a point.
- Due to the qualitative nature of the work, the level of opinion is presented textually rather than quantitatively.

4. Findings

4.1. Important attributes of routine asthma care

A number of logistical points were considered important for routine care:

- Most participants raised the importance of the consistency with who you see for routine care.
- Some participants raised the experience that when they see an asthma specialist (e.g. asthma nurse) rather than a general nurse/GP they are more knowledgeable about asthma.
- Most participants expressed the importance of getting an appointment when required.

The group highlighted the content of routine care appointments that is important to them:

• The importance of the asthma action plan was strongly expressed by the majority of participants: "The creation of a plan. It all comes down to, for us, if we let it slip it will go. And it can spiral out of control and I'll be in hospital for a week."

(Participant 9)

 There were mixed views about asthma reviews, with most participants considering them to be important if done properly. There was concern that not much action or change came from some reviews:

"On the other hand, the six monthly check up with the nurse, I slightly feel is a complete waste of time. And I took, trying to trigger a little bit more action, I took the Asthma UK's printed out from the internet action plan to fill in with her. And she looked at it and said, oh well I don't know how to fill that in, I mean, after all everybody's different. Which I thought was the point of it. But I just said, okay, fine, I'm an intelligent educated woman, I can work it out love, don't worry."

(Participant 2)

- Triggers were raised as an important discussion point for a routine appointment by most participants.
- Most participants raised the importance of an inhaler technique check:

"But in reality nobody has ever in the last 20 years asked to see me using an inhaler and check that I am using it correctly. So, I think although it's a box ticking exercise, there is something about asking your GP practice are you sure that this person knows, have you actually checked while they've been with you."

(Participant 2)

And there's another massive gap as in I believe, we weren't told properly how to use your inhaler until about last year.

(Participant 9)

• The majority of the group stressed the importance of receiving the correct medication and dose for their asthma and symptoms:

"And it was only by accident in '91, when I went to a well man clinic, the person that was doing it was an asthma Nurse Specialist. And that made a huge difference because she then concentrated for the next 12 months on getting a preventer right, etc. etc."

(Participant 2)

"And you know, I was told when I moved out of London that I was on a children's dose of preventer and I shouldn't have never have been on it. I was on it for like 10 years living in London."

(Participant 1)

• Most the group mentioned the importance of specialist diagnosis to ensure that they are on the correct treatment path. The nitric oxide test was discussed as being experienced by a couple of participants, some discussed the trial and error of treatments as a first port of call in a negative way: "And so, we were referred to a consultant at X who said, I'm gonna deal with your allergies and your asthma because they're connected. And the first thing he did was do the nitric oxide testing and actually did a very proper diagnosis to his asthma"

...

"That's the diagnosis, that's the proper diagnosis. Not just a here have an inhaler and see if it gets any better [laughs], you know. Cos, somebody may have taken the flowers that was causing your problem out of the house or, there's all sorts of things that can trigger these."

(Participant 8)

- The assessment of comorbidities like allergies was considered important to some members of the group, and one participant raised this as an important part of diagnosis.
- Some participants discussed importance of a referral to a specialist consultant for a full asthma review and/or access to severe asthma treatments where necessary.
- The participants highlighted the importance of a review of any emergency hospital care with their GP/nurse.
- One of the participants mentioned the need for doctors to speak to the school of a child with severe asthma to discuss their needs.

Further to GP surgeries providing routine care, the availability of nebulisers at surgeries particularly in rural areas was considered important by the group.

4.2. Important attributes of emergency hospital care

On arrival and during stay

 Most respondents raised the correct asthma severity assessment and consequential treatment to be of high importance:

"...But at the hospital, quite often we've gone, been made because their standard operating kind of practices are that you've got to do 10 puffs on your inhaler, then wait and then do that another nine times before they then say, oh no it's quite bad actually. And then, like I say, he's gone on to have five, sixfold ... nebulisers. And they're going, do you know what we can't stabilise this and he's been in and then admitted for a week, a week after that. That was crazy."

(Participant 8)

...

"So, for me that's ... I mean it's challenging. It's almost like it's a near miss really. So, they have been applying an incorrect treatment package for an emergency situation and they've not been quick enough to respond to those challenges..."

(Participant 8)

- One participant discussed the importance of their assessment and treatment by the paramedics before arrival at hospital.
- One participant strongly expressed the importance of allergy assessment on admission to hospital: "Allergies as well. Honestly, it's the most dangerous place we've ever found for eating is in a hospital. The number of times I've been there and then tried to get something to eat and then she's basically just pulled the spoon out of the macaroni and cheese and just stuck it in the chips to try and give me some food. Just stuff like that"

(Participant 9)

Review and discharge

- There was discussion in the group of the following being important to be included in the review at discharge:
 - Why the patient ended up in hospital
 - o Medication review
 - o Inhaler technique check
 - o Referral to a specialist consultant and access to correct treatment
- One participant mentioned the importance of not discharging a patient prematurely and putting them at risk and potentially causing readmission.
- One participant discussed the length of hospital stay as a reflection of how appropriate the treatment received was.

4.3. Important to know about patient to assess care received

• The importance of the age of patient was strongly expressed throughout the group, particularly in relation to children being different and looked at in isolation. Older patients with dementia were also discussed by some of the group as being important to look at in isolation:

"And I'd be intrigued to know how in those situations where we've got elderly care, that there's an asthma care plan in place and how those people have responded to that. And whether or not they're being monitored and how they are being monitored. Are they being monitored on the basis that they're going to express their own difficulty of breathing, or is somebody trained to spot it. And I think I know what the answer to that question is."

(Participant 8)

"Totally. And I think the same applies for young children."

(Participant 7)

• A couple of participants raised the difficulty of dealing with late onset asthma, raising the importance of age and date of diagnosis to identify this group:

"So, I think it's more difficult for people with late onset asthma because I don't know any ... I really don't know any different."

(Participant 7)

• Triggers were raised as important by the majority of participants.

"Triggers. I think it would be nice if they're collecting information about people with asthma and their triggers, it would be good to have that nationally to find out you know is it pollution. Is pollution a trigger. Is smoking kind of thing. Is it dust like"

(Participant 1)

• Some participants mentioned the importance of comorbidities and the symptoms and side effects from the various treatments.

"Relationships to comorbidities. If they've got other things which have developed in their life, it's quite important."

(Participant 8)

- One participant raised the importance of family history.
- Some participants discussed the importance of identifying whether a person with asthma has mental health conditions, dementia and/or learning difficulties.

"...But also at the other end of the scale, if you are moving in to an area where you've got patients who are suffering from dementia or any ... it's a hugely risky area for people. So, if you are under the care of the somebody and you're not necessarily able to control or express the same kind of warning signs that maybe everyone on the table would consider obvious today, wind the clock forward a few years. You're in a care home, okay, and you're starting to struggle with your breathing. Would somebody else spot that? How does somebody else baseline that?"

(Participant 8)

• Lifestyle issues in general were mentioned by one participant, diet was mentioned by another participant, and most the participants discussed smoking as being important to asthma, and discussed the likely underreporting of smoking levels to health care professionals.

4.4. What they would like to come out of the National Asthma Audit and how would they use the data

The importance of breakdown of the results by various patient characteristics was a common theme

• One participant strongly expressed the need for a breakdown of people with mild, moderate and severe asthma and to give a comparison with measures such as peak flow, hospital admissions and elements of care received.

"I would be really interested in seeing kind of a breakdown of the different categories, so mild, moderate and severe. And to see the variation that occurs within that. And then see well based on this is there anything we can break it down to better assist. And even if it's number based. So, a wider range of things and then maybe for your base level asthma. So how bad your asthma is in terms of how bad it can get and then how bad you are at the moment."

...

"But to make that kind of point and to change it, you need to have a look at the discrepancies within that scale itself. So, just looking at say, well these are the people who have said mild, here's their peak flows, sometimes they've been in hospital and then look at the variation across that. And it's gonna be massive I reckon. And then basically kind of have to providence or you know, lack thereof, for change in that system."

(Participant 9)

- Most the group discussed the importance of the audit showing the geographical variation with provision of "gold standard" care package down to county level and individual GP surgeries.
- Many of the respondents discussed the importance of separate reporting for underrepresented groups like children and older people with dementia.

"I think there's a huge amount of detail that can't possibly be included in a small leaflet to be given to patients. And if you are very involved then you're going to want to drill down a lot more. But actually national things give a really wide overview. And my feeling was that perhaps on this edition you'd drill down in to people ... they'd drill down to people with mental health or learning difficulties and that maybe the next one will drill down in to a different area."

(Participant 2)

Some of the participants highlighted areas that an overview would be useful in relation to statistics at a GP practice level. These areas are:

- Number/Percentage of patients who:
 - o Have asthma
 - o Have asthma and have been seen by their GP/Nurse
 - o Have asthma and have received each of the basic elements of care
 - o Have asthma and have attended A&E and/or been admitted to hospital
 - Have asthma and have been admitted to hospital for asthma within a month of attending their GP surgery to discuss their asthma
- How many languages
 - o Can asthma reviews be accommodated in/translated to?
 - o Do they have asthma action plans available in?

"I don't know how ... what detail the audit's gonna go in to. But you'd want to know how many patients have got asthma, have they seen those patients, have they done reviews, have they got plans in place for all of those patients. Cos, they may have lots of asthma patients that they've never actually seen in like five or six years. So, I don't know if you want to collect basic baseline information that would basically give you some idea of what's going on in that practice with regards to asthma care. And then the next level is, okay, when you see those patients what are you doing with them in the time that you see them."

(Participant 8)

"...And how many people go to their GP cos they're worried about their asthma, so have a doctor's appointment. But within a month are admitted to hospital for their asthma. That's gonna give us a fairly clear picture as well as to whether or not the GPs are being effective. And you could break that, then drill that kind of basic data down in to really localised areas. You can start with counties and then you can almost go down to practice level can't you. Surely that's got to be the way that we're actually identifying whether or not what we're doing is working. Cos I don't know that we know yet."

(Participant 2)

One participant strongly expressed the importance of routine care and asthma to save the NHS money:

"... Cos, if we were getting what we needed from our day to day practitioners we wouldn't even go to hospital."

(Participant 4)

Other topics identified as useful to come out of the National Asthma Audit

• One participant strongly expressed the view that the audit will give evidence on the importance of each element of care which may change thinking on what the "gold standard" package of care is: "So, surely the national audit's actually going to be identifying some of that, or it should be. So, does it make an impact if people do actually attend their six monthly reviews or don't. I mean, we assume it's a good thing. But actually is the evidence there that it is..."

(Participant 2)

- Some participants suggested the idea of showing the relevant asthma triggers by geographical location.
- One participant identified the importance of quantifying the gap between preventer inhaler usage and stated adherence particularly for teenagers.
- Some participants discussed a quantification of how far people with asthma are away from hospitals and where the nearest GP surgery with a nebuliser is.

How people with asthma will use the output from the audit

- Some of the group raised the use for informed patient choice between GP practices on where delivers the better asthma care.
- A few of the group discussed the ability to improve patient understanding on what to expect, including carers of people with mental health or learning conditions to ensure that they receive the care they need:

"If you've got a choice of GPs practice to have influence about how one handles it better than the other. I mean that's informed patient choice isn't it. Where you live in the country, how well they respond to asthma is informed patient choice. And so that if you are in the circumstances where you've only got one GPs practice within the locale, you know, then ... and if you can see that they're not particularly great at handling this then when you go for a visit, you are in a position to maybe ask them some questions about, am I not also, not gonna have, let's take that you know the seven things, and they're only giving you four. You can say, where are the other three things please to help me manage my condition."

(Participant 8)

• A small number of participants mentioned the use of the audit output to campaign to improve the local area like the inclusion of nebulisers in local GP surgeries. This included campaigns on behalf of

groups who are not receiving a high standard of care who are not able to campaign for themselves (including people with mental health or learning difficulties, or where English is not their first language):

"Yeah, we're assuming the data is going to start to give us information about what works and what doesn't work. So, then you can look at this place seems to have the gold standard and be most effective, what do we need to do here to get it up to that level. But some of it we do need to actually be quite sure whether it works or not."

(Participant 2)

• A small number of respondents also raised the use of the audit results to raise awareness of the seriousness of asthma in schools.

4.5. Patient level report

The example of the primary care section of a patient level report shown in Appendix 2 was shared with the group. This report section was produced by the AADP as a mockup of what the patient level report could include and look like, and was used to encourage feedback. Feedback was received in a number of areas:

Layout

- The participants found the draft report messy and difficult to follow.
 - o The switch between two and one columns amplified the difficulty to follow.
- One participant highlighted the difficulty to read black text on dark colours (referring to the pie chart).
- One participant expressed their like of the increased font size and emboldening of key points, but was less complementary of the colour changing:

"I think actually the size of the font and actually emboldening key points is really, really important just as your eye moves down the page. But then it's kind of like, I'm not sure about the colour changing..."

(Participant 4)

Charts

- The group were generally positive about the inclusion of charts in the report.
- The majority of participants favoured the bar charts due to their ease of understanding and accessibility.
- Some participants raised the importance of a consistent chart style and colour.
- One participant highlighted the need for the chart scale to go down to zero.
- One participant mentioned that progress overtime should be shown so that an assessment can be made as to whether this is a good or bad result.
- One participant suggested the inclusion of a Venn diagram to show the breakdown of respondent age.

Guidelines

- The inclusion of the guidelines was questioned by many of the participants, showing that the reason of including these needs to me made clear. It was suggested that the report following the guidelines should show whether the guidelines are being met or not, and whether there are any regional and socio-demographic disparities.
- Some members of the group questioned whether the guidelines go far enough (e.g. Annual review too far apart especially with the seasonality of asthma). One participant raised that it is important to explain that these are the current guidelines and unless we meet these there is less chance of increasing them.
- Some of the group questioned the meaning of the National Review of Asthma Deaths (NRAD) line stating that "People at high risk of severe asthma attacks should be monitored more closely, ensuring that their personal asthma action plan (PAAPs) are reviewed and updated at each review". The general consensus was that this is too vague and needs to be more specific to allow monitoring. "More closely, what does that mean? It should be more specific. It should be monitored whenever this happens or monitored at least guarterly or something like that."

(Participant 1)

Recommendations box

- Some of the participants raised that the recommendations box should include what you are going to do about the results and what you're going to look into next.
- One participant mentioned that this could be branded as an action rather than recommendation as this encourages the patient to go and do it.

Pictures

• The participants had mixed views on the pictures, with positive comments in relation to breaking up the text and negative comments in relation to being unnecessary and looking childish. Their use as bullet points in a poster was raised as an idea and also as different visual representations for groups to help identify the relevant bits (e.g. children, people with mental health conditions):

"I think the little logos are nice. Cos, if you use them consistently they will sort of help break up the stats and sort of take your ... people who don't want to read the stats can just read these sort of conclusions."

(Participant 1)

"That clipart thing there, I just don't think so. So, maybe if you use a series of other, kind of these ones like this, perhaps just like in poster bullet points perhaps. Just kind of bringing that up somewhere else. But simply putting in that one there to perhaps just show a review when it's just a check list just seems unnecessary and it seems just sort of childish and cartoonish..."

(Participant 9)

The group put forward a number of suggestions in relation to the report:

• The general consensus in the group was that the report needs to tell a story, starting by expressing the purpose of the report and what you want the patients to do with this information.

"I would have liked it to start off saying, people with asthma should have no symptoms and they should not be experiencing any symptoms because their medication is appropriate for them. And then these standards will achieve that. And then the measure of health whatever data it is you want to put up that measures whether or not you know those standards are leading ... being adhered to and leading to the fact that people with asthma are on the right medication and they're experiencing no symptoms and they're leading a full life."

(Participant 1)

"But the purpose of the audit is not only just to improve your guidelines but also to measure and improve GP practice, yes?"

(Participant 1)

- The need for a glossary of the terminology used like comorbidity, exacerbation, healthcare professional and any acronyms was discussed by some of the participants.
- One participant strongly expressed the need for the inclusion of the number of people the data represents, and another participant mentioned that the larger the number the stronger the campaigning power. When clarified, it was confirmed that the inclusion of the number at the start would be sufficient unless the numbers in each of the sections changed throughout.
- Some of the participants expressed the importance of demonstrating the representativeness of the sample by factors such as geography, ethnicity and age.
- There was consensus among the group that there should be specific sections for children, teenagers and elderly. One participant suggested that this be in the form of one page of infographics highlighting the areas of concern:

"Yeah, cos you'd want to know. So, 75% of the asthma patients had an annual review. Well who are those 75%? Is it elderly and children? Who's the 25%? Are they all teenagers?"

(Participant 1)

"Make sure to separate children and teenagers for difference because they're equally different"

(Participant 9)

 A few participants discussed the release of different report editions, each with a different focus, for example mental health. • Some of the group expressed the importance of the inclusion of breakdown by socio-economic group, ethnicity and first language to ensure engagement with different communities:

"Yeah, how good are they engaging with the different communities. Is there a gap? When you lump all of this data together, they say, yeah we engage with 80% of people. Okay, is that 80% of all people or is that 100% of the white middle class insistent English speaking community and only 40% of the Bangladeshi East End of London community and clearly then there's a gap. And that's unfortunately, that's where unmet need lies isn't it. You know when we merge these data sets together and we go, yeah 80% of us, it's not as bad as 75, is it. Whereas actually there's a whole bunch of people out there"

(Participant 8)

• One participant suggested that up and down arrows could be used to show what measures were positive and negative respectively.

4.6. COPD Infographic

The COPD infographic shown in Appendix 3 was shared with the group. This infographic was intended for a wider audience than asthma patients specifically, and was being shown to the group as an example of what could be produced from the National COPD Audit and could therefore be replicated for the National Asthma Audit. Feedback was received in a number of areas:

Positive comments

- The general opinion in the room was that the COPD infographic is attractive and that they would be drawn to read it.
- Some participants commented that they like how it tells a story.
- One participant mentioned that they like the lungs with the magnifying glass at the top.
- The general consensus in the group was that the four graphs at the top contained important information, and they liked how they were presented, particularly the use of the up and down arrows:

"The things I like to see are the four things up there which are all kind of the immediate things you're drawn to. You know, the infant mortality and stuff these are the biggest kind of selling points I think..."

(Participant 9)

Negative comments

• Other than the top four graphs the majority of participants expressed that the information included was not relevant to them as a patient:

"To be honest, the only thing that I'm interested in this top bit. The rest I kind of not that bothered to know about."

(Participant 1)

- One participant expressed concern that the most important points at the top are smaller than the less important points further down.
- A small number of participants raised the opinion that grey on grey doesn't work well.

The group put forward a number of suggestions in relation to the infographic:

- A small number of participants raised the importance of colour usage and how it could be used to link topics across the infographic thematically. Concern was raised by one participant that red is a danger colour and should be used sparingly to highlight important points; in this example people would be immediately drawn to the "reports" section due to the red colour.
- One participant mentioned that circular representation of the infographic could work.
- One participant mentioned that they would like to know how many practices were audited, but that no further methodological information was needed.
- Similarly as with the report, one participant raised the need for the infographic to tell a story, summarising what has and hasn't improved and what you want the patients to do with this information.
- The general consensus in the group was that the infographic pages should take two forms, one for patients which can be displayed in waiting rooms and one for doctors areas displaying information relevant to them:

"You need to have specific literature targeted at the patient cos then you could send it to every GP practice and the patient participation groups can then give it out to the patients. And make sure it's put out in the GP surgery."

(Participant 1)

• One participant expressed the need for child and teenager specific infographic pages for use on children's wards:

"Then even with targeting the patients as well, it's good to have a split maybe in general and then if you're going to have one in terms of areas, like the children's wards in hospitals. It's good to have one aimed at children and teenagers then. Cos that way if they go in and they see it there, it's a lot easier for the parents to make out, that actually this is the stuff that affects my child. And is probably likely to read it than just a general one with all this information down at the bottom or maybe a bit here, put a bit of children's stuff."

(Participant 9)

• When prompted, most the group said that they would like the same infographics to be used in both the patient level report and infographic page.

4.7. Gathering views and experiences of people with asthma

Although this section of the topic guide was not asked explicitly due to time constraints, the group raised suggestions for a number of questions which could be asked to people with asthma directly to assess if they were receiving good care:

To assess the quality of emergency care the following two questions were mentioned:

- Have you had an emergency episode where your initial treatment has not responded?
- How quickly were you progressed to the next stage of treatment?

To assess the quality of the asthma patients life the following two questions were raised:

- Have you taken antibiotics?
- Have you had a cold?

To assess the persons asthma and their resultant well-being, these three questions were mentioned:

- How is your asthma? On a 10 point scale instead of severe, moderate and mild.
- How is your asthma making you feel?
- How limiting is your asthma?

5. Summary and Conclusions

In summary and conclusion, the following eight key issues stand out:

1. The importance of looking at different groups in the population separately, particularly for children

This was a common theme throughout the group and almost all participants mentioned it's importance in some way. Groups mentioned as important to look at in isolation were children, teenagers, older people with dementia, people with mental health conditions and/or learning difficulties and people for whom English is not their first language.

2. The level of asthma care received in routine appointments is important

Most respondents raised the importance of their routine asthma care. Many respondents strongly expressed the importance of this care being done properly, particularly the asthma review and inhaler technique check.

3. The significance of GP level statistics for patients

This was mentioned by many participants as giving them informed patient choice and a basis to campaign for better care.

4. When receiving emergency care the correct asthma severity assessment and consequential treatment is of high importance

The participants discussed their stories of emergency care where experienced, which resulted in most participants highlighting the importance of correct severity assessment and consequentially the appropriate treatment.

5. The importance of trigger identification for people with asthma

Most participants raised the importance of trigger identification and its role within a routine appointment. Some participants suggested that triggers should be reported at a national level by geographic location.

6. Results of a future National Asthma Audit are useful for people with asthma

A variety of uses of the audit output were mentioned by a range of participants. Potential uses by people with asthma include informing patient choice, ensuring patients know what care they should receive and also to aid campaigning for better care.

7. Positive opinions on a patient focussed report with a clear purpose and meaningful breakdowns

The majority of participants expressed positivity towards a patient focussed report but stressed the need for it to have a clear purpose, with meaningful sections or editions for underrepresented groups like children and people with mental health conditions. Many of the respondents raised concern with some aspects of the design, emphasising the need for a good consistent design throughout.

8. Positive opinions on an infographic page relevant to the patient audience

Almost all participants gave a positive response to the COPD infographic page, but the importance of all information being relevant to the audience was strongly expressed.

Appendix 1: Topic guide

Asthma Audit Development Project

Focus group topic guide: 21st October 2017

Research Aims

To explore and understand:

- What people with asthma think is important to them about their asthma care
- What people with asthma think is important to record in an asthma audit
- Why a national asthma audit is important to people with asthma
- How people with asthma would like the draft patient level report to be
- Where to look to understand the views and experiences of people with asthma (time dependent)

Introduction

- Thank you for agreeing to take part
- Housekeeping: Fire exits and toilets
- Introduce Cat (Senior Insight Analyst) and Andrew (Policy Analyst)
- Introduction to Asthma UK (keep brief as all AUK volunteers)
- Explanation of research
 - o Royal College of Physicians
 - o Introduce Rachael (Project manager)
 - Asthma Audit
- Details of participation
 - Voluntary
 - Confidential
 - o Recording and data protection
 - o Quotes in report: permission form at the end (if not completed at the start)
 - No right or wrong answers
 - o May move topics on as have a lot to cover
 - o Length of group up to 1.5 hours
 - o Travel expenses will be processed at the end
 - o £30 thank you payment voucher to be provided at the end
- Questions?
- Start recording now

Participant background

- Ask each participant to introduce themselves, including:
 - o Their name
 - Where they live
 - o How long they (or their child if applicable) have had asthma
 - o Were they diagnosed with asthma as a child or adult
 - o How they would describe their asthma severity: Severe, Moderate, Mild
 - Where do they receive their day to day asthma care (e.g. GP surgery, specialist centre)

What is important to them about their asthma care/ What do they think is important to be recorded in an asthma audit

Views on what is important to them to be included in their routine care (doctors surgery or specialist centre)

- Annual review
 - Date of review
 - What the review included (items in blue below could be discussed in relation to the annual review or generally about their routine care)
- Medication
 - Inhaler
 - Type of inhaler(s)
 - How many reliever inhalers
 - How many preventer inhalers
 - Inhaler technique check
 - Inhaler dose
 - Preference for a metered dose inhaler
 - Spacer
 - Oral steroid tablets
 - Number of prescriptions
 - Doses
 - Co-morbidities
 - How medications work together
 - Side effects
- Asthma attacks
 - Number/frequency of asthma attacks
 - Hospital attendance due to asthma attack
 - GP follow up (within 2 days)
- Diagnosis
 - Correct
 - Up to date tools/guidance
- Clinical measures
- Other factors
 - Personalised asthma action plan
 - Triggers discussion
 - Referrals to secondary and tertiary care
 - Smoking cessation/medication
- Views on what is important to them in relation to emergency care when admitted to hospital
 - o **Arrival**
 - Arrival time at hospital
 - Treatment administered
 - Physical condition/Asthma severity
 - Pre-hospital treatment
 - Pre-hospital condition
 - During their stay
 - Treatment administered
 - Physical condition
 - Anything else important to their experience
 - Review and discharge
 - Time discharged from hospital (length of stay)
 - Inhaler technique check
 - Assessment of medication

- Assessment of adherence
- Personalised Asthma Action plan
- Triggers and exacerbating factors
- Smoking cessation
- Follow up appointments
- Prescription (e.g. Prednisolone)

What do they think is important for the doctor or nurse to know about the asthma patient to be able to assess the most appropriate care/

What do they think is important to be recorded in the asthma audit about the asthma patient to be able to assess the most appropriate care (*Emergency care or routine care*)

- Demographics
 - o Age
- Are they transitioning from paediatric to adult services
- Gender
- Other health conditions or disabilities
 - o Mental Health
 - Learning disability
- Smoking status
 - o Exposure to second hand smoke
- Asthma details
 - History
 - Medication

Why a national asthma audit is important to them

What does a national asthma audit mean to them

- What information would they like to come out of the audit
- How would they use the asthma audit output/information
 - o Their care
 - o Campaign for service improvement/development

What do they think of this patient level report

- Secondary care (hospital) section
 - Give them a quick minute to look through: If anything jumps out to them feel free to shout out
 - Layout
 - What do they think about the visualisations
 - Type
 - Stacked bar chart
 - o Donut
 - o Pie chart
 - o People size
 - Do they understand the data shown?
 - o Is explanation needed to help understand the charts
 - Would they prefer all the same or a mix of charts?
 - Would they prefer the data to be shown in other ways
 - What do they think about the pictures (e.g. Smoking, Brain)
 - What do you think about the text size and variations throughout?
 - o Content
 - What do they like

- What do they dislike
- Most important topic(s) to be included (doesn't have to be included in this example)
- How would you feel about including a section showing how many asthma patients admitted to hospital died from their asthma
- What do they think of the recommendation(s) box
- What do they think of the mental health/learning disabilities section
- Do they think an introduction is needed to the National Asthma Audit and terms used?
- How do they find the language it is written in
 - How could the language be improved
- Primary care (local doctor) section
 - Layout
 - o Give them a quick minute to look through: If anything jumps out to them feel free to shout out
 - What do they think about the visualisations
 - Type
 - o 3d pie chart
 - o Bar chart
 - Line chart
 - Do they understand the data shown
 - o Is explanation needed to help understand the charts
 - Would they prefer all the same or a mix of charts
 - Would they prefer the data to be shown in other ways
 - What do they think about the pictures (e.g. Brain, Upward arrow, Clipboard)
 - What do you think about the text size and variations throughout?
 - Content
 - What do they like
 - What do they dislike
 - Most important topic(s) to be included (doesn't have to be included in this example)
 - What do they think of the recommendation(s) box
 - What do they think of the standards/guidelines
 - What do they think of the mental health/learning disabilities section
 - Do they think an introduction is needed to the National Asthma Audit and terms used?
 - How do they find the language it is written in
 - How could the language be improved
- Infographic
 - Give them a quick minute to look through: If anything jumps out to them feel free to shout out
 - Would they find a similar thing useful
 - o What do they like
 - What do they dislike
 - o Most important topic(s) to be included

Where else to look to understand views and experiences of people with asthma (Time dependent)

- How do they feel about online surveys
 - How would they like to find out about a survey
 - o How do they think is a good way to get people with asthma to complete a survey
- Do they think that mental health conditions are important in an audit?
 - o How can we encourage people with mental health conditions to tell us about their care

- We want to ensure that we capture the views and experiences of people from different backgrounds
 - What organisations/community groups can we work with to help reach a diverse range of people

Rachael to ask any questions

Close

- Any other thoughts?
- Turn recording off
- Thank participants for their contributions
- Reaffirm confidentiality and anonymity
- Next steps for project
- How to contact research team (Cat Broadbent) in case of any queries or concerns
- Expenses and thank you payments

Appendix 2: Example of the primary care (local doctor) section which was reviewed in the focus group

Example of the primary care (local doctor) section

Annual asthma reviews by a local doctor

Asthma standard or guideline

NICE quality (QS 25) - statement 5: Review

People with asthma receive a structured review at least annually.

National Review of Asthma Deaths (NRAD

People with asthma should have a structured review by a healthcare professional with specialist training in asthma, at least annually. People at high risk of severe asthma attacks should be monitored more closely, ensuring that their personal asthma action plan (PAAPs) are reviewed and updated at each review.

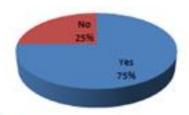
BTS/SIGN (153): British guideline on the management of asthma

In primary care, people with asthma should be reviewed regularly by a nurse or doctor with appropriate training in asthma management. Review should incorporate a written action plan.

Asthma standards and guidelines state that people with asthma should be have their asthma reviewed by their local doctor at least once a year.

Of the asthma patients the National Asthma Audit received information on from local doctors in Wales 75% had had an annual review of their asthma in the last 18 months.

Asthma patients who received an annual asthma review by their local doctor

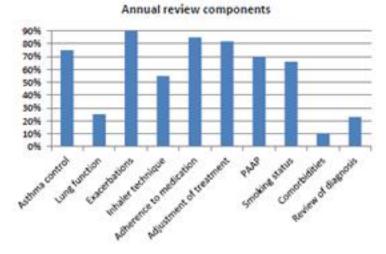


Annual asthma reviews should include ten key components.

The chart below shows you out of the 75% who had an annual review, what components were included in those reviews.



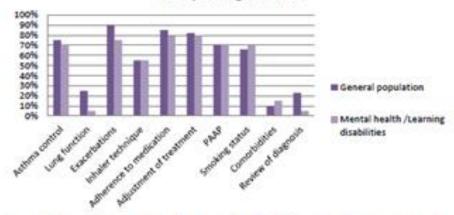
75% of asthma patients had a review of their asthma by a local doctor



Does care differ for people with mental health or learning disabilities?

The graph below shows the difference in the number of annual review components received by the general asthma population compared with those with a recorded mental health or learning disability.

General population vs asthma patients with mental health/learning disabilities

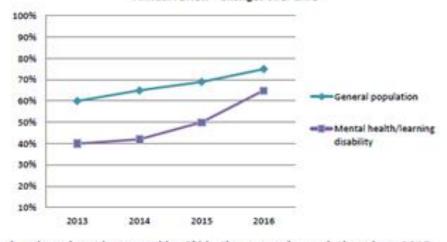


Asthma patients with a mental health or learning disability are less likely to receive a lung function test, a review of their exacerbations or an assessment of their adherence to asthma medication. They are more likely to have a review of their smoking status and comorbidities.

Changes over time

The number of annual asthma reviews, and individual review components being delivered within those reviews, have both increased since 2015.

Annual review - Changes over time



Annual reviews have increased by 6% in the general population since 2015

Recommendation(s)

This box will tell you what recommendation(s) the National Asthma Audit has produced specifically for local doctors and annual asthma reviews.

Appendix 3: National COPD Audit Programme Infographics















Section	Generic Questions					
Variable name	Age_Number	Age_Mean	Age_Stand deviation	Age_Median	Age_Lower Quartile	Age_Upper Quartile
Question number	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1
Variable	age_N	age_mean	age_sd	age_median	age_lq	age_uq
National results	292	42.8	17.7	41.0	28.0	54

Gender_Male_		Gender_Female_			Recorded Mental	Recorded Mental	Recorded Mental	Recorded Learning	Recorded Learning	Recorded Learning
Number	Gender_Male_%	Number	Gender_Female_%	Gender_Total	Health_Number	Health_%	Health_Total	DisNumber	Dis%	DisTotal
Q1.2	Q1.2	Q1.2	Q1.2	Q1.2	Q1.4	Q1.4	Q1.4	Q1.5	Q1.5	Q1.5
gendernumerator1	gendernumpc1	gendernumerator2	gendernumpc2	genderdenominator	q14numerator	q14numpc	q14denominator	q15numerator	q15numpc	q15denominator
74	25.3	218	74.7	292	61	20.9	292	4	1.4	292

				Smoking		Smoking status_									
Smoking status_Never_	Smoking	Smoking status_Ex_		status_Current_	Smoking	Current_Vaping_	Smoking status_	Smoking status_Not	Smoking status_Not		Exposure		Exposure		Exposure SHS_Not
Number	status_Never_%	Number	Smoking status_Ex_%	Number	status_Current_%	Number	Current_Vaping_%	recorded_Number	recorded_%	Smoking status_Total	SHS_No_Number	Exposure SHS_No_%	SHS_Yes_Number	Exposure SHS_Yes_%	recorded_Number
Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.7	Q1.7	Q1.7	Q1.7	Q1.7
q16numerator1	q16numpc1	q16numerator2	q16numpc2	q16numerator3	q16numpc3	q16numerator4	q16numpc4	q16numerator5	q16numpc5	q16denominator	q17numerator0	q17numpc0	q17numerator1	q17numpc1	q17numerator2
134	46.5	54	18.8	81	28.1		1 0.3	18	6.3	3 28	8 8	8 30.4	61	21.1	. 140

Appendix 27: AADP secondary care adult clincial pilot results

		Pre-hospital care								
Exposure SHS_Not recorded_%		Pre- hosp_steroids_Yes_ Number	Pre-hosp_steroids_ Yes_%	Pre-hosp_steroids_ Yes_Total						
Q1.7	Q1.7	Q2.1	Q2.1	Q2.1						
q17numpc2	q17denominator	q21anumerator	q21anumpc	q21adenominator						
48.4	289	102	36.3	781						

						Acute presentation	n and admission				
	Pre-hosp_β2		Pre-hosp_Oxygen_Yes_		Pre-			Heart rate_Standard			Heart rate_Upper
agonist_Yes_Number	agonist_Yes_%	agonist_Total	Number	hosp_Oxygen_Yes_%	hosp_Oxygen_Total	Heart rate_Number	Heart rate_Mean	deviation	Heart rate_Median	quartile	quartile
Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.1
q21bnumerator	q21bnumpc	q21bdenominator	q21cnumerator	q21cnumpc	q21cdenominator	q31_N	q31_mean	q31_sd	q31_median	q31_lq	q31_uq
192	67.4	285	80	29.3	273	287	104.7	18.4	104.0	93.0	118.0

		Resp rate_Standard		Resp rate_Lower	Resp rate_Upper			PEF arrival_Standard		PEF arrival_Lower	PEF arrival_Upper			PEF predicted_	
Resp rate_Number	Resp rate_Mean	deviation	Resp rate_Median	quartile	quartile	PEF arrival_Number	PEF arrival_Mean	deviation	PEF arrival_Median	quartile	quartile	PEF predicted_Number	PEF predicted_Mean	Standard deviation	PEF predicted_Median
Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3
q32_N	q32_mean	q32_sd	q32_median	q32_lq	q32_uq	q33a_N	q33a_mean	q33a_sd	q33a_median	q33a_lq	q33a_uq	q33b_N	q33b_mean	q33b_sd	q33b_median
2	88 24.4	6.6	23.0	20.0	27.5	215	224.5	90.8	210.0	160.0	280.0	87	431.1	92.	2 430.0

PEF predicted_Lower	PEF predicted_Upper			PEF Prev.B_Standard		PEF Prev.B_Lower	PEF Prev.B_Upper	PEF Not		PEF Not			Sp02_Standard		
quartile	quartile	PEF Prev.B_Number	PEF Prev.B_Mean	deviation	PEF Prev.B_Median	quartile	quartile	recorded_Number	PEF Not recorded_%	recorded_Total	Sp02_Number	Sp02_Mean	deviation	Sp02_Median	Sp02_Lower quartile
Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4
q33b_lq	q33b_uq	q33c_N	q33c_mean	q33c_sd	q33c_median	q33c_lq	q33c_uq	q33dnumerator	q33dnumpc	q33ddenominator	q34a_N	q34a_mean	q34a_sd	q34a_median	q34a_lq
360.0	0 450.0	89	365.4	104.6	370.0	300.0	420.0	67	22.9	292	288	95.4	4.	96	5.0 94.0

	Sp02_Not		Sp02_Not	Supp.Oxy_No, room	Supp.Oxy_No, room	Supp.Oxy_Yes_		Supp.Oxy_Not	Supp.Oxy_Not		Supp.Oxy_Flow	Supp.Oxy_Flow	Supp.Oxy_Flow rate_	Supp.Oxy_Flow	Supp.Oxy_Flow
Sp02_Upper quartile	recorded_Number	Sp02_Not recorded_%	recorded_Total	air_Number	air_%	Number	Supp.Oxy_Yes_%	recorded_Number	recorded_%	Supp.Oxy_Total	rate_Yes	rate_Mean	Standard deviation	rate_Median	rate_Lower quartile
Q3.4	Q3.4	Q3.4	Q3.4	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1
q34a_uq	q34bnumerator	q34bnumpc	q34bdenominator	q341anumerator0	q341anumpc0	q341anumerator1	q341anumpc1	q341anumerator2	q341anumpc2	q341adenominator	q341b_N	q341b_mean	q341b_sd	q341b_median	q341b_lq
98.	0 4	1.4	292	206	71.5	69	24.0	13	4.5	288	50	4.5	2.	6	4.0 2.0

Supp.Oxy_Flow				Supp.Oxy_Fi02_	Supp.Oxy_Fi02_		Oxygen presYes,				Oxygen presYes, not			Oxygen presNo,	Oxygen presNo,
rate_Upper quartile	Supp.Oxy_Fi02_Yes	Mean	Standard deviation	Median	Lower quartile	Upper quartile	admin_Number	admin_%	admin_Total	admin_Number	admin_%	admin_Total	admin_Number	admin_%	admin_Total
Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5
g341b_ug	q341c_N	q341c_mean	q341c_sd	q341c_median	q341c_lq	q341c_uq	q35anumerator	q35anumpc	q35adenominator	q35bnumerator	q35bnumpc	q35bdenominator	q35cnumerator	q35cnumpc	q35cdenominator
	6.0	60.0	36.8	37.5	26.0	100.0	95	32.5	292	45	15.4	292		22	7.5 292

					Time from arrival to										
			Time from arrival to	Time from arrival to	first administration of	first administration of	first administration of	first administration of	First admin	First admin	First admin			Time from arrival to	Time from arrival to
Oxygen presNo, not	Oxygen presNo, not	Oxygen presNo, not					steroids	steroids	steroids_Not	steroids_Not	steroids_Not	First admin steroids_	First admin	First admin β2 agonist	
admin_Number	admin_%	admin_Total	steroids_Number	steroids (Hours)_Mean	Standard deviation	(Hours)_ Median	(Hours) _Lower quartile	(Hours) _Upper quartile	administered_Number	administered_%	recorded_Number	Not recorded_%	steroids_Total	_Number	(Hours) _Mean
Q3.5	Q3.5	Q3.5	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.7	Q3.7
q35dnumerator	q35dnumpc	q35ddenominator	timtoster_N	timtoster_mean	timtoster_sd	timtoster_median	timtoster_lq	timtoster_uq	q36cnumerator1	q36cnumpc1	q36cnumerator2	q36cnumpc2	q36cdenominator	timtobeta_N	timtobeta_mean
8	8 30.1	292	205	3.8	10.7	1.5	0.5	3.6	8	3.4	23	9.7	237	190	1.6

									Length of stay						Discharge and review
Time from arrival to First															
admin β2 agonist	Time from arrival to	Time from arrival to	Time from arrival to	First admin β2	First admin β2	First admin β2									
(Hours)_	First admin β2 agonist	First admin β2 agonist	First admin B2 agonist	agonist_Not	agonist_Not	agonist_Not	First admin β2 agonist_	First admin β2			Length of stay_		Length of stay_Lower	Length of stay_Upper	Date & Time discharge_
Standard deviation	(Hours) _Median	(Hours) _Lower quartile	(Hours) _Upper quartile	administered_Number	administered_%	recorded_Number	Not recorded_%	agonist_Total	Length of stay_Number	Length of stay_Mean	Standard deviation	Length of stay_Median	quartile	quartile	Not yet discNumber
Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q4.1
timtobeta_sd	timtobeta_median	timtobeta_lq	timtobeta_uq	q37cnumerator1	q37cnumpc1	q37cnumerator2	q37cnumpc2	q37cdenominator	los_N	los_mean	los_sd	los_median	los_lq	los_uq	q41cnumerator
21.1	0.7	0.3	2.0		1.5	10	4.9	204	244	75.0	55.8	67.5	33.0	108.0	20

v															
Date & Time discharge_	Date & Time discharge_	Discharge	Discharge	Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_	Discharge bundle_Self-	Discharge bundle_Self-					DB_Med.classes_	
Not yet disc%	Not yet discTotal	bundle_No_Number	bundle_No_%	BTS_Number	BTS_%	non-BTS_Number	Yes, non-BTS_%	disNumber	dis%	Discharge bundle_Total	DB_Inhaler_Number	DB_Inhaler_%	DB_Inhaler_Total	Number	DB_Med.classes_%
Q4.1	Q4.1	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q41cnumpc	q41cdenominator	q42numerator0	q42numpc0	q42numerator1	q42numpc1	q42numerator2	q42numpc2	q42numerator3	q42numpc3	q42denominator	q421anumerator	q421anumpc	q421adenominator	q421bnumerator	q421bnumpc
6.9	9 290	108	37.5	75	26.0	98	34.0	7	2.4	288	188		71.5 263	1	184 70.5

DB_Med.classes_				DB_Import.Ad_			DB_PAAP_Modified_N		DB_PAAP_Modified_To	DB_ PAAP_Issued	DB_PAAP_Issued	DB_PAAP_Issued			
Total	DB_Doses_Number	DB_Doses_%	DB_Doses_Total	Number	DB_Import.Ad_%	DB_Import.Ad_Total	umber	DB_PAAP_Modified_%	tal	new_Number	new_%	new_Total	DB_Triggers_Number	DB_Triggers_%	DB_Triggers_Total
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q421bdenominator	q421cnumerator	q421cnumpc	q421cdenominator	q421dnumerator	q421dnumpc	q421ddenominator	q421enumerator	q421enumpc	q421edenominator	q421fnumerator	q421fnumpc	q421fdenominator	q421gnumerator	q421gnumpc	q421gdenominator
261	. 178	68.5	260	179	69.	257	55	28.2	195	:	137 5	9.3 231	178	69.5	256

DD Savelile -	DD Countries	DD Countries	DB_Follow-up_	DD Fallers on	DD 5-ll-	DB_Follow-up_	DD 5-11	DD Sellenner				Don't Marketon	Devel Marchine	Dood Marchine
DB_Smoking cessation_Number			Community_2 days_Number		DB_Follow-up_ Community_2 days_Total			DB_Follow-up_ Specialist_4 weeks_Total	DB_Other_Number	DB_Other_%		PredMgs/day_ Number		PredMgs/day_ Standard Deviation
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.3	Q4.3	Q4.3
q421hnumerator	q421hnumpc	q421hdenominator	q421inumerator	q421inumpc	q421idenominator	q421jnumerator	q421jnumpc	q421jdenominator	q421knumerator	q421knumpc	q421kdenominator	q43a_N	q43a_mean	q43a_sd
79	45.7	173	146	57.3	255	195	73.0	267	4	2 19.2	219	23	4 37.4	5.8

PredMgs/day_	PredMgs/day_	PredMgs/day_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo, completed	PredNo, completed	PredNo, completed			
Median	Lower quartile	Upper quartile	Number	Mean	Standard Deviation	Median	Lower quartile	Upper quartile	during hosp_Number	during hosp_%	during hosp_Total	PredNo_Number	PredNo_%	PredNo_Total
Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3
g43a_median	q43a_lq	q43a_uq	q43b_N	q43b_mean	q43b_sd	q43b_median	q43b_lq	q43b_uq	q43cnumerator	q43cnumpc	q43cdenominator	q43dnumerator	q43dnumpc	q43ddenominator
	40.0 40.0	40.0	225	6.2	5.0	5.0	4.	0 7.0	21	8.2	256	21	. 8	.2 256

Section	Admissions, staffir	ng levels and genera	l organisation of ca	ire		
Variable name	asthma	asthma admissions_2016_		asthma admissions_2016_	asthma admissions_2016_	Emergency coded asthma admissions_2016_ Upper quartile
Question number	Q1.3	Q1.3	Q1.3	Q1.3	Q1.3	Q1.3
Variable	q13_N	q13_mean	q13_sd	q13_median	q13_lq	q13_uq
National results	27	376.1	373.5	326	201	397

Emergency coded asthma admissions_discharged from resp. ward_Number	asthma admissions_ discharged from resp.	from resp. ward_	asthma admissions_ discharged from resp.	asthma admissions_ discharged from resp.	Emergency coded asthma admissions_ discharged from resp. ward_Upper quartile
Q1.4	Q1.4	Q1.4	Q1.4	Q1.4	Q1.4
q14_N	q14_mean	q14_sd	q14_median	q14_lq	q14_uq
25	89.1	58.1	81	43	11:

Organisation of ac	ute respiratory care												
Resp. Nurse			Resp. Nurse		Resp. Nurse	Resp. Nurse		Resp. Nurse				Resp. Nurse	
available_No_	Resp. Nurse	Resp. Nurse	available)_Monday_Nu	Resp. Nurse	available)_Monday_	available)_Tuesday_Nu	Resp. Nurse	available)_Tuesday_Tot	Resp. Nurse available)_	Resp. Nurse available)_	Resp. Nurse available)_	available)_Thursday_N	Resp. Nurse
Number	available)_No_%	available)_No_Total	mber	available)_Monday_%	Total	mber	available)_Tuesday_%	al	Wednesday_Number	Wednesday_%	Wednesday_Total	umber	available)_Thursday_%
Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8
q28anumerator	q28anumpc	q28adenominator	q28bnumerator	q28bnumpc	q28bdenominator	q28cnumerator	q28cnumpc	q28cdenominator	q28dnumerator	q28dnumpc	q28ddenominator	q28enumerator	q28enumpc
0	0	29	29	100	29	28	96.6	29	29	100	29	29	100

Resp. Nurse	Resp. Nurse		Resp. Nurse	Resp. Nurse		Resp. Nurse	Resp. Nurse		Resp. Nurse				
available)_Thursday_To				available)_Saturday_Nu		available)_Saturday_To			available)_Sunday_				If yes, responsible for
tal			Total		available)_Saturday_%					Asthma lead_Number	Asthma lead_%		training_Number
Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.8	Q2.10	Q2.10	Q2.10	Q2.10.1
q28edenominator	q28fnumerator	q28fnumpc	q28fdenominator	q28gnumerator	q28gnumpc	q28gdenominator	q28hnumerator	q28hnumpc	q28hdenominator	q210numerator	q210numpc	q210denominator	q2101numerator
29	9 28	96.6	29	9	31.0	29	7	24.1	29	24	82.8	29	14

		Integrating care	across primary and	secondary sectors									
If yes, responsibl	e for If yes, responsible for training_Total	Specialist Resp. Service_Severe r Asthma_Not provided_Number	Specialist Resp. Service_Severe Asthma_Not provided_%	Service_Severe	Service_Severe Asthma_Hosp.	Asthma_Comm.based_N	Specialist Resp. Service_Severe Asthma_Comm.based_%	Specialist Resp. Service_Severe Asthma_Single team_Number			MDT meetings_No_ Number	MDT meetings_No_%	MDT meetings_No_Total
Q2.10.1	Q2.10.1	Q5.2	Q5.2	Q5.2	Q5.2	Q5.2	Q5.2	Q5.2	Q5.2	Q5.2	Q5.5	Q5.5	Q5.5
q2101numpc	q2101denominator	q52anumerator0	q52anumpc0	q52anumerator1	q52anumpc1	q52anumerator2	q52anumpc2	q52anumerator3	q52anumpc3	q52adenominator	q55anumerator	q55anumpc	q55adenominator
	58.3	24	3 10).3 23	79.3	1	3.4	. 2	6.9	29	10	34.5	29

MDT meetings_Resp.	MDT meetings_Resp.
ConsNumber	Cons%
Q5.5	Q5.5
q55bnumerator	q55bnumpc
21	100

MDT meetings_Resp.	MDT meetings_Spec.	MDT meetings_Spec.	MDT meetings_Spec.	MDT meetings_GP_		MDT	MDT meetings_Hosp.	MDT meetings_Hosp.	MDT meetings_Hosp.
ConsTotal	Trainee_Number	Trainee_%	Trainee_Total	Number	MDT meetings_GP_%	meetings_GP_Total	nurse_Number	nurse_%	nurse_Total
Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5
q55bdenominator	q55cnumerator	q55cnumpc	q55cdenominator	q55dnumerator	q55dnumpc	q55ddenominator	q55enumerator	q55enumpc	q55edenominator
21	7	33.3	21	1	5	20	18	90	20

						MDT		MDT	MDT		MDT
MDT meetings_Comm.	MDT meetings_Comm.	MDT meetings_Comm.	MDT meetings_Physio_	MDT	MDT meetings_Physio_	meetings_Dietician_Nu	MDT	meetings_Dietician_Tot	meetings_Matron_	MDT	meetings_Matron_Tota
Nurse_Number	nurse_%	nurse_Total	Number	meetings_Physio_%	Total	mber	meetings_Dietician_%	al	Number	meetings_Matron_%	I
Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5
q55fnumerator	q55fnumpc	q55fdenominator	q55gnumerator	q55gnumpc	q55gdenominator	q55hnumerator	q55hnumpc	q55hdenominator	q55inumerator	q55inumpc	q55idenominator
3	15	20	9	45	20	2	10	20	1	5	20

MDT	MDT	MDT	MDT	MDT	MDT	MDT		MDT	MDT		MDT
			meetings_Thoracic			meetings_Psychologist_N		meetings_Psychologist_T	meetings_Radiologist_N	MDT	meetings_Radiologist_T
care_Number	care_%	care_Total	surgeon_Number	surgeon_%	surgeon_Total	umber	meetings_Psychologist_%	otal	umber	meetings_Radiologist_%	otal
Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5
q55jnumerator	q55jnumpc	q55jdenominator	q55knumerator	q55knumpc	q55kdenominator	q55Inumerator	q55Inumpc	q55ldenominator	q55mnumerator	q55mnumpc	q55mdenominator
	1 5	20	0	0	20	3	15	20	2	10	20

MDT		MDT				MDT		MDT		MDT		MDT
meetings_Pharmacist_N	MDT	meetings_Pharmacist_T	MDT meetings_Other_	MDT	MDT meetings_Other_	frequency_Weekly_	MDT	frequency_Fortnightly_N	MDT	frequency_Monthly_Nu	MDT	frequency_Quarterly_N
umber	meetings_Pharmacist_%	otal	Number	meetings_Other_%	Total	Number	frequency_Weekly_%	umber	frequency_Fortnightly_%	mber	frequency_Monthly_%	umber
Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5	Q5.5.1	Q5.5.1	Q5.5.1	Q5.5.1	Q5.5.1	Q5.5.1	Q5.5.1
q55nnumerator	q55nnumpc	q55ndenominator	q55onumerator	q55onumpc	q55odenominator	q551anumerator1	q551anumpc1	q551anumerator2	q551anumpc2	q551anumerator3	q551anumpc3	q551anumerator4
3	15	20	6	33.3	18	7	33.3	1	4.8	9	42.9	1

				Patient and carer	engagement				
						Formal survey seeking			
MDT	MDT frequency_Other_	MDT		patient/carer views_	patient/carer views_	patient/carer views_ >4	patient/carer views_ >4	patient/carer views_ 3-	patient/carer views_ 3-
frequency_Quarterly_%	Number	frequency_Other_%	MDT frequency_Total	Continuous_Number	Continuous_%	times a year_Number	times a years_%	4 times a year_Number	4 times a year_%
Q5.5.1	Q5.5.1	Q5.5.1	Q5.5.1	Q6.1	Q6.1	Q6.1	Q6.1	Q6.1	Q6.1
q551anumpc4	q551anumerator5	q551anumpc5	q551adenominator	q61numerator1	q61numpc1	q61numerator2	q61numpc2	q61numerator3	q61numpc3
4.8	3	14.3	21	0	0	1	3.5	1	3.5

		Formal survey seeking	Formal survey seeking										
Formal survey seeking	Formal survey seeking	patient/carer views_	patient/carer views_	Formal survey seeking	Formal survey seeking	Formal survey seeking					Strategic group for	Strategic group for	
patient/carer views_ 1-	patient/carer views_ 1-	Less than once a	Less than once a	patient/carer views_	patient/carer views_	patient/carer views_	Strategic group for resp.	Strategic group for	Strategic group for resp.	Strategic group for	resp. services_Not	resp. services_Not	Strategic group for
2 times a year_Number	2 times a year_%	year_Number	year_%	Never_Number	Never_%	Total	services_No_Number	resp. services_No_%	services_Yes_Number	resp. services_Yes_%	known_Number	known_%	resp. services_Total
Q6.1	Q6.1	Q6.1	Q6.1	Q6.1	Q6.1	Q6.1	Q6.2	Q6.2	Q6.2	Q6.2	Q6.2	Q6.2	Q6.2
q61numerator4	q61numpc4	q61numerator5	q61numpc5	q61numerator6	q61numpc6	q61denominator	q62numerator0	q62numpc0	q62numerator1	q62numpc1	q62numerator2	q62numpc2	q62denominator
3	10.3	9	31.0	15	51.7	29	5	17.2	21	72.4	3	10.3	29

Patient rep. on		Patient rep. on		Patient rep. on	Patient rep. on		Patient access to	Patient access to	Patient access to	Patient access to	Patient access to	Patient access to	Patient access to
strategic	Patient rep. on	strategic	Patient rep. on	strategic group_Not	strategic group_Not	Patient rep. on	healthcare	healthcare	healthcare	healthcare	healthcare records_Not	healthcare records_Not	healthcare
group_No_Number	strategic group_No_%	group_Yes_Number	strategic group_Yes_%	known_Number	known_%	strategic group_Total	records_No_Number	records_No_%	records_Yes_Number	records_Yes_%	known_Number	known_%	records_Total
Q6.2.1	Q6.2.1	Q6.2.1	Q6.2.1	Q6.2.1	Q6.2.1	Q6.2.1	Q6.3	Q6.3	Q6.3	Q6.3	Q6.3	Q6.3	Q6.3
q621numerator0	q621numpc0	q621numerator1	q621numpc1	q621numerator2	q621numpc2	q621denominator	q63numerator0	q63numpc0	q63numerator1	q63numpc1	q63numerator2	q63numpc2	q63denominator
13	61.9	6	28.6	2	9.5	21	10	34.5	18	62.1	1	3.5	29

							Transitional care		
Link with patient support/engagement group_No_Number	support/engagement	support/engagement	Link with patient support/engagement	group_Not	support/engagement		Transitional care_YP has full record_Number		Transitional care_YP has full record_Total
Q6.4	Q6.4	Q6.4	Q6.4	Q6.4	Q6.4	Q6.4	Q7.1	Q7.1	Q7.1
q64numerator0	q64numpc0	q64numerator1	q64numpc1	q64numerator2	q64numpc2	q64denominator	q71anumerator	q71anumpc	q71adenominator
11	37.9	18	62.1	0	0	29	14	50	

has same record as	has same record as	has same record as	has transitional care	has transitional care	has transitional care	Transitional care_YP has named case worker_Number	has named case	Transitional care_YP has named case worker_Total
Q7.1	Q7.1	Q7.1	Q7.1	Q7.1	Q7.1	Q7.1	Q7.1	Q7.1
q71bnumerator	q71bnumpc	q71bdenominator	q71cnumerator	q71cnumpc	q71cdenominator	q71dnumerator	q71dnumpc	q71ddenominator
14	53.8	26	11	42.3	26	6	24	25

Section	Generic questions					
Variable name	Age_Number	Age_Mean	Age_Stand deviation	Age_Median	Age_Lower Quartile	Age_Upper Quartile
Question number	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1
Variable	age_N	age_mean	age_sd	age_median	age_lq	age_uq
National results	263	2.7	1.3	3	2	4

Gender_Male_		Gender_Female_			Recorded Learning	Recorded Learning	Recorded Learning
Number	Gender_Male_%	Number	Gender_Female_%	Gender_Total	DisNumber	Dis%	DisTotal
Q1.2	Q1.2	Q1.2	Q1.2	Q1.2	Q1.4	Q1.4	Q1.4
gendernumerator1	gendernumpc1	gendernumerator2	gendernumpc2	genderdenominator	q14numerator	q14numpc	q14denominator
191	72.6	72	27.4	263	5	19	264

Exposure		Exposure		Exposure SHS_Not	Exposure SHS_Not	
SHS_No_Number	Exposure SHS_No_%	SHS_Yes_Number	Exposure SHS_Yes_%	recorded_Number	recorded_%	Exposure SHS_Total
Q1.5	Q1.5	Q1.5	Q1.5	Q1.5	Q1.5	Q1.5
q15numerator0	q15numpc0	q15numerator1	q15numpc1	q15numerator2	q15numpc2	q15denominator
123	46.6	37	14.0	104	39.4	264

Pre-hospital care								
Pre-						Pre-		
hosp_steroids_Yes_	Pre-hosp_steroids_	Pre-hosp_steroids_Yes	Pre-hosp_β2	Pre-hosp_β2	Pre-hosp_β2	hosp_Oxygen_Yes_	Pre-	Pre-
Number	Yes_%	Total	agonist_Yes_Number	agonist_Yes_%	agonist_Total	Number	hosp_Oxygen_Yes_%	hosp_Oxygen_Total
Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1
q21anumerator	q21anumpc	q21adenominator	q21bnumerator	q21bnumpc	q21bdenominator	q21cnumerator	q21cnumpc	q21cdenominator
30	15.6	250	181	69.3	261	23	9.3	248

Acute presentation	n and admission							
		Heart rate_Standard		Heart rate_Lower	Heart rate_Upper			Resp rate_Standard
Heart rate_Number	Heart rate_Mean	deviation	Heart rate_Median	quartile	quartile	Resp rate_Number	Resp rate_Mean	deviation
Q3.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.2	Q3.2	Q3.2
q31_N	q31_mean	q31_sd	q31_median	q31_lq	q31_uq	q32_N	q32_mean	q32_sd
263	145.9	19.4	146	134	154	264	42.5	9.5

	Resp rate_Lower	Resp rate_Upper			Sp02_Standard				Sp02_Not		Sp02_Not
Resp rate_Median	quartile	quartile	Sp02_Number	Sp02_Mean	deviation	Sp02_Median	Sp02_Lower quartile	Sp02_Upper quartile	recorded_Number	Sp02_Not recorded_%	recorded_Total
Q3.2	Q3.2	Q3.2	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3
q32_median	q32_lq	q32_uq	q33a_N	q33a_mean	q33a_sd	q33a_median	q33a_lq	q33a_uq	q33bnumerator	q33bnumpc	q33bdenominator
41.5	36	48	264	94.6	3.4	95	93	97	0	0	264

Supp.Oxy_No, room	Supp.Oxy_No, room	Supp.Oxy_Yes_			Supp.Oxy_Not						Supp.Oxy_Flow	Supp.Oxy_Flow
air_Number	air_%	Number	Supp.Oxy_Yes_%	recorded_Number	recorded_%	Supp.Oxy_Total	rate_Yes	rate_Mean	Standard deviation	rate_Median	rate_Lower quartile	rate_Upper quartile
Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1
q331anumerator0	q331anumpc0	q331anumerator1	q331anumpc1	q331anumerator2	q331anumpc2	q331adenominator	q331b_N	q331b_mean	q331b_sd	q331b_median	q331b_lq	q331b_uq
252	95.5	9	3.4	3	1.1	264	15	7.1	3.9	6	4	10

	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Oxygen presYes,	Oxygen presYes,	Oxygen presYes,	Oxygen presYes, not	Oxygen pres. Yes, not	Oxygen pres. Yes, not	Oxygen pres. No,	Oxygen presNo,	Oxygen presNo,
Supp.Oxy_Fi02_Yes		Standard deviation	Median	Lower quartile	Upper quartile		admin_%					admin_Number		admin_Total
Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.3.1	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4
q331c_N	q331c_mean	q331c_sd	q331c_median	q331c_lq	q331c_uq	q34anumerator	q34anumpc	q34adenominator	q34bnumerator	q34bnumpc	q34bdenominator	q34cnumerator	q34cnumpc	q34cdenominator
2	2 60	45.3	60	2	8 9	2 37	14.0	264	4	1.5	264	56	21.2	264

		Í							
Oxygen presNo, not	Oxygen presNo, not	Oxygen presNo, not	Steroids		Steroids		Steroids adminNot	Steroids adminNot	
admin_Number	admin_%	admin_Total	adminNo_Number	Steroids adminNo_%	adminYes_Number	Steroids adminYes_%	recorded_Number	recorded_%	Steroids adminTotal
Q3.4	Q3.4	Q3.4	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5
q34dnumerator	q34dnumpc	q34ddenominator	q35numerator0	q35numpc0	q35numerator1	q35numpc1	q35numerator2	q35numpc2	q35denominator
148	56.1	264	57	22.2	198	77.0	2	0.8	257

		Time from arrival to		Time from arrival to	Time from arrival to					
Time from arrival to	Time from arrival to	First admin β2 agonist	Time from arrival to	First admin β2 agonist	First admin β2 agonist	First admin β2	First admin β2	First admin β2	First admin β2	
First admin β2 agonist	First admin β2 agonist	(Hours)_	First admin β2 agonist	(Hours)_Lower	(Hours) _Upper	agonist_Not	agonist_Not	agonist_Not	agonist_	First admin β2
_Number	(Hours)_Mean	Standard deviation	(Hours)_Median	quartile	quartile	administered_Number	administered_%	recorded_Number	Not recorded_%	agonist_Total
Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6
timtobeta_N	timtobeta_mean	timtobeta_sd	timtobeta_median	timtobeta_lq	timtobeta_uq	q36cnumerator1	q36cnumpc1	q36cnumerator2	q36cnumpc2	q36cdenominator
172	2.3	13.2	0.725	0.3	1.6	2	1.1	6	3.3	18

Length of stay						Review and dischar	ge			
		Length of stay		Length of stav	Length of stay		Date & Time	Date & Time		
Length of stay	Length of stay	(Hours)_	Length of stay			Date & Time discharge_	discharge_	discharge_	Discharge	Discharge
(Hours)_Number	(Hours) _Mean	Standard deviation	(Hours) _Median	quartile	quartile	Not yet discNumber	Not yet disc%	Not yet discTotal	bundle_No_Number	bundle_No_%
Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q4.1	Q4.1	Q4.1	Q4.2	Q4.2
los_N	los_mean	los_sd	los_median	los_lq	los_uq	q41cnumerator	q41cnumpc	q41cdenominator	q42numerator0	q42numpc0
236	24.1	22.4	20.6	13.525	31.0	21	8.0	262	119	46.7

				Discharge	Discharge							
Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_	bundle_Parental-	bundle_Parental-	Discharge				DB_Med.classes_		DB_Med.classes_
BTS_Number	BTS_%	non-BTS_Number	Yes, non-BTS_%	disNumber	dis%	bundle_Total	DB_Inhaler_Number	DB_Inhaler_%	DB_Inhaler_Total	Number	DB_Med.classes_%	Total
Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2						
q42numerator1	q42numpc1	q42numerator2	q42numpc2	q42numerator3	q42numpc3	q42denominator	q421anumerator	q421anumpc	q421adenominator	q421bnumerator	q421bnumpc	q421bdenominator
21	8.2	115	45.1	0	0	255	170	70.5	241	153	66.	2 231

				DB_Import.Ad_			DB_PAAP_Modified_N		DB_PAAP_Modified_T		DB_PAAP_Issued				
DB_Doses_Number	DB_Doses_%		DB_Doses_Total	Number	DB_Import.Ad_%	DB_Import.Ad_Total	umber	DB_PAAP_Modified_%	otal	new_Number	new_%	new_Total	DB_Triggers_Number	DB_Triggers_%	DB_Triggers_Total
q421cnumerator	q421cnumpc		q421cdenominator	q421dnumerator	q421dnumpc	q421ddenominator	q421enumerator	q421enumpc	q421edenominator	q421fnumerator	q421fnumpc	q421fdenominator	q421gnumerator	q421gnumpc	q421gdenominator
14	13	64.1	22:	3 75	34.2	219	11	5.4	203	98		43.75 224	8:	1	34.8 233

DB_Follow-up_		DB_Follow-up_	DB_Follow-up_		DB_Follow-up_						
	DB_Follow-up_	Community_2	Paed.clinic_4	DB_Follow-up_	Paed.clinic_4	DB_Follow-up_	DB_Follow-up_	DB_Follow-up_			
days_Number	Community_2 days_%	days_Total	weeks_Number	Paed.clinic_4 weeks_%	weeks_Total	Paed.specNumber	Paed.spec%	Paed.specTotal	DB_Other_Number	DB_Other_%	DB_Other_Total
q421inumerator	q421inumpc	q421idenominator	q421jnumerator	q421jnumpc	q421jdenominator	q421knumerator	q421knumpc	q421kdenominator	q421Inumerator	q421Inumpc	q421Idenominator
97	41.6	233	43	19.4	222	13	5.8	224	32	14.2	226

				PredNo, completed	PredNo, completed	
PredNo_Number	PredNo_%	PredYes_Number	PredYes_%	during hosp_Number	during hosp_%	PredTotal
q43numerator0	q43numpc0	q43numerator1	q43numpc1	q43numerator2	q43numpc2	q43denominator
62	25	17/	70.2	12	18	

Section	Generic questions					
Variable name	Age_Number	Age_Mean	Age_Stand deviation	Age_Median	Age_Lower Quartile	Age_Upper Quartile
Question number	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1
Variable	age_N	age_mean	age_sd	age_median	age_lq	age_uq
National results	144	9.0	2.8	9	7	11

Gender_Male_		Gender_Female_			CAMHS		
Number	Gender_Male_%	Number	Gender_Female_%	Gender_Total	referral_Number	CAMHS referral_%	CAMHS referral_Total
Q1.2	Q1.2	Q1.2	Q1.2	Q1.2	Q1.4	Q1.4	Q1.4
gendernumerator1	gendernumpc1	gendernumerator2	gendernumpc2	genderdenominator	q14numerator	q14numpc	q14denominator
86	59.7	58	40.3	144	2	1.4	139

Recorded Learning	Recorded Learning	Recorded Learning	Exposure		Exposure		Exposure SHS_Not	Exposure SHS_Not	
DisNumber	Dis%	DisTotal	SHS_No_Number	Exposure SHS_No_%	SHS_Yes_Number	Exposure SHS_Yes_%	recorded_Number	recorded_%	Exposure SHS_Total
Q1.5	Q1.5	Q1.5	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6
q15numerator	q15numpc	q15denominator	q16numerator0	q16numpc0	q16numerator1	q16numpc1	q16numerator2	q16numpc2	q16denominator
3	2.2	139	53	37.1	26	18.2	64	44.8	143

Pre-hospital care								
Pre-								
hosp_steroids_Yes_	Pre-hosp_steroids_	Pre-hosp_steroids_	Pre-hosp_β2	Pre-hosp_β2	Pre-hosp_β2	Pre-hosp_Oxygen_Yes_	Pre-	Pre-
Number	Yes_%	Yes_Total	agonist_Yes_Number	agonist_Yes_%	agonist_Total	Number	hosp_Oxygen_Yes_%	hosp_Oxygen_Total
Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1
q21anumerator	q21anumpc	q21adenominator	q21bnumerator	q21bnumpc	q21bdenominator	q21cnumerator	q21cnumpc	q21cdenominator
35	24.5	143	108	75.5	143	10	7.1	141

Acute presentatio	Acute presentation and admission												
		Heart rate_Standard		Heart rate_Lower	Heart rate_Upper			Resp rate_Standard		Resp rate_Lower	Resp rate_Upper		
Heart rate_Number	Heart rate_Mean	deviation	Heart rate_Median	quartile	quartile	Resp rate_Number	Resp rate_Mean	deviation	Resp rate_Median	quartile	quartile		
Q3.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.1	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2		
q31_N	q31_mean	q31_sd	q31_median	q31_lq	q31_uq	q32_N	q32_mean	q32_sd	q32_median	q32_lq	q32_uq		
143	125.0	18.4	125	113	140	143	31.0	8.7	28	24	36		

		PEF arrival_Standard		PEF arrival_Lower	PEF arrival_Upper			PEF predicted_		PEF predicted_Lower	PEF predicted_Upper
PEF arrival_Number	PEF arrival_Mean	deviation	PEF arrival_Median	quartile	quartile	PEF predicted_Number	PEF predicted_Mean	Standard deviation	PEF predicted_Median	quartile	quartile
Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3
q33a_N	q33a_mean	q33a_sd	q33a_median	q33a_lq	q33a_uq	q33b_N	q33b_mean	q33b_sd	q33b_median	q33b_lq	q33b_uq
17	179.7	65.6	180	150	220	10	277.2	100.1	297	200	358

		PEF Prev.B_Standard		PEF Prev.B_Lower	PEF Prev.B_Upper	PEF Not		PEF Not
PEF Prev.B_Number	PEF Prev.B_Mean	deviation	PEF Prev.B_Median	quartile	quartile	recorded_Number	PEF Not recorded_%	recorded_Total
Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3	Q3.3
q33c_N	q33c_mean	q33c_sd	q33c_median	q33c_lq	q33c_uq	q33dnumerator	q33dnumpc	q33ddenominator
3	323.3	30.6	330	290	350	125	87.4	143

		Sp02_Standard				Sp02_Not		Sp02_Not	Supp.Oxy_No, room	Supp.Oxy_No, room
Sp02_Number	Sp02_Mean	deviation	Sp02_Median	Sp02_Lower quartile	Sp02_Upper quartile	recorded_Number	Sp02_Not recorded_%	recorded_Total	air_Number	air_%
Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4	Q3.4.1	Q3.4.1
q34a_N	q34a_mean	q34a_sd	q34a_median	q34a_lq	q34a_uq	q34bnumerator	q34bnumpc	q34bdenominator	q341anumerator0	q341anumpc0
	141 9	14.5	95	93	97	3	2.1	144	128	90.1

Supp.Oxy_Yes_		Supp.Oxy_Not	Supp.Oxy_Not		Supp.Oxy_Flow	Supp.Oxy_Flow	Supp.Oxy_Flow rate_	Supp.Oxy_Flow	Supp.Oxy_Flow	Supp.Oxy_Flow
Number	Supp.Oxy_Yes_%	recorded_Number	recorded_%	Supp.Oxy_Total	rate_Yes	rate_Mean	Standard deviation	rate_Median	rate_Lower quartile	rate_Upper quartile
Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1
q341anumerator1	q341anumpc1	q341anumerator2	q341anumpc2	q341adenominator	q341b_N	q341b_mean	q341b_sd	q341b_median	q341b_lq	q341b_uq
11	7.7	3	2.1	142	12	7.2	4.4	5.5	4.5	10

	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Supp.Oxy_Fi02_	Oxygen presYes,	Oxygen presYes,	Oxygen presYes,	Oxygen presYes, not	Oxygen presYes, not	Oxygen presYes, not
Supp.Oxy_Fi02_Yes	Mean	Standard deviation	Median	Lower quartile	Upper quartile	admin_Number	admin_%	admin_Total	admin_Number	admin_%	admin_Total
Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.4.1	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5
q341c_N	q341c_mean	q341c_sd	q341c_median	q341c_lq	q341c_uq	q35anumerator	q35anumpc	q35adenominator	q35bnumerator	q35bnumpc	q35bdenominator
1	85		85	85	85	15	10.4	144	1	0.7	144

								Time from arrival to	Time from arrival to	Time from arrival to	Time from arrival to
						Time from arrival to					first administration of
Oxygen presNo,	Oxygen presNo,	Oxygen presNo,	Oxygen presNo, not	Oxygen presNo, not	Oxygen presNo, not	first administration of	first administration of	steroids (Hours)_	steroids	steroids	steroids
admin_Number	admin_%	admin_Total	admin_Number	admin_%	admin_Total	steroids_Number	steroids (Hours) _Mean	Standard deviation	(Hours) _Median	(Hours) _Lower quartile	(Hours) _Upper quartile
Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.6
q35cnumerator	q35cnumpc	q35cdenominator	q35dnumerator	q35dnumpc	q35ddenominator	timtoster_N	timtoster_mean	timtoster_sd	timtoster_median	timtoster_lq	timtoster_uq
37	25.7	144	71	49.3	144	118	1.2	5.7	1.0	0.45	2

							Time from arrival to				
first administration of	first administration of	first administration of	first administration of		Time from arrival to	Time from arrival to	First admin β2 agonist	Time from arrival to	Time from arrival to	Time from arrival to	First admin β2
steroids_Not	steroids_Not	steroids_Not	steroids_Not	first administration of	First admin β2	First admin β2 agonist	(Hours)_	First admin β2 agonist	First admin β2 agonist	First admin β2 agonist	agonist_Not
administered_Number	administered_%	recorded_Number	recorded_%	steroids_Total	agonist_Number	(Hours) _Mean	Standard deviation	(Hours) _Median	(Hours) _Lower quartile	(Hours) _Upper quartile	administered_Number
Q3.6	Q3.6	Q3.6	Q3.6	Q3.6	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7	Q3.7
q36cnumerator1	q36cnumpc1	q36cnumerator2	q36cnumpc2	q36cdenominator	timtobeta_N	timtobeta_mean	timtobeta_sd	timtobeta_median	timtobeta_lq	timtobeta_uq	q37cnumerator1
5	3.9	4	3.1	128	92	0.4	3.4	0.5	0.2	0.9	1

				Length of stay					
First admin β2	First admin β2	First admin β2				Length of stay			
agonist_Not	agonist_Not	agonist_Not	First admin β2		Length of stay	(Hours)_	Length of stay	Length of stay	Length of stay
administered_%	recorded_Number	recorded_%	agonist_Total	Length of stay_Number	(Hours) _Mean	Standard deviation	(Hours) _Median	(Hours) _Lower quartile	(Hours) _Upper quartile
Q3.7	Q3.7	Q3.7	Q3.7	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1	Q1.3, Q4.1
q37cnumpc1	q37cnumerator2	q37cnumpc2	q37cdenominator	los_N	los_mean	los_sd	los_median	los_lq	los_uq
1.0	2	2.1	96	131	34.6	29.3	24.8	13.9	45.7

Review and discha	ırge										
Date & Time	Date & Time	Date & Time							Discharge	Discharge	
discharge_	discharge_	discharge_	Discharge	Discharge	Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_Yes,	Discharge bundle_	bundle_Self/Parental-	bundle_Self/Parental-	Discharge
Not yet discNumber	Not yet disc%	Not yet discTotal	bundle_No_Number	bundle_No_%	BTS_Number	BTS_%	non-BTS_Number	Yes, non-BTS_%	disNumber	dis%	bundle_Total
Q4.1	Q4.1	Q4.1	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2
q41cnumerator	q41cnumpc	q41cdenominator	q42numerator0	q42numpc0	q42numerator1	q42numpc1	q42numerator2	q42numpc2	q42numerator3	q42numpc3	q42denominator
11	7.6	144	58	41.7	25	18.0	56	40.3	0	0	139

			DB_Med.classes_		DB_Med.classes_				DB_Import.Ad_		
DB_Inhaler_Number	DB_Inhaler_%	DB_Inhaler_Total	Number	DB_Med.classes_%	Total	DB_Doses_Number	DB_Doses_%	DB_Doses_Total	Number	DB_Import.Ad_%	DB_Import.Ad_Total
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q421anumerator	q421anumpc	q421adenominator	q421bnumerator	q421bnumpc	q421bdenominator	q421cnumerator	q421cnumpc	q421cdenominator	q421dnumerator	q421dnumpc	q421ddenominator
106	80.9	131	90	70.9	127	80	63.5	126	67	53.2	126

DB_PAAP_Modified_N		DB_PAAP_Modified_To	DB_PAAP_Issued	DB_PAAP_Issued	DB_PAAP_Issued				DB_Smoking	DB_Smoking	DB_Smoking
umber	DB_PAAP_Modified_%	tal	new_Number	new_%	new_Total	DB_Triggers_Number	DB_Triggers_%	DB_Triggers_Total	cessation_Number	cessation_%	cessation_Total
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q421enumerator	q421enumpc	q421edenominator	q421fnumerator	q421fnumpc	q421fdenominator	q421gnumerator	q421gnumpc	q421gdenominator	q421hnumerator	q421hnumpc	q421hdenominator
24	22.6	106	47	41.6	113	59	46.5	127	7	7.3	96

DB_Follow-up_		DB_Follow-up_	DB_Follow-up_		DB_Follow-up_						
						DB_Follow-up_	DB_Follow-up_	DB_Follow-up_			
days_Number	Community_2 days_%	days_Total	weeks_Number	Paed.clinic_4 weeks_%	weeks_Total	Paed.specNumber	Paed.spec%	Paed.specTotal	DB_Other_Number	DB_Other_%	DB_Other_Total
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q421inumerator	q421inumpc	q421idenominator	q421jnumerator	q421jnumpc	q421jdenominator	q421knumerator	q421knumpc	q421kdenominator	q421Inumerator	q421Inumpc	q421Idenominator
39	30.2	129	29	23.4	124	32	26.4	121	22	18.6	118

PredMgs/kgs per	PredMgs/kgs per	PredMgs/kgs per	Pred. Mgs/kgs per	PredMgs/kgs per	PredMgs/kgs per						
I .	l .					PredMgs/day_	Pred. Mgs/day	PredMgs/day_	PredMgs/day_	PredMgs/day_	PredMgs/day_
					Upper quartile			Standard Deviation			Upper quartile
Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3
q43a_N	q43a_mean	q43a_sd	q43a_median	q43a_lq	q43a_uq	q43b_N	q43b_mean	q43b_sd	q43b_median	q43b_lq	q43b_uq
16	1.45	0.6		1	2	100			40	30	40

PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo.Days_	PredNo, completed	PredNo, completed	PredNo, completed			
Number	Mean	Standard Deviation	Median	Lower quartile	Upper quartile	during hosp_Number	during hosp_%	during hosp_Total	PredNo_Number	PredNo_%	PredNo_Total
Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3	Q4.3
q43c_N	q43c_mean	q43c_sd	q43c_median	q43c_lq	q43c_uq	q43dnumerator	q43dnumpc	q43ddenominator	q43enumerator	q43enumpc	q43edenominator
93	2.3	1.6	2	1	3	23	18.5	124	18	15.1	119

Section	Admissions, staffin	g levels and genera	al organisation of c	are							
	,	8	8								1
							Emergency paediatric	Emergency paediatric	Emergency paediatric	Emergency naediatric	Emergency paediatric
	Paediatric emergency	resporatory coded		resporatory coded	resporatory coded	resporatory coded					
	admissions_2016_										
Variable name	Number	Mean	Standard deviation	Median	Lower quartile	Upper quartile	Number	Mean	Standard deviation	Median	Lower quartile
Question number	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1	Q1.1	Q1.2	Q1.2	Q1.2	Q1.2	Q1.2
Variable	q11_N	q11_mean	q11_sd	q11_median	q11_lq	q11_uq	q12_N	q12_mean	q12_sd	q12_median	q12_lq
National results	20	6209.0	5316.0	4990	3312	6781	20	1465	1002.9	1199.5	756

Emergency paediatric resporatory coded admissions_2016_ Upper quartile
Q1.2
q12_uq

Paediatric emergency coded asthma admissions_2016_ Number	Paediatric emergency coded asthma admissions_2016_ Mean	coded asthma admissions_2016_			Paediatric emergency coded asthma admissions_2016_ Upper quartile
Q1.3	Q1.3	Q1.3	Q1.3	Q1.3	Q1.3
q13_N	q13_mean	q13_sd	q13_median	q13_lq	q13_uq
18	352.8	425.7	181 5	141	318

							Emergency paediatric	Emergency paediatric
Emergency paediatric	coded asthma	coded asthma	coded asthma					
coded asthma	admissions_	admissions_	admissions_					
admissions_discharged	admissions_	admissions_discharged	admissions_	admissions_	admissions_	discharged from resp.	discharged from resp.	discharged from resp.
from resp.	discharged from resp.	from resp. ward_	discharged from resp.	discharged from resp.	discharged from resp.	ward_No dedicated	ward_No dedicated	ward_No dedicated
ward_Number	ward_Mean	Standard deviation	ward_Median	ward_Lower quartile	ward_Upper quartile	resp. ward_Number	resp. ward_%	resp. ward_Total
Q1.4								
q14a_N	q14a_mean	q14a_sd	q14a_median	q14a_lq	q14a_uq	q14bnumerator	q14bnumpc	q14bdenominator
5	0.2	0.4	0	0	0	19	100	19

		Medical beds used for		Medical beds used for	Medical beds used for						
Medical beds used for	Medical beds used for	paediatric	Medical beds used for	paediatric	paediatric						
paediatric	paediatric	patients_Standard	paediatric	patients_Lower	patients_Upper	Paediatric HDUs_No	Paediatric HDUs_No	Paediatric HDUs_No	Paediatric HDUs_Paed.	Paediatric HDUs_Paed.	Paediatric HDUs_Paed.
patients_Number	patients_Mean	deviation	patients_Median	quartile	quartile	HDU_Number	HDU_%	HDU_Total	HDU_Number	HDU_%	HDU_Total
Q1.5	Q1.5	Q1.5	Q1.5	Q1.5	Q1.5	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6
q15_N	q15_mean	q15_sd	q15_median	q15_lq	q15_uq	q16anumerator	q16anumpc	q16adenominator	q16bnumerator	q16bnumpc	q16bdenominator
21	34.9	27.9	28	22	34	. 6	30	20	16	72.7	22

			Paediatric								
			HDUs_Mixed	Paediatric	Paediatric						
Paediatric HDUs_Med.	Paediatric HDUs_Med.	Paediatric HDUs_Med.	med./surg	HDUs_Mixed	HDUs_Mixed	Paediatric HDUs_Resp.	Paediatric HDUs_Resp.	Paediatric HDUs_Resp.	Paediatric HDUs_Other	Paediatric HDUs_Other	Paediatric HDUs_Other
HDU_Number	HDU_%	HDU_Total	HDU_Number	med./surg HDU_%	med./surg HDU_Total	HDU_Number	med./surg HDU_%	med./surg HDU_Total	HDU_Number	med./surg HDU_%	med./surg HDU_Total
Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6	Q1.6
q16cnumerator	q16cnumpc	q16cdenominator	q16dnumerator	q16dnumpc	q16ddenominator	q16enumerator	q16enumpc	q16edenominator	q16fnumerator	q16fnumpc	q16fdenominator
5	25	20	8	38.1	21	0	0	19	1	5	20

| Paediatric HDU |
|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|----------------|
| beds_0_Number | beds_0_% | beds_1_Number | beds_1_% | beds_2_Number | beds_2_% | beds_3_Number | beds_3_% | beds_4_Number | beds_4_% | beds_5_Number | beds_5_% |
| Q1.7 |
| q17numerator0 | q17numpc0 | q17numerator1 | q17numpc1 | q17numerator2 | q17numpc2 | q17numerator3 | q17numpc3 | q17numerator4 | q17numpc4 | q17numerator5 | q17numpc5 |
| 3 | 13.0 | 3 | 13.0 | 3 | 13.0 | 2 | 8.7 | 6 | 26.1 | 2 | 8.7 |

Paediatric HDU	Paediatric HDU										
beds_6_Number	beds_6_%	beds_7_Number	beds_7_%	beds_8_Number	beds_8_%	beds_9_Number	beds_9_%	beds_10_Number	beds_10_%	beds_>10_Number	beds_>10_%
Q1.7	Q1.7										
q17numerator6	q17numpc6	q17numerator7	q17numpc7	q17numerator8	q17numpc8	q17numerator9	q17numpc9	q17numerator10	q17numpc10	q17numerator11	q17numpc11
	0 0	1	4.3	0	0	2	8.7	0	0	1	4.3

Paediatric HDU											
beds_Total	PICU beds_0_Number	PICU beds_0_%	PICU beds_1_Number	PICU beds_1_%	PICU beds_2_Number	PICU beds_2_%	PICU beds_3_Number	PICU beds_3_%	PICU beds_4_Number	PICU beds_4_%	PICU beds_5_Number
Q1.7	Q1.8	Q1.8	Q1.8								
q17denominator	q18numerator0	q18numpc0	q18numerator1	q18numpc1	q18numerator2	q18numpc2	q18numerator3	q18numpc3	q18numerator4	q18numpc4	q18numerator5
23	18	78.3	0	0	0	0	0	0	0	0	0

PICU beds_5_%	PICUbeds_6_Number	PICU beds_6_%
Q1.8	Q1.8	Q1.8
q18numpc5	q18numerator6	q18numpc6
	nl o	(

								PICU		
PICU beds_7_Number	PICU beds_7_%	PICU beds_8_Number	PICU beds_8_%	PICU beds_9_Number	PICU beds_9_%	PICU beds_10_Number	PICU beds_10_%	beds_>10_Number	PICU beds_>10_%	PICU beds_Total
Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8	Q1.8
q18numerator7	q18numpc7	q18numerator8	q18numpc8	q18numerator9	q18numpc9	q18numerator10	q18numpc10	q18numerator11	q18numpc11	q18denominator
0	0	2	8.7	1	4.3	0	0	2	8.7	23

Control for DEM	Contract for DEM	Contain for DEM	DIGIT		DIGIT t	DIGIT - A	DIGIT	DIGIT t	DIGIT t	DIGIT	DIGIT
1 °	*	.,	PICU outreach service Days None N	DICII outroach							PICU outreach
Number				service Days None %	service_Days_None_To				service_Days_ Tuesday_Number	service_Days_ Tuesday_%	service_Days_ Tuesday_Total
						·=	·-	•=	·-		
Q1.9	Q1.9	Q1.9	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10
q19numerator	q19numpc	q19denominator	q110anumerator	q110anumpc	q110adenominator	q110bnumerator	q110bnumpc	q110bdenominator	q110cnumerator	q110cnumpc	q110cdenominator
21	95.5	22	12	57.1	21	6	28.6	21	6	30	20

PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach
									service_Days_		service_Days_
Wednesday_Number	Wednesday_%	Wednesday_Total	Thursday_Number	Thursday_%	Thursday_Total						Saturday_Total
Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10
q110dnumerator	q110dnumpc	q110ddenominator	q110enumerator	q110enumpc	q110edenominator	q110fnumerator	q110fnumpc	q110fdenominator	q110gnumerator	q110gnumpc	q110gdenominator
6	28.6	21	6	28.6	21	. 6	28.6	21	6	28.6	21

								Paediatric intake		
						Paediatric intake	Paediatric intake	rota_staff members	Paediatric intake	Paediatric intake
PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	PICU outreach	rota_staff members	rota_staff members	WTE_Cons.Acute	rota_staff members	rota_staff members
service_Days_	service_Days_	service_Days_	service_Days_	service_Days_	service_Days_	WTE_Cons.Acute	WTE_Cons.Acute	PaedStandard	WTE_Cons.Acute	WTE_Cons.Acute
Sunday_Number	Sunday_%	Sunday_Total	Runs overnight_Number	Runs overnight_%	Runs overnight_Total	PaedNumber	PaedMean	Deviation	PaedMedian	PaedLower quartile
Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.10	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11
q110hnumerator	q110hnumpc	q110hdenominator	q110inumerator	q110inumpc	q110idenominator	q111a_N	q111a_mean	q111a_sd	q111a_median	q111a_lq
6	28.6	21	6	28.6	21	22	9.4	3.5	10	8

									Paediatric intake		Paediatric intake
Paediatric intake	rota_staff members	Paediatric intake	rota_staff members								
rota_staff members	WTE_Paed.Intensive	rota_staff members	WTE_Paed.Intensive								
WTE_Cons.Acute	WTE_Spec.Resp.Paed	WTE_Spec.Resp.Paed	WTE_Spec.Resp.Paed	WTE_Spec.Resp.Paed	WTE_Spec.Resp.Paed	WTE_Spec.Resp.Paed	WTE_Paed.Intensive	WTE_Paed.Intensive	Care ConsStandard	WTE_Paed.Intensive	Care ConsLower
PaedUpper quartile	Number	Mean	Standard Deviation	_Median	Lower quartile	Upper quartile	Care ConsNumber	Care ConsMean	Deviation	Care ConsMedian	quartile
Q1.11											
q111a_uq	q111b_N	q111b_mean	q111b_sd	q111b_median	q111b_lq	q111b_uq	q111c_N	q111c_mean	q111c_sd	q111c_median	q111c_lq
11	22	1.5	2.9	0	0	2	21	0.3	1.3	C	0

									Paediatric intake		
Paediatric intake			Paediatric intake				Paediatric intake		rota_staff members	Paediatric intake	Paediatric intake
rota_staff members	Paediatric intake	Paediatric intake	rota_staff members	Paediatric intake	Paediatric intake	Paediatric intake	rota_Staff member	Paediatric intake	WTE_Paed. SpR/Spec.	rota_staff members	rota_staff members
WTE_Paed.Intensive	rota_staff members	rota_staff members	WTE_Other spec. paed.	rota_staff members	rota_staff members	rota_staff members	WTE_Paed. SpR/Spec.	rota_staff members	trainee &	WTE_Paed. SpR/Spec.	WTE_Paed. SpR/Spec.
Care ConsUpper	WTE_Other spec. paed.	WTE_Other spec. paed.	ConsStandard	WTE_Other spec. paed.	WTE_Other spec. paed.	WTE_Other spec. paed.	trainee &	WTE_Paed. SpR/Spec.	fellowStandard	trainee &	trainee &
quartile	ConsNumber	ConsMean	Deviation	ConsMedian	ConsLower quartile	ConsUpper quartile	fellow_Number	trainee & fellow_Mean	Deviation	fellow_Median	fellowLower quartile
Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11	Q1.11
q111c_uq	q111d_N	q111d_mean	q111d_sd	q111d_median	q111d_lq	q111d_uq	q111e_N	q111e_mean	q111e_sd	q111e_median	q111e_lq
0	19	1.9	3.4	0	0	2	18	8.6	4.4	8.2	6

	Frequency of	Frequency of	Frequency of		Frequency of			Frequency of	Frequency of	Frequency of	
Paediatric intake	paediatric patient	paediatric patient	paediatric patient	Frequency of	paediatric patient	Frequency of	Frequency of	paediatric patient	paediatric patient	paediatric patient	Frequency of
rota_staff members	review by senior	review by senior	review by senior	paediatric patient	review by senior	paediatric patient	paediatric patient	review by senior	review by senior	review by senior	paediatric patient
WTE_Paed. SpR/Spec.	decision maker_On	decision maker_On	decision maker_On	review by senior	decision maker_On	review by senior	review by senior	decision maker_On	decision maker_On	decision maker_On	review by senior
trainee &	weekdays_Twice	weekdays_Twice	weekdays_Daily_Numb	decision maker_On	weekdays_Other_Num	decision maker_On	decision maker_On	weekends_Twice	weekends_Twice	weekends_Daily_Num	decision maker_On
fellow_Upper quartile	daily_Number	daily_%	er	weekdays_Daily_%	ber	weekdays_Other_%	weekdays_Total	daily_Number	daily_%	ber	weekends_Daily_%
Q1.11	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12	Q1.12
q111e_uq	q112anumerator1	q112anumpc1	q112anumerator2	q112anumpc2	q112anumerator3	q112anumpc3	q112adenominator	q112cnumerator1	q112cnumpc1	q112cnumerator2	q112cnumpc2
12	8	34.8	8	34.8	7	30.4	23	7	30.4	10	43.5

Frequency of											
paediatric patient	Frequency of	Frequency of									
review by senior	paediatric patient	paediatric patient									
decision maker_On	review by senior	review by senior	Staff in paediatric		Staff in paediatric		Staff in paediatric				
weekends_Other_Num	decision maker_On	decision maker_On	team_WTE_FY1_Numb	Staff in paediatric	team_WTE_FY1_Stand	team_WTE_FY1_Media	team_WTE_FY1_Lower	team_WTE_FY1_Upper	team_WTE_FY2_Numb	Staff in paediatric	team_WTE_FY2_Stand
ber	weekends_Other_%	weekends_Total	er	team_WTE_FY1_Mean	ard Deviation	n	quartile	quartile	er	team_WTE_FY2_Mean	ard Deviation
Q1.12	Q1.12	Q1.12	Q1.13								
q112cnumerator3	q112cnumpc3	q112cdenominator	q113a_N	q113a_mean	q113a_sd	q113a_median	q113a_lq	q113a_uq	q113b_N	q113b_mean	q113b_sd
6	26.1	23	21	1	1.3	1	0	1	22	1.9	1.2

Chaff in an adiabata	Chaff in an addated	Chaff in an addated	Chaff in an adiabata	Chaff in an addated	Chaff in an adiabat	Chaff in an adiabata	Chaff in an addated	Chaff in an addahala	Chaff in an adiabata	Chaff in an adiabata	Chaff in an adiabata
Staff in paediatric team WTE FY2 Media		· ·		· ·	Staff in paediatric team WTE CT1/CT2 S		Staff in paediatric team WTE CT1/CT2 L	· ·		· ·	Staff in paediatric team WTF_ST3&above
n					tandard Deviation			pper quartile	_Number	_Mean	_Standard Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113b_median	q113b_lq	q113b_uq	q113c_N	q113c_mean	q113c_sd	q113c_median	q113c_lq	q113c_uq	q113d_N	q113d_mean	q113d_sd
2	1	3	18	5.4	4.4	6.5	1	8	22	7.1	4.4

											ļ
					Staff in paediatric						Staff in paediatric
Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric		team_WTE_Associate	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric		team_WTE_Staff
team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_Associate	team_WTE_Associate	SpecStandard	team_WTE_Associate	team_WTE_Associate	team_WTE_Associate	team_WTE_Staff	team_WTE_Staff	grade_Standard
_Median	_Lower quartile	_Upper quartile	SpecNumber	SpecMean	Deviation	SpecMedian	SpecLower quartile	SpecUpper quartile	grade_Number	grade_Mean	Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113d_median	q113d_lq	q113d_uq	q113e_N	q113e_mean	q113e_sd	q113e_median	q113e_lq	q113e_uq	q113f_N	q113f_mean	q113f_sd
7	3.9	11	22	0.7	1.0	0	0	1	21	1.4	2.9

											Staff in paediatric
Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	team_WTE_Resp.Nurse
team_WTE_Staff	team_WTE_Staff	team_WTE_Staff	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	ConsStandard
grade_Median	grade_Lower quartile	grade_Upper quartile	t_Number	t_Mean	t_Standard Deviation	t_Median	t_Lower quartile	t_Upper quartile	ConsNumber	ConsMean	Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113f_median	q113f_lq	q113f_uq	q113g_N	q113g_mean	q113g_sd	q113g_median	q113g_lq	q113g_uq	q113h_N	q113h_mean	q113h_sd
0	0	1	23	0.3	0.7	0	0	0.1	23	0.1	0.6

					Staff in paediatric						Staff in paediatric
Staff in paediatric	team_WTE_Paed.Nurs	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	team_WTE_Asthma				
team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	e Cons_Standard	team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	team_WTE_Asthma	team_WTE_Asthma	Nurse SpecStandard
ConsMedian	ConsLower quartile	ConsUpper quartile	e ConsNumber	e Cons_Mean	Deviation	e Cons_Median	e ConsLower quartile	e ConsUpper quartile	Nurse SpecNumber	Nurse SpecMean	Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13							
q113h_median	q113h_lq	q113h_uq	q113i_N	q113i_mean	q113i_sd	q113i_median	q113i_lq	q113i_uq	q113j_N	q113j_mean	q113j_sd
0	0	0	23	0	0	0	0	0	23	0.6	0.7

Staff in paediatric team_WTE_Asthma	team_WTE_Asthma	team_WTE_Asthma Nurse SpecUpper	team_WTE_Other	Staff in paediatric	spec. resp. nurse_Standard	team_WTE_Other spec. resp.	team_WTE_Other spec. resp.	team_WTE_Other spec. resp.		Staff in paediatric team_WTE_Other spec. paed.	Staff in paediatric team_WTE_Other spec. paed. nurse_Standard Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113j_median 0.6		q113j_uq	q113k_N 23		q113k_sd	q113k_median 0.2		q113k_uq	q113I_N	q113I_mean	q113I_sd 2.0

											ļ
Staff in paediatric	Staff in paediatric	Staff in paediatric			Staff in paediatric				Staff in paediatric		Staff in paediatric
team_WTE_Other	team_WTE_Other	team_WTE_Other	Staff in paediatric	Staff in paediatric	team_WTE_Physiother	Staff in paediatric	Staff in paediatric	Staff in paediatric	team_WTE_Research	Staff in paediatric	team_WTE_Research
spec. paed.	spec. paed.	spec. paed.	team_WTE_Physiother	team_WTE_Physiother	apist_Standard	team_WTE_Physiother	team_WTE_Physiother	team_WTE_Physiother	registrar/fellow_Numb	team_WTE_Research	registrar/fellow_Stand
nurse_Median	nurse_Lower quartile	nurse_Upper quartile	apist_Number	apist_Mean	Deviation	apist_Median	apist_Lower quartile	apist_Upper quartile	er	registrar/fellow_Mean	ard Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113l_median	q113l_lq	q113l_uq	q113m_N	q113m_mean	q113m_sd	q113m_median	q113m_lq	q113m_uq	q113n_N	q113n_mean	q113n_sd
0	0	2	22	1.0	0.9	1	0	1.8	23	0.0	0.2

Staff in paediatric	Staff in paediatric	Staff in paediatric			Staff in paediatric						Staff in paediatric
team_WTE_Research	team_WTE_Research	team_WTE_Research	Staff in paediatric	Staff in paediatric	team_WTE_Research	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	Staff in paediatric	team_WTE_Resp.paed.
registrar/fellow_Media	registrar/fellow_Lower	registrar/fellow_Upper	team_WTE_Research	team_WTE_Research	nurse_Standard	team_WTE_Research	team_WTE_Research	team_WTE_Research	team_WTE_Resp.paed.	team_WTE_Resp.paed.	cons_Standard
n	quartile	quartile	nurse_Number	nurse_Mean	Deviation	nurse_Median	nurse_Lower quartile	nurse_Upper quartile	cons_Number	cons_Mean	Deviation
Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13	Q1.13
q113n_median	q113n_lq	q113n_uq	q113o_N	q113o_mean	q113o_sd	q113o_median	q113o_lq	q113o_uq	q113p_N	q113p_mean	q113p_sd
0	0	0	21	0.3	0.6	0	0	0.6	23	1.1	1.5

								Unfilled posts in	Unfilled posts in		Unfilled posts in
Staff in paediatric	paediatric	paediatric	Unfilled posts in	paediatric							
team WTE Resp.paed.	team WTE Resp.paed.	team WTE Resp.paed.	team WTE Other Nu	team WTE Other Me	team WTE Other Sta	team WTE Other Me	team WTE Other Low	team WTE Other Upp	team WTE FY1 Numb	paediatric	team WTE FY1 Stand
conse_Median	cons_Lower quartile	cons_Upper quartile	mber	an	ndard Deviation	dian	er quartile	er quartile	er	team_WTE_FY1_Mean	ard Deviation
Q1.13	Q1.13	Q1.14	Q1.14	Q1.14							
q113p_median	q113p_lq	q113p_uq	q113q_N	q113q_mean	q113q_sd	q113q_median	q113q_lq	q113q_uq	q114a_N	q114a_mean	q114a_sd
0	0	2	19	0.8	2.3	0	0	0	19	0.1	0.2

Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in		Unfilled posts in	Unfilled posts in	Unfilled posts in				
paediatric	paediatric	paediatric	paediatric	Unfilled posts in	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric
team_WTE_FY1_Media	team_WTE_FY1_Lower	team_WTE_FY1_Upper	team_WTE_FY2_Numb	paediatric	team_WTE_FY2_Stand	team_WTE_FY2_Media	team_WTE_FY2_Lower	team_WTE_FY2_Upper	team_WTE_CT1/CT2_N	team_WTE_CT1/CT2_	team_WTE_CT1/CT2_S
n	quartile	quartile	er	team_WTE_FY2_Mean	ard Deviation	n	quartile	quartile	umber	Mean	tandard Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114a_median	q114a_lq	q114a_uq	q114b_N	q114b_mean	q114b_sd	q114b_median	q114b_lq	q114b_uq	q114c_N	q114c_mean	q114c_sd
0	0	0	19	0.1	0.3	0	0	0	18	0.4	0.6

											Unfilled posts in
Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled postsin	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric
paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Associate
team_WTE_CT1/CT2_	team_WTE_CT1/CT2_L	team_WTE_CT1/CT2_U	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_ST3&above	team_WTE_Associate	team_WTE_Associate	SpecStandard
Median	ower quartile	pper quartile	_Number	_Mean	_Standard Deviation	_Median	_Lower quartile	_Upper quartile	SpecNumber	SpecMean	Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114c_median	q114c_lq	q114c_uq	q114d_N	q114d_mean	q114d_sd	q114d_median	q114d_lq	q114d_uq	q114e_N	q114e_mean	q114e_sd
0	0	1	19	1.1	1.0	1	0	2	20	0.1	0.2

					Unfilled posts in						
Unfilled posts in	Unfilled posts in	Unfilled postsin	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in
paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Staff	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric
team_WTE_Associate	team_WTE_Associate	team_WTE_Associate	team_WTE_Staff	team_WTE_Staff	grade_Standard	team_WTE_Staff	team_WTE_Staff	team_WTE_Staff	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis
SpecMedian	SpecLower quartile	SpecUpper quartile	grade_Number	grade_Mean	Deviation	grade_Median	grade_Lower quartile	grade_Upper quartile	t_Number	t_Mean	t_Standard Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114e_median	q114e_lq	q114e_uq	q114f_N	q114f_mean	q114f_sd	q114f_median	q114f_lq	q114f_uq	q114g_N	q114g_mean	q114g_sd
0	0	0	20	0.2	0.5	0	0	0	21	0.0	0.2

					Unfilled posts in						Unfilled posts in
Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric
paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Resp.Nurse	paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Paed.Nurs
team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Physiologis	team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	ConsStandard	team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	team_WTE_Resp.Nurse	team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	e Cons_Standard
t_Median	t_Lower quartile	t_Upper quartile	ConsNumber	ConsMean	Deviation	ConsMedian	ConsLower quartile	ConsUpper quartile	e ConsNumber	e Cons_Mean	Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114g_median	q114g_lq	q114g_uq	q114h_N	q114h_mean	q114h_sd	q114h_median	q114h_lq	q114h_uq	q114i_N	q114i_mean	q114i_sd
0	0	0	22	0	0	0	0	0	23	0	0

											Unfilled posts in
					Unfilled posts in		Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric
Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	paediatric	paediatric	paediatric	paediatric	team_WTE_Other
paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Asthma	paediatric	team_WTE_Asthma	team_WTE_Asthma	team_WTE_Other	team_WTE_Other	spec. resp.
team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	team_WTE_Paed.Nurs	team_WTE_Asthma	team_WTE_Asthma	Nurse SpecStandard	team_WTE_Asthma	Nurse SpecLower	Nurse SpecUpper	spec. resp.	spec. resp.	nurse_Standard
e Cons_Median	e ConsLower quartile	e ConsUpper quartile	Nurse SpecNumber	Nurse SpecMean	Deviation	Nurse SpecMedian	quartile	quartile	nurse_Number	nurse_Mean	Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114i_median	q114i_lq	q114i_uq	q114j_N	q114j_mean	q114j_sd	q114j_median	q114j_lq	q114j_uq	q114k_N	q114k_mean	q114k_sd
0	0	0	23	0.1	0.1		0	0	23	0	(

					Unfilled posts in						ļ
Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	Unfilled posts in	Unfilled posts in			Unfilled posts in
paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Other	paediatric	paediatric	paediatric	Unfilled posts in	Unfilled posts in	paediatric
team_WTE_Other	team_WTE_Other	team_WTE_Other	team_WTE_Other	team_WTE_Other	spec. paed.	team_WTE_Other	team_WTE_Other	team_WTE_Other	paediatric	paediatric	team_WTE_Physiother
spec. resp.	spec. resp.	spec. resp.	spec. paed.	spec. paed.	nurse_Standard	spec. paed.	spec. paed.	spec. paed.	team_WTE_Physiother	team_WTE_Physiother	apist_Standard
nurse_Median	nurse_Lower quartile	nurse_Upper quartile	nurse_Number	nurse_Mean	Deviation	nurse_Median	nurse_Lower quartile	nurse_Upper quartile	apist_Number	apist_Mean	Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114k_median	q114k_lq	q114k_uq	q114l_N	q114l_mean	q114l_sd	q114l_median	q114l_lq	q114l_uq	q114m_N	q114m_mean	q114m_sd
0	0	0	22	0	0	0	0	0	22	0	0

			Unfilled posts in		Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in			Unfilled posts in
Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	paediatric	paediatric	paediatric	paediatric	Unfilled posts in	Unfilled posts in	paediatric
paediatric	paediatric	paediatric	team_WTE_Research	paediatric	team_WTE_Research	team_WTE_Research	team_WTE_Research	team_WTE_Research	paediatric	paediatric	team_WTE_Research
team_WTE_Physiother	team_WTE_Physiother	team_WTE_Physiother	registrar/fellow_Numb	team_WTE_Research	registrar/fellow_Stand	registrar/fellow_Media	registrar/fellow_Lower	registrar/fellow_Upper	team_WTE_Research	team_WTE_Research	nurse_Standard
apist_Median	apist_Lower quartile	apist_Upper quartile	er	registrar/fellow_Mean	ard Deviation	n	quartile	quartile	nurse_Number	nurse_Mean	Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114m_median	q114m_lq	q114m_uq	q114n_N	q114n_mean	q114n_sd	q114n_median	q114n_lq	q114n_uq	q114o_N	q114o_mean	q114o_sd
0	0	0	22	0	0	0	0	0	21	0	0

					Unfilled posts in						
Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	paediatric	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in	Unfilled posts in
paediatric	paediatric	paediatric	paediatric	paediatric	team_WTE_Resp.paed.	paediatric	paediatric	paediatric	paediatric	paediatric	paediatric
team_WTE_Research	team_WTE_Research	team_WTE_Research	team_WTE_Resp.paed.	team_WTE_Resp.paed.	cons_Standard	team_WTE_Resp.paed.	team_WTE_Resp.paed.	team_WTE_Resp.paed.	team_WTE_Other_Nu	team_WTE_Other_Me	team_WTE_Other_Sta
nurse_Median	nurse_Lower quartile	nurse_Upper quartile	cons_Number	cons_Mean	Deviation	conse_Median	cons_Lower quartile	cons_Upper quartile	mber	an	ndard Deviation
Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14	Q1.14
q114o_median	q114o_lq	q114o_uq	q114p_N	q114p_mean	q114p_sd	q114p_median	q114p_lq	q114p_uq	q114q_N	q114q_mean	q114q_sd
0	0	0	22	0.0	0.2	0	0	0	21	0.1	0.4

Unfilled posts in	Unfilled posts in	Unfilled posts in									
paediatric	paediatric	paediatric	Smoking cessation	Smoking cessation	Smoking cessation	Smoking cessation	Smoking cessation		Smoking cessation		Smoking cessation
team_WTE_Other_Me	team_WTE_Other_Low	team_WTE_Other_Upp	service_No service	service_No service	service_Less than 0.5	service_Less than 0.5	service_0.5	Smoking cessation	service_1	Smoking cessation	service_2
dian	er quartile	er quartile	available_Number	available_%	WTE_Number	WTE_%	WTE_Number	service_ 0.5 WTE_%	WTE_Number	service_1 WTE_%	WTE_Number
Q1.14	Q1.14	Q1.14	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15
q114q_median	q114q_lq	q114q_uq	q115numerator1	q115numpc1	q115numerator2	q115numpc2	q115numerator3	q115numpc3	q115numerator4	q115numpc4	q115numerator5
0	0	0	8	40	0	0	0	0	3	15	0

	Smoking cessation		Smoking cessation		Smoking cessation			Smoking-cessation	Smoking-cessation	Smoking-cessation	Smoking-cessation
Smoking cessation		Smoking cessation				Smoking cessation			•		pharmacotherapies_16
service_2 WTE_%	WTE_Number	service_3 WTE_%	WTE_Number	service_4+ WTE_%	WTE_Number	service_Other WTE_%	service_Total	years_Number	years_%	years_Total	years_Number
Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.15	Q1.16	Q1.16	Q1.16	Q1.16
q115numpc5	q115numerator6	q115numpc6	q115numerator7	q115numpc7	q115numerator8	q115numpc8	q115denominator	q116anumerator	q116anumpc	q116adenominator	q116bnumerator
	0 1		0	0	8	40	20	8	44.4	18	6

										If yes, smoking-
								If yes, smoking-cessation	If yes, smoking-	cessation
Smoking-cessation	pharmacotherapies_Nic	cessation	pharmacotherapies_Ni							
pharmacotherapies_16	pharmacotherapies_16	pharmacotherapies_Ad	pharmacotherapies_Ad	pharmacotherapies_Ad	pharmacotherapies_No	pharmacotherapies_No	pharmacotherapies_No	otine	pharmacotherapies_Ni	cotine
years_%	years_Total	ults only_Number	ults only_%	ults only_Total	pharmacy_Number	pharmacy_%	pharmacy_Total	replacement_Number	cotine replacement_%	replacement_Total
Q1.16	Q1.16	Q1.16								
q116bnumpc	q116bdenominator	q116cnumerator	q116cnumpc	q116cdenominator	q116dnumerator	q116dnumpc	q116ddenominator	q116enumerator	q116enumpc	q116edenominator
40	15	8	53.3	15	0	0	13	7	63.6	11

			Organisation of ac	ute respiratory car	e						
If yes, smoking-	If yes, smoking-	If yes, smoking-	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric
cessation	cessation	cessation	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory
pharmacotherapies_Ot	pharmacotherapies_Ot	pharmacotherapies_Ot	consultant_Days_None	consultant_Days_None	consultant_Days_None	consultant_Days_Mon	consultant_Days_Mon	consultant_Days_Mon	consultant_Days_Tues	consultant_Days_Tues	consultant_Days_Tues
her_Number	her_%	her_Total	_Number	_%	_Total	day_Number	day_%	day_Total	day_Number	day_%	day_Total
Q1.16	Q1.16	Q1.16	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1
q116fnumerator	q116fnumpc	q116fdenominator	q21anumerator	q21anumpc	q21adenominator	q21bnumerator	q21bnumpc	q21bdenominator	q21cnumerator	q21cnumpc	q21cdenominator
1	14.3	7	10	43.5	23	9	56.3	16	8	53.3	15

On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric
respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory
consultant_Days_Wed	consultant_Days_Wed	consultant_Days_Wed	consultant_Days_Thurs	consultant_Days_Thurs	consultant_Days_Thurs	consultant_Days_Frida	consultant_Days_Frida	consultant_Days_Frida	consultant_Days_Satur	consultant_Days_Satur	consultant_Days_Satur
nesday_Number	nesday_%	nesday_Total	day_Number	day_%	day_Total	y_Number	у_%	y_Total	day_Number	day_%	day_Total
Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1	Q2.1
q21dnumerator	q21dnumpc	q21ddenominator	q21enumerator	q21enumpc	q21edenominator	q21fnumerator	q21fnumpc	q21fdenominator	q21gnumerator	q21gnumpc	q21gdenominator
8	53.3	15	9	60	15	8	53.3	15	5	35.7	14

									Senior decision maker	Senior decision maker	Senior decision maker
									from resp. team	from resp. team	from resp. team
On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	On-call paediatric	undertake ward rounds	undertake ward rounds	undertake ward rounds
respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	respiratory	of new asthma	of new asthma	of new asthma
consultant_Days_Sund	consultant_Days_Sund	consultant_Days_Sund	consultant_Days_In	consultant_Days_In	consultant_Days_In	consultant_Days_Out	consultant_Days_Out	consultant_Days_Out	patients_MAU/ad.war	patients_MAU/ad.war	patients_MAU/ad.war
ay_Number	ay_%	ayy_Total	hours_Number	hours_%	hours_Total	of hours_Number	of hours_%	of hours_Total	d_Days_None_Number	d_Days_None_%	d_Days_None_Total
Q2.1	Q2.1	Q2.1	Q2.1.1	Q2.1.1	Q2.1.1	Q2.1.1	Q2.1.1	Q2.1.1	Q2.2	Q2.2	Q2.2
q21hnumerator	q21hnumpc	q21hdenominator	q211anumerator	q211anumpc	q211adenominator	q211bnumerator	q211bnumpc	q211bdenominator	q22anumerator	q22anumpc	q22adenominator
6	42.9	14	9	42.9	21	6	30	20	10	47.6	21

Senior decision maker			Senior decision maker			Senior decision maker		Senior decision maker	Senior decision maker		Senior decision maker
from resp. team	Senior decision maker	Senior decision maker	from resp. team	Senior decision maker	Senior decision maker	from resp. team	Senior decision maker	from resp. team	from resp. team	Senior decision maker	from resp. team
undertake ward rounds	from resp. team	from resp. team	undertake ward rounds	from resp. team	from resp. team	undertake ward rounds	from resp. team	undertake ward rounds	undertake ward rounds	from resp. team	undertake ward rounds
of new asthma	undertake ward rounds	undertake ward rounds	of new asthma	undertake ward rounds	undertake ward rounds	of new asthma	undertake ward rounds	of new asthma	of new asthma	undertake ward rounds	of new asthma
patients_MAU/ad.war	of new asthma	of new asthma	patients_MAU/ad.war	of new asthma	of new asthma	patients_MAU/ad.war	of new asthma	patients_MAU/ad.war	patients_MAU/ad.war	of new asthma	patients_MAU/ad.war
d_Days_Monday_Num	patients_MAU/ad.war	patients_MAU/ad.war	d_Days_Tuesday_Num	patients_MAU/ad.war	patients_MAU/ad.war	d_Days_Wednesday_N	patients_MAU/ad.war	d_Days_Wednesday_T	d_Days_Thursday_Nu	patients_MAU/ad.war	d_Days_Thursday_Tota
ber	d_Days_Monday_%	d_Days_Monday_Total	ber	d_Days_Tuesday_%	d_Days_Tuesday_Total	umber	d_Days_Wednesday_%	otal	mber	d_Days_Thursday_%	I
Q2.2											
q22bnumerator	q22bnumpc	q22bdenominator	q22cnumerator	q22cnumpc	q22cdenominator	q22dnumerator	q22dnumpc	q22ddenominator	q22enumerator	q22enumpc	q22edenominator
6	46.2	13	5	38.5	13	5	38.5	13	5	38.5	13

Senior decision maker			Senior decision maker		Senior decision maker	Senior decision maker					
from resp. team	Senior decision maker	Senior decision maker	from resp. team	Senior decision maker	from resp. team	from resp. team	Senior decision maker				
undertake ward rounds	from resp. team	from resp. team	undertake ward rounds	from resp. team	undertake ward rounds	undertake ward rounds	from resp. team				
of new asthma	undertake ward rounds	undertake ward rounds	of new asthma	undertake ward rounds	of new asthma	of new asthma	undertake ward rounds				
patients_MAU/ad.war	of new asthma	of new asthma	patients_MAU/ad.war	of new asthma	patients_MAU/ad.war	patients_MAU/ad.war	of new asthma				
d_Days_Friday_Numbe	patients_MAU/ad.war	patients_MAU/ad.war	d_Days_Saturday_Num	patients_MAU/ad.war	d_Days_Saturday_Tota	d_Days_Sunday_Numb	patients_MAU/ad.war	patients_MAU/ad.war	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D
r	d_Days_Friday_%	d_Days_Friday_Total	ber	d_Days_Saturday_%	I	er	d_Days_Sunday_%	d_Days_Sunday_Total	ays_None_Number	ays_None_%	ays_None_Total
Q2.2	Q2.3	Q2.3	Q2.3								
q22fnumerator	q22fnumpc	q22fdenominator	q22gnumerator	q22gnumpc	q22gdenominator	q22hnumerator	q22hnumpc	q22hdenominator	q23anumerator	q23anumpc	q23adenominator
6	46.2	13	3	23.1	13	3	23.1	13	11	52.4	21

						Senior decision maker					
Senior decision maker	from resp. team	Senior decision maker									
from resp. team	undertake ward rounds	from resp. team									
undertake ward rounds	of new asthma	undertake ward rounds									
of new asthma	patients_Resp.ward_D	of new asthma									
patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	ays_Wednesday_Num	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D	patients_Resp.ward_D
ays_Monday_Number	ays_Monday_%	ays_Monday_Total	ays_Tuesday_Number	ays_Tuesday_%	ays_Tuesday_Total	ber	ays_Wednesday_%	ays_Wednesday_Total	ays_Thursday_Number	ays_Thursday_%	ays_Thursday_Total
Q2.3											
q23bnumerator	q23bnumpc	q23bdenominator	q23cnumerator	q23cnumpc	q23cdenominator	q23dnumerator	q23dnumpc	q23ddenominator	q23enumerator	q23enumpc	q23edenominator
3	27.3	11	3	27.3	11	3	27.3	11	3	27.3	11

									Senior decision maker		Senior decision maker
Senior decision maker	from resp. team	Senior decision maker	from resp. team								
from resp. team	undertake ward rounds	from resp. team	undertake ward rounds								
undertake ward rounds	of new asthma	undertake ward rounds	of new asthma								
of new asthma	patients_Other	of new asthma	patients_Other								
patients Resp.ward D	wards Days None Nu	patients Other	wards Days None Tot								
ays_Friday_Number	ays_Friday_%	ays_Friday_Total	ays_Saturday_Number	ays_Saturday_%	ays_Saturday_Total	ays_Sunday_Number	ays_Sunday_%	ays_Sunday_Total	mber	wards_Days_None_%	al
Q2.3	Q2.4	Q2.4	Q2.4								
q23fnumerator	q23fnumpc	q23fdenominator	q23gnumerator	q23gnumpc	q23gdenominator	q23hnumerator	q23hnumpc	q23hdenominator	q24anumerator	q24anumpc	q24adenominator
3	27.3	11	2	18.2	11	2	18.2	11	9	42.9	21

| Senior decision maker |
|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|
| from resp. team |
| undertake ward rounds |
| of new asthma |
| patients_Other |
| wards_Days_Monday_ | wards_Days_Monday_ | wards_Days_Monday_ | wards_Days_Tuesday_ | wards_Days_Tuesday_ | wards_Days_Tuesday_ | wards_Days_Wednesd | wards_Days_Wednesd | wards_Days_Wednesd | wards_Days_Thursday | wards_Days_Thursday | wards_Days_Thursday |
| Number | % | Total | Number | % | Total | ay_Number | ay_% | ay_Total | _Number | _% | _Total |
| Q2.4 |
| q24bnumerator | q24bnumpc | q24bdenominator | q24cnumerator | q24cnumpc | q24cdenominator | q24dnumerator | q24dnumpc | q24ddenominator | q24enumerator | q24enumpc | q24edenominator |
| 6 | 50 | 12 | 5 | 41.7 | 12 | 5 | 41.7 | 12 | 5 | 41.7 | 12 |

Senior decision maker		Senior decision maker		Senior decision maker							
from resp. team	Senior decision maker	from resp. team	Senior decision maker	from resp. team							
undertake ward rounds	from resp. team	undertake ward rounds	from resp. team	undertake ward rounds	Respiratory nurse	Respiratory nurse	Respiratory nurse				
of new asthma	undertake ward rounds	of new asthma	undertake ward rounds	of new asthma	available to review	available to review	available to review				
patients_Other	of new asthma	patients_Other	patients_Other	patients_Other	patients_Other	patients_Other	of new asthma	patients_Other	asthma paed.	asthma paed.	asthma paed.
wards_Days_Friday_Nu	patients_Other	wards_Days_Friday_To	wards_Days_Saturday_	wards_Days_Saturday_	wards_Days_Saturday_	wards_Days_Sunday_N	patients_Other	wards_Days_Sunday_T	patients_Days_None_	patients_Days_None_	patients_Days_None_T
mber	wards_Days_Friday_%	tal	Number	%	Total	umber	wards_Days_Sunday_%	otal	Number	%	otal
Q2.4	Q2.5	Q2.5	Q2.5								
q24fnumerator	q24fnumpc	q24fdenominator	q24gnumerator	q24gnumpc	q24gdenominator	q24hnumerator	q24hnumpc	q24hdenominator	q25anumerator	q25anumpc	q25adenominator
6	50	12	1	8.3	12	1	8.3	12	5	21.7	23

Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse
available to review	available to review	available to review	available to review	available to review	available to review	available to review	available to review	available to review	available to review	available to review	available to review
asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.	asthma paed.
patients_Days_Monda	patients_Days_Monda	patients_Monday_Tota	patients_Days_Tuesda	patients_Days_Tuesda	patients_Days_Tuesda	patients_Days_Wedne	patients_Days_Wedne	patients_Days_Wedne	patients_Days_Thursda	patients_Days_Thursda	patients_Days_Thursda
y_Number	у_%	I	y_Number	у_%	y_Total	sday_Number	sday_%	sday_Total	y_Number	y_%	y_Total
Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5
q25bnumerator	q25bnumpc	q25bdenominator	q25cnumerator	q25cnumpc	q25cdenominator	q25dnumerator	q25dnumpc	q25ddenominator	q25enumerator	q25enumpc	q25edenominator
15	83.3	18	15	83.3	18	15	88.2	17	11	68.8	16

Respiratory nurse	Respiratory nurse	Respiratory nurse	Respiratory nurse							
available to review	available to review	available to review	available to review							
asthma paed.	asthma paed.	asthma paed.	asthma paed.	Patients with access to	Patients with access to					
patients_Days_Friday_	patients_Days_Friday_	patients_Days_Friday_	patients_Days_Saturda	patients_Days_Saturda	patients_Days_Saturda	patients_Days_Sunday	patients_Days_Sunday	patients_Days_Sunday	paediatric respiratory	paediatric respiratory
Number	%	Total	y_Number	y_%	y_Total	_Number	_%	_Total	nurse_None_Number	nurse_None_%
Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.5	Q2.6	Q2.6
q25fnumerator	q25fnumpc	q25fdenominator	q25gnumerator	q25gnumpc	q25gdenominator	q25hnumerator	q25hnumpc	q25hdenominator	q26anumerator0	q26anumpc0
12	70.6	17	0	0	17	0	0	17	6	26.1

		Patients with access to									
		paediatric respiratory	Patients with access to								
		nurse_Only those	paediatric respiratory								
Patients with access to	Patients with access to	under	nurse_Only those	Patients with access to	Patients with access to	Patients with access to					
paediatric respiratory	paediatric respiratory	resp.physician_Numbe	under	paediatric respiratory	paediatric respiratory	paediatric respiratory	Asthma		Asthma		Asthma lead_Hospital
nurse_All_Number	nurse_All_%	r	resp.physician_%	nurse_Other_Number	nurse_Other_%	nurse_Total	lead_No_Number	Asthma lead_No_%	lead_Yes_Number	Asthma lead_Yes_%	wide lead_Number
Q2.6	Q2.7	Q2.7	Q2.7	Q2.7	Q2.7						
q26anumerator1	q26anumpc1	q26anumerator2	q26anumpc2	q26anumerator3	q26anumpc3	q26adenominator	q27numerator0	q27numpc0	q27numerator1	q27numpc1	q27numerator2
14	60.9	0	0	3	13.0	23	4	18.2	16	72.7	2

									Managing respirat	ory failure - emerg	ency oxygen therap
						16	16				
Asthma lead_Hospital		If yes, responsible for		If yes, responsible for training Yes, but also	If yes, responsible for	Paediatric oxygen	Paediatric oxygen	Paediatric oxygen			
				training_Yes_Number		'			policy_No_Number	, .	policy_Yes_Number
wide lead_%	Astiilia leau_Total	training_NO_Number	training_NO_%	training_res_ivumber	training_res_/	auuit_ivuilibei	auuit_%	training_rotal	policy_No_Nullibel	policy_NO_%	policy_res_ivullibel
Q2.7	Q2.7	Q2.7.1	Q2.7.1	Q2.7.1	Q2.7.1	Q2.7.1	Q2.7.1	Q2.7.1	Q3.1	Q3.1	Q3.1
q27numpc2	q27denominator	q271numerator0	q271numpc0	q271numerator1	q271numpc1	q271numerator2	q271numpc2	q271denominator	q31numerator0	q31numpc0	q31numerator1
9.1	22	6	31.6	12	63.2	1	5.3	19	6	26.1	12

у											
						Paper or electronic	Paper or electronic	Paper or electronic			Paediatric med.
	Paediatric oxygen			Paper or electronic	Paper or electronic	prescribing_Electronic	prescribing_Electronic	prescribing_Electronic	Paper or electronic		chart/record with place
Paediatric oxygen	policy_Not	Paediatric oxygen	Paediatric oxygen	prescribing_Paper	prescribing_Paper	partially	partially	fully	prescribing_Electronic	Paper or electronic	to prescribe
policy_Yes_%	known_Number	policy_Not known_%	policy_Total	only_Number	only_%	implemented_Number	implemented_%	implemented_Number	fully implemented_%	prescribing_Total	oxygen_Yes_Number
Q3.1	Q3.1	Q3.1	Q3.1	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.2	Q3.3
q31numpc1	q31numerator2	q31numpc2	q31denominator	q32numerator1	q32numpc1	q32numerator2	q32numpc2	q32numerator3	q32numpc3	q32denominator	q33numerator
52.2	5	21.7	23	11	47.8	6	26.1	6	26.1	23	17

											Monitoring chart which
Paediatric med.	Paediatric med.	Monitoring chart which			Monitoring chart which	allows the follow to be					
chart/record with place	chart/record with place	allows the follow to be	Monitoring chart which	Monitoring chart which	allows the follow to be	recorded: Monitoring					
to prescribe	to prescribe	recorded:	allows the follow to be	allows the follow to be	recorded: Target	recorded: Target	recorded: Target	recorded: Actual	recorded: Actual	recorded: Actual	of emergency
oxygen_Yes_%	oxygen_Total	None_Number	recorded: None_%	recorded: None_Total	saturation_Number	saturation_%	saturation_Total	saturation_Number	saturation_%	saturation_Total	oxyNumber
Q3.3	Q3.3	Q3.4									
q33numpc	q33denominator	q34anumerator	q34anumpc	q34adenominator	q34bnumerator	q34bnumpc	q34bdenominator	q34cnumerator	q34cnumpc	q34cdenominator	q34dnumerator
77.3	22	5	41.7	12	10	50	20	22	100	22	22

	Monitoring chart which										Oxygen training
Monitoring chart which	allows the follow to be							Oxygen training		Oxygen training	programme_Monitorin
allows the follow to be	recorded: Monitoring	Oxygen training		Oxygen training	Oxygen training	Oxygen training	Oxygen training	programme_Pres.emer	Oxygen training	programme_Pres.emer	g emergency oxy.
recorded: Monitoring	of emergency	programme_None_Nu	Oxygen training	programme_None_Tot	programme_Not	programme_Not	programme_Not	gency oxy.	programme_Pres.emer	gency oxy.	Nurses and
of emergency oxy%	oxyTotal	mber	programme_None_%	al	known_Number	known_%	known_Total	doctors_Number	gency oxy. doctors_%	doctors_Total	other_Number
Q3.4	Q3.4	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5	Q3.5
q34dnumpc	q34ddenominator	q35anumerator	q35anumpc	q35adenominator	q35bnumerator	q35bnumpc	q35bdenominator	q35cnumerator	q35cnumpc	q35cdenominator	q35dnumerator
95.7	23	7	43.8	16	8	61.5	13	2	20	10	6

		Integrating care a	cross primary and s	econdary sectors							
											ļ
Oxygen training	Oxygen training							MDT			MDT
programme_Monitorin	programme_Monitorin	Paediatric specialist		Paediatric specialist				meetings_Paed.resp.co	MDT	MDT	meetings_Spec.trainee
g emergency oxy.	g emergency oxy.	asthma	Paediatric specialist	asthma	MDT meetings_No_		MDT	ns_	meetings_Paed.resp.co	meetings_Paed.resp.co	_
Nurses and other_%	Nurses and other_Total	service_Yes_Number	asthma service_Yes_%	service_Yes_Total	Number	MDT meetings_No_%	meetings_No_Total	Number	ns_%	ns_Total	Number
Q3.5	Q3.5	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2
q35dnumpc	q35ddenominator	q41numerator	q41numpc	q41denominator	q42anumerator	q42anumpc	q42adenominator	q42bnumerator	q42bnumpc	q42bdenominator	q42cnumerator
46.2	13	15	65.2	23	15	65.2	23	8	66.7	12	1

		MDT						MDT			MDT
MDT	MDT	meetings_Paed.asthma	MDT	MDT				meetings_Hosp.asthma	MDT	MDT	meetings_Hosp.paed.
meetings_Spec.trainee	meetings_Spec.trainee	nurse spec	meetings_Paed.asthma	meetings_Paed.asthma	MDT meetings_GP_		MDT	nurse spec	meetings_Hosp.asthma	meetings_Hosp.asthma	nurse_
_%	_Total	Number	nurse spec_%	nurse spec_Total	Number	MDT meetings_GP_%	meetings_GP_Total	Number	nurse spec%	nurse specTotal	Number
Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2
q42cnumpc	q42cdenominator	q42dnumerator	q42dnumpc	q42ddenominator	q42enumerator	q42enumpc	q42edenominator	q42fnumerator	q42fnumpc	q42fdenominator	q42gnumerator
9.1	11	7	58.3	12	0	0	11	4	36.4	11	2

		MDT						MDT			
MDT	MDT	meetings_Physiothera	MDT	MDT	MDT		MDT	meetings_Psychologist	MDT	MDT	MDT
meetings_Hosp.paed.	meetings_Hosp.paed.	pist_	meetings_Physiothera	meetings_Physiothera	meetings_Dietician_	MDT	meetings_Dietician_To	_	meetings_Psychologist	meetings_Psychologist	meetings_Radiologist_
nurse_%	nurse_Total	Number	pist_%	pist_Total	Number	meetings_Dietician_%	tal	Number	_%	_Total	Number
Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2
q42gnumpc	q42gdenominator	q42hnumerator	q42hnumpc	q42hdenominator	q42inumerator	q42inumpc	q42idenominator	q42jnumerator	q42jnumpc	q42jdenominator	q42knumerator
20	10	4	40	10	0	0	10	2	20	10	0

		MDT									
MDT		meetings_Pharmacistt	MDT	MDT				MDT		MDT	MDT
meetings_Radiologist_	meetings_Radiologist_	_	meetings_Pharmacist_	meetings_Pharmacist_	MDT meetings_Other_	MDT	MDT	frequency_Weekly_	MDT	frequency_Fortnightly_	frequency_Fortnightly_
%	Total	Number	%	Total	Number	meetings_Other_%	meetings_Other_Total	Number	frequency_Weekly_%	Number	%
Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1
q42knumpc	q42kdenominator	q42Inumerator	q42Inumpc	q42ldenominator	q42mnumerator	q42mnumpc	q42mdenominator	q421anumerator1	q421anumpc1	q421anumerator2	q421anumpc2
0	10	1	10	10	4	44.4	9	3	37.5	1	12.5

							Patient and carer engagement					
									Formal survey seeking			
MDT		MDT	MDT	MDT			Formal survey seeking	Formal survey seeking	patient/carer views_	Formal survey seeking	Formal survey seeking	
frequency_Monthly_N	MDT	frequency_Quarterly_	frequency_Quarterly_	frequency_Other_	MDT		patient/carer views_	patient/carer views_	>4 times a	patient/carer views_	patient/carer views_ 3-	
umber	frequency_Monthly_%	Number	%	Number	frequency_Other_%	MDT frequency_Total	Continuous_Number	Continuous_%	year_Number	>4 times a years_%	4 times a year_Number	
Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q4.2.1	Q5.1	Q5.1	Q5.1	Q5.1	Q5.1	
q421anumerator3	q421anumpc3	q421anumerator4	q421anumpc4	q421anumerator5	q421anumpc5	q421adenominator	q51numerator1	q51numpc1	q51numerator2	q51numpc2	q51numerator3	
2	25	1	12.5	1	12.5	8	9	40.9	1	4.5	1	

			Formal survey seeking	Formal survey seeking							
Formal survey seeking	Formal survey seeking	Formal survey seeking	patient/carer views_	patient/carer views_	Formal survey seeking	Formal survey seeking	Formal survey seeking	Strategic group for		Strategic group for	
patient/carer views_ 3-	patient/carer views_ 1-	patient/carer views_1-	Less than once a	Less than once a	patient/carer views_	patient/carer views_	patient/carer views_	paed.	Strategic group for	paed.	Strategic group for
4 times a year_%	2 times a year_Number	2 times a year_%	year_Number	year_%	Never_Number	Never_%	Total	services_No_Number	paed. services_No_%	services_Yes_Number	paed. services_Yes_%
Q5.1	Q5.1	Q5.1	Q5.1	Q5.1	Q5.1	Q5.1	Q5.1	Q5.2	Q5.2	Q5.2	Q5.2
q51numpc3	q51numerator4	q51numpc4	q51numerator5	q51numpc5	q51numerator6	q51numpc6	q51denominator	q52numerator0	q52numpc0	q52numerator1	q52numpc1
4.5	4	18.2	4	18.2	3	13.6	22	2	8.7	19	82.6

Strategic group for	Strategic group for		Patient rep. on		Patient rep. on		Patient rep. on	Patient rep. on		Paediatric patient	Paediatric patient
			· ·	Patient rep. on			strategic group Not	strategic group_Not		access to healthcare	access to healthcare
· –	· –				•	strategic group_Yes_%				records_No_Number	records_No_%
Q5.2	Q5.2	Q5.2	Q5.2.1	Q5.2.1	Q5.2.1	Q5.2.1	Q5.2.1	Q5.2.1	Q5.2.1	Q5.3	Q5.3
q52numerator2	q52numpc2	q52denominator	q521numerator0	q521numpc0	q521numerator1	q521numpc1	q521numerator2	q521numpc2	q521denominator	q53numerator0	q53numpc0
2	8.7	23	12	60	4	20	4	20	20	5	21.7

		Paediatric patient							Link with patient		
Paediatric patient	Paediatric patient	access to healthcare	Paediatric patient	Paediatric patient	Link with patient	Link with patient	Link with patient	Link with patient	support/engagement	Link with patient	Link with patient
access to healthcare	access to healthcare	records_Not	access to healthcare	access to healthcare	support/engagement	support/engagement	support/engagement	support/engagement	group_Not	support/engagement	support/engagement
records_Yes_Number	records_Yes_%	known_Number	records_Not known_%	records_Total	group_No_Number	group_No_%	group_Yes_Number	group_Yes_%	known_Number	group_Not known_%	group_Total
Q5.3	Q5.3	Q5.3	Q5.3	Q5.3	Q5.4						
q53numerator1	q53numpc1	q53numerator2	q53numpc2	q53denominator	q54numerator0	q54numpc0	q54numerator1	q54numpc1	q54numerator2	q54numpc2	q54denominator
15	65.2	3	13.0	23	12	52.2	6	26.1	5	21.7	23

Transitional care	Transitional care												
Transitional care_YP			Transitional care_GP	Transitional care_GP	Transitional care_GP	Transitional care_YP	Transitional care_YP	Transitional care_YP	Transitional care_YP	Transitional care_YP	Transitional care_YP		
	_	_		has same record as YP_%		has transitional care plan_Number		has transitional care plan_Total	has named case worker_Number	has named case worker_%	has named case worker_Total		
q61anumerator	q61anumpc	q61adenominator	q61bnumerator	q61bnumpc	q61bdenominator	q61cnumerator	q61cnumpc	q61cdenominator	q61dnumerator	q61dnumpc	q61ddenominator		
15	65.2	23	14	60.9	23	10	43.5	23	8	34.8	23		