

Paediatric intensive care admissions of preterm children born <32 weeks gestation: a national retrospective cohort study using data linkage

Tim J van Hasselt , ¹ Chris Gale , ² Cheryl Battersby , ² Peter J Davis, ³ Elizabeth Draper, ¹ Sarah E Seaton , ¹ On behalf of the United Kingdom Neonatal Collaborative and the Paediatric Critical Care Society Study Group (PCCS-SG)

► Additional supplemental material is published online only. To view, please visit the journal online (http://dx.doi. org/10.1136/archdischild-2023-325970).

¹Department of Population Health Sciences, University of Leicester, Leicester, UK ²Neonatal Medicine, School of Public Health, Faculty of Medicine, Imperial College London, London, UK ³Paediatric Intensive Care Unit, Bristol Royal Hospital for Children, University Hospitals Bristol NHS Foundation Trust, Bristol, UK

Correspondence to

Dr Tim J van Hasselt,
Department of Population
Health Sciences, University of
Leicester, Leicester LE1 7RH, UK;
t.vanhasselt@nhs.net

Received 15 June 2023 Accepted 19 October 2023

ABSTRACT

Objective Survival of babies born very preterm (<32 weeks gestational age) has increased, although pretermborn children may have ongoing morbidity. We aimed to investigate the risk of admission to paediatric intensive care units (PICUs) of children born very preterm following discharge home from neonatal care.

Design Retrospective cohort study, using data linkage of National Neonatal Research Database and the Paediatric Intensive Care Audit Network datasets.

Setting All neonatal units and PICUs in England and Wales.

Patients Children born very preterm between 1 January 2013 and 31 December 2018 and admitted to neonatal units

Main outcome measures Admission to PICU after discharge home from neonatal care, before 2 years of age.

Results Of the 40 690 children discharged home from neonatal care, there were 2308 children (5.7%) with at least one admission to PICU after discharge. Of these children, there were 1901 whose first PICU admission after discharge was unplanned.

The percentage of children with unplanned PICU admission varied by gestation, from 10.2% of children born <24 weeks to 3.3% born at 31 weeks. Following adjustment, unplanned PICU admission was associated with lower gestation, male sex (adjusted OR (aOR) 0.79), bronchopulmonary dysplasia (aOR 1.37), necrotising enterocolitis requiring surgery (aOR 1.39) and brain injury (aOR 1.42). For each week of increased gestation, the aOR was 0.90.

Conclusions Most babies born <32 weeks and discharged home from neonatal care do not require PICU admission in the first 2 years. The odds of unplanned admissions to PICU were greater in the most preterm and those with significant neonatal morbidity.



© Author(s) (or their employer(s)) 2023. Re-use permitted under CC BY. Published by BMJ.

To cite: van Hasselt TJ, Gale C, Battersby C, et al. Arch Dis Child Fetal Neonatal Ed Epub ahead of print: [please include Day Month Year]. doi:10.1136/archdischild-2023-325970

BACKGROUND

In recent decades, survival following very preterm birth (<32 weeks gestation) has increased, both within the UK¹ and in other high-income countries.² Despite improvements in survival, very pretermborn children experience ongoing morbidity³ and increased frequency of hospitalisation.⁴ A proportion of children born preterm become critically ill after discharge from neonatal care, and require admission to paediatric intensive care units (PICUs),

WHAT IS ALREADY KNOWN ON THIS TOPIC

- ⇒ Children born very preterm (<32 weeks gestational age) are at increased risk of morbidity in childhood.
- ⇒ For such children, survival has increased, particularly for babies born <28 weeks.
- ⇒ It is not known what proportion of very preterm babies discharged from neonatal units will need further intensive care.

WHAT THIS STUDY ADDS

- ⇒ 2308 babies (5.7%) born very preterm and discharged home from neonatal care were subsequently admitted to paediatric intensive care units (PICUs) before the age of 2 years, and the majority of these admissions were unplanned.
- ⇒ The percentage of children with unplanned PICU admission following neonatal discharge varied by gestation, from 10.2% of babies born <24 weeks to 3.3% born at 31 weeks.
- ⇒ Most unplanned PICU admissions of children born very preterm and discharged home occur in the first few months after neonatal discharge.
- Major neonatal morbidities (brain injury, severe necrotising enterocolitis and bronchopulmonary dysplasia) were associated with unplanned PICU admission after neonatal discharge.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

- ⇒ Neonatal healthcare professionals may be able to identify babies at highest risk of being admitted to PICU when planning neonatal discharge.
- ⇒ Appropriate, sensitive and tailored discussion with families may improve understanding of this risk; we plan to work with families to develop infographics to assist these discussions.

particularly due to respiratory viruses in the first years of life. Previous estimates using Office for National Statistics (ONS) data suggest that up to 20%–25% of babies born <25 weeks in the UK may be admitted to PICU in the first 2 years of life, however to our knowledge no study has been published within the UK or internationally using





Original research

data linkage of national neonatal and PICU datasets to examine PICU admissions of preterm-born children.

The risk of requiring PICU admission after neonatal discharge for very preterm-born children is uncertain, as are factors that affect this risk. Understanding these may be helpful for counselling families of very preterm children, and for neonatal discharge planning. We aimed to use a novel linkage of the National Neonatal Research Database (NNRD) and Paediatric Intensive Care Audit Network (PICANet) datasets to investigate the risk of, and risk factors associated with, admission to PICU after neonatal discharge for babies born at 22–31 weeks gestational age.

METHODS

We identified all babies born at <32 weeks gestational age from 1 January 2013 to 31 December 2018, and admitted for neonatal care in England and Wales. All babies born at this gestation should all receive neonatal care and therefore have associated NNRD data. We excluded babies born <22 weeks, and those whose neonatal admissions were recorded as occurring after day 1. Each child was followed up until 2 years of age (a period when children are particularly susceptible to respiratory infections) to investigate if they were admitted to PICUs in England and Wales.

We identified children discharged home from neonatal units, and classified subsequent PICU admission in two ways: (1) 'PICU admission from home' was identified as PICU admission at least 24 hours following neonatal discharge home; (2) 'unplanned PICU admissions', a subset of (1), excluding elective PICU admissions such as those for planned surgery.

Previous PICU admissions during the neonatal stay, with return to the neonatal unit, were not examined, although these children were included within the cohort if they were subsequently discharged home from the neonatal unit.

Information about the care the babies received in the neonatal unit was provided by the NNRD, which captures demographic and clinical data related to neonatal unit admissions, daily clinical care and discharges from all neonatal units in England since 2012 and Wales since 2013.⁷ This was linked with PICU admissions provided by the PICANet, a national audit database of demographic and clinical data collected from every PICU admission across the UK and Ireland, with complete coverage for England and Wales from 2003.⁸ Data are submitted to PICANet within three months of a child's discharge, and subsequently data are subject to validation before any analysis.

Personal identifiers for babies and children (NHS number; date of birth; surname; postcode) were provided by NNRD and PICANet to NHS Digital (now NHS England) who identified children common to both cohorts and provided pseudonymised linked data. Over 99% of children had unique NHS numbers in both datasets, for whom probabilistic linkage was not required. Linked information on deaths in the first 2 years of life was provided by the ONS.

Statistical analysis

We performed descriptive statistics of the cohort, presenting frequencies with percentages for categorical variables, mean with SD for parametric variables and median with IQR for non-parametric variables. We calculated postmenstrual age (PMA) at neonatal discharge and at first unplanned PICU admission.

Our primary analysis was a logistic regression model predicting unplanned PICU admission using characteristics from the neonatal period. Variables were selected for inclusion by a multidisciplinary advisory panel (online supplemental table 1). The selected variables were: gestation, sex, small for gestational age (SGA) and the major neonatal morbidities of bronchopulmonary dysplasia (BPD) requiring oxygen at 36 weeks, severe necrotising enterocolitis (NEC) requiring surgery⁹ and brain injury. Brain injury included grade III/IV intraventricular haemorrhage, periventricular leukomalacia, hypoxic ischaemic encephalopathy, meningitis and seizures after exclusions for congenital or inherited causes. ¹⁰ The major neonatal morbidities also reflected the core outcome set for neonatal research. ¹¹

Implausible birth weights over 3 SD from the median for gestation and sex, 12 or <300 g/>2500 g, were excluded from multivariable models. SGA was defined as birth weight <10th centile for gestation and sex using centiles defined elsewhere. 13 14

Gestational age in completed weeks was modelled linearly, with sensitivity analysis re-fitting the model using categorisation of gestation at birth to investigate the robustness of this approach. The primary model and sensitivity analyses were compared using the Akaike Information Criterion (AIC) for model quality, ¹⁵ Hosmer-Lemeshow ¹⁶ and link tests ¹⁷ for model fit, Brier's score ¹⁸ and c-statistic for predictive ability ¹⁹ and variance inflation factor for multicollinearity. ²⁰

Repeat PICU admission in the first 2 years was explored graphically using a Sankey diagram, created using SankeyMATIC (sankeymatic.com).

RESULTS

There were 46 698 children born <32 weeks between 2013 and 2018 and admitted to neonatal units within England and Wales. After exclusions for neonatal admission after day 1 (n=13), and gestation <22 weeks (n=1), there were 46 684 children (figure 1). In total, 3929 babies died in neonatal care and 2065 were discharged to receive ongoing care in other settings. Of the 40 690 children discharged home from neonatal care, 2308 children (5.7%) had at least one admission to PICU after discharge, comprising 3270 PICU admissions in total.

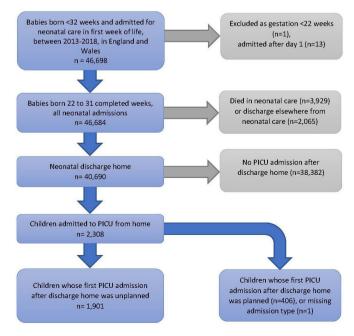


Figure 1 Study flowchart. PICU, paediatric intensive care unit.

Table 1 Birth and neonatal characteristics of cohort: children born <32 weeks between 2013 and 2018 in England and Wales and discharged home from neonatal care

		All children discharged home from neonatal (Children admitted to PICU after discharge home	
		N	%	N	%
Total		40 690		2308	
Sex	Male	22 014	54.1	1384	60.0
	Female	18 653	45.8	921	39.9
	Missing	23	0.1	3	0.1
Gestation	<24	529	1.3	72	3.1
	24	1416	3.5	175	7.6
	25	2034	5.0	205	8.9
	26	2804	6.9	252	10.9
	27	3733	9.2	236	10.2
	28	5103	12.5	287	12.4
	29	6121	15.0	312	13.5
	30	8129	20.0	369	16.0
	31	10 821	26.6	400	17.3
Birth weight (g)	Mean (SD)	1246 (361.2)	-	1131 (377.9)	_
	Small for gestational age (<10th centile)	3246	8.0	220	9.5
	Missing	164	0.4	16	0.7
Multiple birth	Singleton	30 047	73.8	1725	74.7
	Twin	9692	23.8	542	23.5
	Triplet and above	950	2.3	41	1.8
	Missing	1	0.0	0	0
Antenatal steroids	None	2076	5.1	132	5.7
	Complete	28 753	70.7	1608	69.7
	Incomplete	7382	18.1	422	18.3
	Missing	2479	6.1	146	6.3
Mode of delivery	Spontaneous vaginal delivery	14 305	35.2	885	38.3
	Instrumental vaginal delivery	1071	2.6	43	1.9
	Elective caesarean section	2760	6.8	116	5.0
	Emergency caesarean section	20 876	51.3	1157	50.1
	Missing	1678	4.1	107	4.6
BPD	None	28 814	70.8	1229	53.3
	Present	11 659	28.7	1059	45.9
	Missing	217	0.5	20	0.9
Severe NEC	Present	658	1.6	106	4.6
Brain injury	Present	2438	6.0	263	11.4

BPD requiring oxygen at 36 weeks corrected gestational age. Severe NEC requiring surgery.

BPD, bronchopulmonary dysplasia; NEC, necrotising enterocolitis; PICU, paediatric intensive care unit.

All PICU admissions

The subgroup of preterm-born children admitted to PICU had a greater proportion of males (60.0% vs 54.1%), lower birth weight (mean 1131 g vs 1246 g), lower gestational age at birth (median 28 weeks vs 29 weeks), and a greater proportion with neonatal morbidity such as brain injury (11.4% vs 6.0%) compared with the overall cohort of babies discharged home (table 1).

The observed percentage of children who were admitted to PICU after discharge varied by gestational age at birth (table 2), from 13.6% of children born <24 weeks to 3.7% of children born at 31 weeks.

Examining the first PICU admission after neonatal discharge home (figure 2, a Sankey diagram in which the width of the connections represents the number of children), the primary diagnosis on admission was most commonly respiratory disease (n=1436, 62.2%) followed by infection (n=223, 9.7%) and cardiovascular disease (n=196, 8.5%).

Overall, 507 (22.0%) children admitted to PICU from home had at least one further PICU admission before the age of 2, this increased to 31.9% for children born <24 weeks. Respiratory conditions were consistently the most common cause of admission (figure 2).

The observed mortality within PICU for children of all gestations was 2.4% (n=56).

Unplanned PICU admissions

The majority of first admissions to PICU following neonatal discharge were unplanned (n=1901, 82.4%). The percentage of children discharged home who had subsequent unplanned PICU admissions varied by gestation, from 10.2% of children discharged born <24 weeks to 3.3% of those born 31 weeks (table 2). As gestational age at birth increased, neonatal discharge occurred at an earlier PMA, as did the first unplanned PICU admission (table 2). Among unplanned admissions, 431 (22.7%)

Original research

Table 2 Number and percentage of children with one or more PICU admissions after discharge home from the neonatal unit, and PMA at neonatal discharge and at first unplanned PICU admission, by gestational age at birth

Gestational age at	PICU admission from home, including planned and discharged home unplanned (% of children from neonatal care discharged home)		Unplanned PIC from home (% discharged ho	of children	PMA at neonatal discharge (weeks)	PMA at first unplanned PICU admission (weeks)		
birth (weeks)	N	N	%	N %		Median (IQR)	Median (IQR)	
<24	529	72	13.6	54	10.2	41.9 (40.1–44.7)	50.9 (45.4–63.4)	
24	1416	175	12.4	137	9.7	41.1 (39.1–43.6)	52.3 (44.7–68.9)	
25	2034	205	10.1	165	8.1	40.0 (38.3–42.1)	48.1 (42.7–62.3)	
26	2804	252	9.0	196	7.0	39.0 (37.4–41.0)	48.5 (42.6–72.4)	
27	3733	236	6.3	186	5.0	38.1 (36.7–40.0)	45.0 (40.1–66.7)	
28	5103	287	5.6	232	4.5	37.4 (36.1–39.3)	43.4 (38.4–55.6)	
29	6121	312	5.1	264	4.3	36.7 (35.6–38.3)	42.9 (39.0–58.8)	
30	8129	369	4.5	313	3.9	36.3 (35.4–37.6)	40.1 (37.6–50.0)	
31	10 821	400	3.7	354	3.3	35.9 (35.1–37.0)	39.9 (37.6–47.0)	
Total	40 690	2308	5.7	1901	4.7	36.9 (35.7–38.9)	43.9 (39.0–58.6)	
PICU, paediatric intensive care unit; PMA, postmenstrual age.								

children had further PICU admissions (including planned or unplanned), again this was highest in those born at <24 weeks (33.3%).

A total of 40 290 children were included in logistic regression analysis after exclusions for missing variable data (0.98% missing) (table 1). Following adjustment, unplanned PICU admission was associated with lower gestation at birth, male sex, BPD, severe NEC and brain injury (table 3). Of the neonatal morbidities, brain injury had the greatest increase in adjusted OR (aOR 1.42) followed by severe NEC (aOR 1.39) then BPD (aOR 1.37).

We compared model predictions for unplanned PICU admission by gestation and morbidity with overall observed percentages (online supplemental figure 1). Increases in predicted risk of unplanned PICU admission appeared to relate to the total number of neonatal morbidities, increasing to 16.8% (95% CI 12.9% to 20.8%) for children born <24 weeks with BPD, NEC and brain injury.

Results remained consistent in planned sensitivity analyses in which gestational age was modelled as a category; and

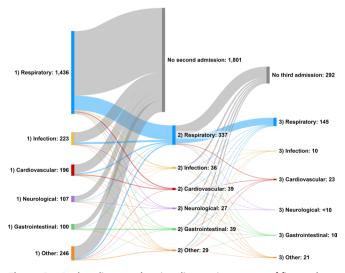


Figure 2 Sankey diagram showing diagnostic category of first and subsequent paediatric intensive care unit admissions after neonatal discharge home of children born <32 weeks.

after exclusion of the 760 (1.9%) children with any congenital anomaly (online supplemental tables 2–4). Values for AIC showed little change, indicating similar model quality. Tests of the primary model showed moderate predictive ability (c-statistic 0.614, Brier score 0.042) and acceptable model fit and collinearity (online supplemental table 5).

DISCUSSION

In this work, we explored the probability of PICU admission for children who were born very preterm and admitted for neonatal care in England and Wales between 2013 and 2018. The observed percentage of children admitted to PICU after neonatal discharge home was highly correlated with gestational age; ranging from 13.6% in those born at <24 weeks gestational age to 3.7% of those born at 31 weeks gestational age. The majority (82%) of these PICU admissions were unplanned, and hence unexpected for the families of these children.

Between a quarter and third of babies born extremely preterm develop BPD,²¹ which can result in respiratory impairment²² and susceptibility to respiratory infections into childhood.⁵ We found that BPD was associated with unplanned PICU admission. Given that the largest percentage of PICU admissions and readmissions were for respiratory disease, the continued high rates of

Table 3 Logistic regression analysis for unplanned PICU admission for children discharged home from neonatal care, using gestation as continuous variable (n=40 290)

Variables		Adjusted OR (95% CI)	P value
Gestation at birth	(weeks)	0.90 (0.88 to 0.92)	<0.001
Sex	Male	Reference	_
	Female	0.79 (0.72 to 0.87)	<0.001
Small for gestational age	Present	1.17 (1.00 to 1.39)	0.057
BPD	Present	1.37 (1.22 to 1.54)	<0.001
Severe NEC	Present	1.39 (1.06 to 1.82)	0.019
Brain injury	Present	1.42 (1.21 to 1.66)	<0.001

BPD requiring oxygen at 36 weeks corrected gestational age. Severe NEC requiring surgery.

BPD, bronchopulmonary dysplasia; NEC, necrotising enterocolitis; PICU, paediatric intensive care unit.

BPD in very preterm survivors is concerning.²³ Unplanned PICU admission was also associated with neonatal brain injury and severe NEC. This is consistent with previous studies that have demonstrated that the number of neonatal morbidities is associated with death and disability in preterm-born children.²⁴ 25

Respiratory admissions to PICU are most commonly due to viral illnesses, such as bronchiolitis. ²⁶ Selective passive immunisation to respiratory syncytial virus (RSV) has been recommended since 2010 in the UK, ²⁷ however our results may prompt neonatal services to consider whether additional measures around neonatal discharge planning may help families prepare for and potentially prevent respiratory viral infections. This may include guidance on how best to access healthcare, ²⁸ open access to paediatric assessment units or neonatal outreach nurse services. The first unplanned PICU admission tended to occur relatively soon after neonatal discharge; healthcare professionals and families may wish to know that the first few months after going home are the highest risk for very preterm-born children.

We observed that a relatively high proportion of children admitted to PICU required subsequent PICU readmission before the age of 2 years (22.0%), and this percentage increased further for the most preterm children. Previous studies have described readmission rates of 15.5% of children readmitted to PICU within 1 year, ²⁹ increasing to 21.7% for children with complex chronic disease, ³⁰ which was similar to that observed in our study.

The mortality rate within PICU was relatively low (2.4%), and similar to the rate previously observed using PICANet data (2.8%) for preterm-born children admitted with respiratory failure.²⁶

The majority of PICU admissions (82% of first admissions) after neonatal discharge were unplanned. Both neonatal and PICU care are associated with symptoms of post-traumatic stress in parents, ^{31–33} and unexpected admissions to another intensive care environment may exacerbate this. Moreover, families may find the PICU environment very different to the neonatal environment they had been used to.³⁴ We are not aware of any literature examining the effects of preparing families for the possibility of PICU admission, however information from our work regarding the potential for PICU admission could be made available to clinicians to share with families who wish to know this.

Strengths and limitations

The major strength of this study is the use of a novel, large, linked national dataset. Given the data quality and completeness of the NNRD and PICANet, and the use of NHS numbers, this should allow high levels of linkage success, 8 35 although we cannot quantify this.

Limited research has explored this area previously. Aggregate data from NNRD and PICANet been used to estimate PICU admission during the neonatal stay for children born very preterm (1.7%–5.5% of children),³⁶ however PICU admissions after neonatal discharge were not examined. A conference abstract reported estimates of extreme preterm-born children requiring PICU admission between birth and 2 years using PICANet and ONS summary data (up to 20%–25% of children born <25 weeks).⁶ However, 30% of PICU admissions had missing data for gestation, leading to a high degree of uncertainty regarding these estimates, in addition this study included children admitted to PICU during their neonatal stay, unlike our study. The use of patient-level data linkage in our study enables us to identify patient journeys of preterm-born children

so provides more robust estimates, and allows adjustment for neonatal factors.

While the datasets we have used cover a wide geographical area, there may be some children whose PICU admission was outside of England or Wales and therefore missed. In addition, a small number of children (0.4%) died outside of neonatal care or PICUs within England and Wales.

While tests of model fit and multicollinearity were satisfactory, our model had only moderate predictive ability for unplanned PICU admission despite a large dataset and selection of significant neonatal morbidity. Therefore, there may have been unmeasured factors which affect the risk of respiratory infections requiring PICU admission, such as the season of discharge, individuals' RSV prophylaxis or smoking within the home.³⁷

Implications for future research

Our dataset does not include babies born since the 2019 changes in UK neonatal management of babies born 22–23 weeks.³⁸ Future work using more recent data may increase understanding of the intensive care needs of this relatively small population, and any effect on overall PICU admissions.

This study did not examine children who were discharged directly from neonatal units to ongoing care in inpatient wards, high dependency units or PICUs (n=2065). These children, particularly those requiring ongoing critical care, are likely to have a greater degree of morbidity, and we intend to study this important group further.

Future research could also examine the impact of early neonatal discharge, and the season of discharge, particularly with regard to subsequent respiratory admissions, and assess the effectiveness of interventions to prevent such illnesses such as RSV prophylaxis for high-risk children.

CONCLUSIONS

The majority of babies born <32 weeks and discharged home from neonatal care do not require PICU admission in the first 2 years of life. However, unexpected admissions to PICU are more common in the most preterm-born children, and especially those with brain injury, severe NEC or BPD. More work is required to understand the impact of morbidity and multimorbidity in the very preterm population.

The main driver of PICU admissions is respiratory illness, mostly occurring in the first few months following neonatal discharge, while a considerable proportion of children require multiple PICU admissions for respiratory disease in the first 2 years.

Our results provide data to support neonatal healthcare professionals in identifying babies with the greatest risk of PICU admission after neonatal discharge. We plan to work with families and healthcare professionals to develop resources to aid discussions around this risk.

Twitter Chris Gale @DrCGale and Cheryl Battersby @DrCBattersby

Acknowledgements The authors thank all UK Neonatal Collaborative neonatal units that agreed to the inclusion of data in the National Neonatal Research Database (NNRD) from their patients to be used in this work. The NNRD is a UK Health Research Authority-approved research database; the Chief Investigator for the NNRD is Professor Neena Modi. The authors would like to thank all paediatric intensive care units in England and Wales who allowed their data to be used in this study. PICANet is commissioned by the Healthcare Quality Improvement Partnership (HQIP) as part of the National Clinical Audit and Patient Outcomes Programme (NCAPOP). HQIP is led by a consortium of the Academy of Medical Royal Colleges, the Royal College of Nursing and National Voices. Its aim is to promote quality improvement in patient outcomes, and in particular, to increase the impact that clinical audit, outcome review programmes and registries have on healthcare quality

Original research

in England and Wales. HQIP holds the contract to commission, manage and develop the NCAPOP, comprising around 40 projects covering care provided to people with a wide range of medical, surgical and mental health conditions. The programme is funded by NHS England, the Welsh Government and, with some individual projects, other devolved administrations and crown dependencies www.hgip.org.uk/nationalprogrammes. Support with the data extraction and linkage was kindly provided by Kayleigh Ougham (NNRD), Lee Norman (PICANet) and NHS Digital. The authors would like to thank the other members of the study advisory group (Professor Jennifer J Kurinczuk, Dr Jonathan Cusack, Dr Patrick Davies, Dr Nicola Mackintosh and Dr Joseph Manning). The authors would also like to acknowledge and thank the support of Bliss, the charity for babies born premature or sick, ADAPT Prem Babies-Leicestershire, the local charity supporting parents and families with premature and poorly babies and The Smallest Things, the volunteer-run charity promoting health and well-being for preterm babies and their families. In particular, the authors would like to acknowledge the children who have contributed data to our research and their families; and the support and input from the families who have contributed so much to this project during the patient and public involvement meetings.

Collaborators Matthew Babirecki (Airedale General Hospital), Rebecca Kettle (Alder Hey), Anand Kamalanathan (Arrowe Park Hospital), Clare Cane (Barnet Hospital), Kavi Aucharaz (Barnsley District General Hospital), Rathod Poorva (Basildon Hospital), Maninder Bal (Basingstoke & North Hampshire Hospital), L M Wong (Bassetlaw District General Hospital), Anita Mittal (Bedford Hospital), Penny Broggio (Birmingham City Hospital), Pinki Surana (Birmingham Heartlands Hospital), Matt Nash (Birmingham Women's Hospital), Sam Wallis (Bradford Royal Infirmary), Ahmed Hassan (Broomfield Hospital, Chelmsford), Karin Schwarz (Calderdale Royal Hospital), Shu-Ling Chuang (Chelsea & Westminster Hospital), Penelope Young (Chesterfield & North Derbyshire Royal Hospital), Ramona Onita (Colchester General Hospital), Mani Kandasamy (Conquest Hospital), Stephen Brearey (Countess of Chester Hospital), Morris/Siramhatia (Croydon University Hospital), Yee Aung (Cumberland Infirmary), Bharath Gowda (Darent Valley Hospital), Mehdi Garbash (Darlington Memorial Hospital), Alex Allwood (Derriford Hospital), Pauline Adiotomre (Diana Princess of Wales Hospital), Nigel Brooke (Doncaster Royal Infirmary), Claire Hollinsworh (Dorset County Hospital), Toria Klutse (East Surrey Hospital), Sonia Spathis (Epsom General Hospital), Sathish Krishnan (Frimley Park Hospital), Samar Sen (Furness General Hospital), Alaa Ghoneem (George Eliot Hospital), Jennifer Holman (Gloucester Royal Hospital), Daniel Dogar (Good Hope Hospital), Girish Gowda (Great Western Hospital), Karen Turnock (Guy's & St Thomas' Hospital), Sobia Balal (Harrogate District Hospital), Cath Seagrave (Hereford County Hospital), Tristan Bate (Hillingdon Hospital), Hilary Dixon (Hinchingbrooke Hospital), Narendra Aladangady (Homerton Hospital), Hassan Gaili (Hull Royal infirmary), Matthew James (Ipswich Hospital), M Lal (James Cook University Hospital), Oluseun Tayo (James Paget Hospital), Abraham Isaac (Kettering General Hospital), Carolina Zorro (Kings College Hospital), Dhaval Dave (King's Mill Hospital), Jonathan Filkin (Kingston Hospital), Savi Sivashankar (Lancashire Women and Newborn Centre), Hannah Shore (Leeds General Infirmary), Jo Behrsin (Leicester General Hospital), Jo Behrsin (Leicester Royal Infirmary), Michael Grosdenier (Leighton Hospital), Ruchika Gupta (Lincoln County Hospital), Ather Ahmed (Lister Hospital), Alison Bedford Russell (Liverpool Women's Hospital), Jennifer Birch (Luton & Dunstable Hospital), Surendran Chandrasekaran (Macclesfield District General Hospital), Ashok Karupaiah (Manor Hospital), Ghada Ramadan (Medway Maritime Hospital), I Misra (Milton Keynes General Hospital), Chris Knight (Musgrove Park Hospital), Richard Heaver (New Cross Hospital), Mohammad Alam (Newham General Hospital), Prakash Thiagaraian (Nobles Hospital), Muthukumar (Norfolk & Norwich University Hospital), Tiziana Fragapane (North Devon District Hospital), Bivan Saha (North Manchester General Hospital), Cheentan Singh (North Middlesex University Hospital), Nick Barnes (Northampton General Hospital), Sangeeta Tiwary (Northumbria Specialist Emergency Care Hospital), Richard Nicholl (Northwick Park Hospital), Dush Batra (Nottingham City Hospital), Dush Batra (Nottingham University Hospital (QMC)), Victoria Nesbitt (Ormskirk District General Hospital), Amit Gupta (Oxford University Hospitals, John Radcliffe Hospital), Katharine McDevitt (Peterborough City Hospital), Ruchika Gupta (Pilgrim Hospital), David Gibson (Pinderfields General Hospital), Peter Mcewan (Poole General Hospital), Sanath Reddy (Princess Alexandra Hospital), Mark Johnson (Princess Anne Hospital), Aesha Mohammedi (Princess Royal Hospital), Patrica Cowley (Princess Royal Hospital), Rashmi Gandhi (Princess Royal University Hospital), Charlotte Groves (Queen Alexandra Hospital), Lidia Tyszcuzk (Queen Charlotte's Hospital), Shilpa Ramesh (Queen Elizabeth Hospital, Gateshead), Salamatu Jalloh (Queen Elizabeth Hospital, King's Lynn), Julia Croft (Queen Elizabeth Hospital, Woolwich), Bushra Abdul-Malik (Queen Elizabeth the Queen Mother Hospital), Dominic Muogbo (Queen's Hospital, Burton on Trent), Ambalika Das (Queen's Hospital, Romford), Khalid Mannan (Queen's Hospital, Romford), Rajiv Chaudhary (Rosie Maternity Hospital, Addenbrookes), Soma Sengupta (Rotherham District General Hospital), Christos Zipitis (Royal Albert Edward Infirmary), Kemy Naidoo (Royal Berkshire Hospital), Archana Mishra (Royal Bolton Hospital), Chris Warren (Royal Cornwall Hospital), Nigel Ruggins (Royal Derby Hospital), Chrissie Oliver (Royal Devon & Exeter Hospital), Lucinda Winckworth (Royal Hampshire County Hospital), Joanne Fedee (Royal Lancaster Infirmary), Anitha Vayalakkad (Royal Oldham Hospital), Richa Gupta (Royal Preston Hospital), Lee Abbott (Royal Stoke

University Hospital), Ben Obi (Royal Surrey County Hospital), Aesha Mohammedi (Royal Sussex County Hospital), Rebecca Winterson (Royal United Hospital), Naveen Athiraman (Royal Victoria Infirmary), Anjali Pektar (Russells Hall Hospital), Jim Baird (Salisbury District Hospital), Adedayo Owoeye (Scarborough General Hospital), Umapathee Maiuran (Scunthorpe General Hospital), Richard Lindley (Sheffield Children's Hospital), Vineet Gupta (Southend Hospital), Faith Emery (Southmead Hospital), Donovan Duffy (St George's Hospital), Salim Yasin (St Helier Hospital), Hannah Shore (St James University Hospital), Akinsola Ogundiya (St Mary's Hospital, IOW), Lidia Tyszcuzk (St Mary's Hospital, London), Ngozi Edi-Osagie (St Mary's Hospital, Manchester), Pamela Cairns (St Michael's Hospital), Vennila Ponnusamy (St Peter's Hospital), Victoria Sharp (St Richard's Hospital), Carrie Heal (Stepping Hill Hospital), Sanjay Salgia (Stoke Mandeville Hospital), Imran Ahmed (Sunderland Royal Hospital), Helen Purves (Tameside General Hospital), Porus Bastani (The Jessop Wing, Sheffield), Eleanor Bond (The Royal Free Hospital), Divyen Shah (The Royal London Hospital—Constance Green), Esther Morris (Torbay Hospital), Se-Yeon Park (Tunbridge Wells Hospital), Giles Kendall (University College Hospital), Puneet Nath (University Hospital Coventry), Igor Fierens (University Hospital Lewisham), Mehdi Garbash (University Hospital of North Durham), Hari Kumar (University Hospital of North Tees), Peter Curtis (Victoria Hospital, Blackpool), Delyth Webb (Warrington Hospital), Bird (Warwick Hospital), Sankara Narayanan (Watford General Hospital), Yee Aung (West Cumberland Hospital), Eleanor Hulse (West Middlesex University Hospital), Tayyaba Aamir (West Suffolk Hospital), Angela Yannoulias (Wexham Park Hospital), Caroline Sullivan (Whipps Cross University Hospital), Ros Garr (Whiston Hospital), Wynne Leith (Whittington Hospital), Shaveta Mulla (William Harvey Hospital). Anna Gregory (Worcestershire Royal Hospital). Edward Yates (Worthing Hospital), Abijeet Godhamgaonkar (Wythenshawe Hospital), Megan Eaton (Yeovil District Hospital), Sundeep Sandhu (York District Hospital), Arun Ramachandran (Singleton Hospital), Abby Parish (Princess of Wales Hospital), Anitha James (The Grange University Hospital), Ian Barnard (Glan Clwyd Hospital), Artur Abelian (Wrexham Maelor Hospital), Shakir Saeed (Ysbyty Gwynedd), Nitin Goel (University Hospital of Wales), David Deekollu (Prince Charles Hospital), Prem Pitchaikani (Glangwili General Hospital).

Contributors TJvH designed the study, and undertook analysis under the supervision of SES, CG and ED. CB and PJD provided clinical interpretation and review of manuscript. All authors contributed to the interpretation, revised the manuscript critically and approved the final version for submission. SES, as supervisor of TJvH, had access to all data and responsibility for the project including decision for publication, and is the guarantor for this paper.

Funding SES (Advanced Fellowship: NIHR300579), CB (Advanced Fellowship: NIHR300617) and TJvH (Doctoral Research Fellowship: NIHR301761) are funded by the National Institute for Health Research (NIHR) for this research project. The views expressed in this publication are those of the authors and not necessarily those of the NIHR, NHS or the UK Department of Health and Social Care.CG is supported by the Medical Research Council through a Clinician Scientist Fellowship (MR/ N008405/1, MR/V036866/1) and this supported his salary over the time spent on this study. He has also received grants and funding from the NIHR, Action Medical Research, Chiesi Pharmaceuticals and the Canadian Institute for Health Research (CIHR)

Disclaimer The views expressed in this publication are those of the authors and not necessarily those of the NIHR, NHS or the UK Department of Health and Social Care.

Competing interests None declared.

Patient consent for publication Not applicable.

Ethics approval Research ethical approval was provided by the East of England committee (reference: 20/EE/0220) and the Confidentiality Advisory Group (20/CAG/0110). This study made use of routinely collected data and therefore no participants were actively recruited.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data may be obtained from a third party and are not publicly available. Data may be obtained from a third party and are not publicly available. PICANet data may be requested from the data controller, the Healthcare Quality Improvement Partnership (HQIP). A Data Access Request Form can be obtained from https://www.hqip.org.uk/national-programmes/accessing-ncapop-data/%23.XQeml_IKhjU.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/licenses/by/4.0/.

ORCID iDs

Tim J van Hasselt http://orcid.org/0000-0001-6157-1083 Chris Gale http://orcid.org/0000-0003-0707-876X Cheryl Battersby http://orcid.org/0000-0002-2898-553X Sarah E Seaton http://orcid.org/0000-0001-8711-4817

REFERENCES

- 1 Santhakumaran S, Statnikov Y, Gray D, et al. Survival of very preterm infants admitted to neonatal care in England 2008-2014: time trends and regional variation. Arch Dis Child Fetal Neonatal Ed 2018;103:F208-15.
- 2 Helenius K, Sjörs G, Shah PS, et al. Survival in very preterm infants: an international comparison of 10 national neonatal networks. *Pediatrics* 2017;140:e20171264.
- 3 Stensvold HJ, Klingenberg C, Stoen R, et al. Neonatal morbidity and 1-year survival of extremely preterm infants. Pediatrics 2017;139:e20161821.
- 4 Coathup V, Boyle E, Carson C, et al. Gestational age and hospital admissions during childhood: population based, record linkage study in England (TIGAR study). BMJ 2020;371:m4075.
- 5 Ghazaly M, Nadel S. Characteristics of children admitted to intensive care with acute bronchiolitis. Eur J Pediatr 2018;177:913–20.
- 6 Yates H. G50 admission of ex-premature babies to PICU in their first two years of life. Archives of Disease in Childhood 2013;98:A27.
- 7 Modi N. Information technology infrastructure, quality improvement and research: the UK national neonatal research database. *Transl Pediatr* 2019;8:193–8.
- 8 Paediatric Intensive Care National Audit Universities of Leicester and Leeds. PICANet: Annual Reporting and Publications. 2023. Available: https://www.picanet. org.uk/annual-reporting-and-publications/2023
- 9 Battersby C, Longford N, Mandalia S, et al. Incidence and enteral feed antecedents of severe neonatal necrotising enterocolitis across neonatal networks in England, 2012-13: a whole-population surveillance study. Lancet Gastroenterol Hepatol 2017;2:43–51.
- 10 Gale C, Statnikov Y, Jawad S, et al. Neonatal brain injuries in England: population-based incidence derived from routinely recorded clinical data held in the National Neonatal Research Database. Arch Dis Child Fetal Neonatal Ed 2018;103:F301–6.
- 11 Webbe JWH, Duffy JMN, Afonso E, et al. Core outcomes in neonatology: development of a core outcome set for neonatal research. Arch Dis Child Fetal Neonatal Ed 2020;105:425–31.
- 12 Manktelow BN, Seaton SE, Field DJ, et al. Population-based estimates of in-unit survival for very preterm infants. Pediatrics 2013;131:e425–32.
- 13 Norris T, Seaton SE, Manktelow BN, et al. Updated birth weight centiles for England and Wales. Arch Dis Child Fetal Neonatal Ed 2018;103:F577–82.
- 14 Cole TJ, Williams AF, Wright CM, et al. Revised birth Centiles for weight, length and head circumference in the UK-WHO growth charts. Ann Hum Biol 2011;38:7–11.
- 15 Burnham KP, Anderson DR. Model selection and multimodel inference: a practical information-theoretic approach. 2.2 ed. New York, NY: New York, NY: Springer New York. 2003.
- 16 Hosmer DW, Lemesbow S. Goodness of fit tests for the multiple logistic regression model. Comm in Stats - Theory & Methods 1980;9:1043–69.

- 17 Pregibon D. Goodness of link tests for generalized linear models. Applied Statistics 1980:29:15.
- 18 Brier GW. Verification of forecasts expressed in terms of probability. Mon Wea Rev 1950;78:1–3.
- 19 Hosmer DW, Lemeshow S, Sturdivant RX. Applied logistic regression. 2013.
- 20 Midi H, Sarkar SK, Rana S. Collinearity diagnostics of binary logistic regression model. Journal of Interdisciplinary Mathematics 2010;13:253–67.
- 21 Shah PS, Lui K, Sjörs G, et al. Neonatal outcomes of very low birth weight and very preterm neonates: an international comparison. J Pediatr 2016;177:144–52.
- 22 Moschino L, Bonadies L, Baraldi E. Lung growth and pulmonary function after prematurity and bronchopulmonary dysplasia. *Pediatr Pulmonol* 2021;56:3499–508.
- 23 Avila-Alvarez A, Zozaya C, Pértega-Diaz S, et al. Temporal trends in respiratory care and bronchopulmonary dysplasia in very preterm infants over a 10-year period in Spain. Arch Dis Child Fetal Neonatal Ed 2022;107:143–9.
- 24 Bassler D, Stoll BJ, Schmidt B, et al. Using a count of neonatal morbidities to predict poor outcome in extremely low birth weight infants: added role of neonatal infection. Pediatrics 2009;123:313–8.
- 25 Farooqi A, Hägglöf B, Sedin G, et al. Impact at age 11 years of major neonatal morbidities in children born extremely preterm. Pediatrics 2011;127:e1247–57.
- 26 O'Donnell DR, Parslow RC, Draper ES. Deprivation, ethnicity and prematurity in infant respiratory failure in PICU in the UK. Acta Paediatr 2010;99:1186–91.
- 27 Joint Committee on Vaccination and Immunisation. Joint Committee on vaccination and Immunisation statement on immunisation for respiratory syncytial virus. 2010. Available: https://webarchive.nationalarchives.gov.uk/ukgwa/20130107105354/http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@ab/documents/digitalasset/dh_120395.pdf2023
- 28 Ingram JC, Powell JE, Blair PS, et al. Does family-centred neonatal discharge planning reduce healthcare usage? A before and after study in South West England. BMJ Open 2016;6:e010752.
- 29 Kane JM, Hall M, Cecil C, et al. Resources and costs associated with repeated admissions to PICUs. Crit Care Explor 2021;3:e0347.
- 30 Chan T, Rodean J, Richardson T, et al. Pediatric critical care resource use by children with medical complexity. J Pediatr 2016;177:197–203.
- 31 Bronner MB, Knoester H, Bos AP, et al. Follow-up after paediatric intensive care treatment: parental posttraumatic stress. Acta Paediatr 2008;97:181–6.
- 32 Feeley N, Zelkowitz P, Cormier C, et al. Posttraumatic stress among mothers of very low birthweight infants at 6 months after discharge from the neonatal intensive care unit. Appl Nurs Res 2011;24:114–7.
- 33 Malouf R, Harrison S, Burton HAL, *et al.* Prevalence of anxiety and post-traumatic stress (PTS) among the parents of babies admitted to neonatal units: a systematic review and meta-analysis. *EClinicalMedicine* 2022;43:101233.
- 34 Evans R, Madsen B. Culture clash: transitioning from the neonatal intensive care unit to the pediatric intensive care unit. *Newborn and Infant Nursing Reviews* 2005:5:188–93.
- 35 Battersby C, Statnikov Y, Santhakumaran S, et al. The United Kingdom national neonatal research database: a validation study. PLoS ONE 2018;13:e0201815.
- 36 Hua X, Petrou S, Coathup V, et al. Gestational age and hospital admission costs from birth to childhood: a population-based record linkage study in England. Arch Dis Child Fetal Neonatal Ed 2023;108:485–91.
- 37 Jones LL, Hashim A, McKeever T, et al. Parental and household smoking and the increased risk of bronchitis, bronchiolitis and other lower respiratory infections in infancy: systematic review and meta-analysis. Respir Res 2011;12:5.
- 38 British Association of Perinatal Medicine. Perinatal management of extreme preterm birth before 27 weeks of gestation, A BAPM framework for practice. 2019. Available: https://www.bapm.org/resources/80-perinatal-management-of-extreme-pretermbirth-before-27-weeks-of-gestation-20192023

Supplementary Table 1 – Members of multi-disciplinary advisory panel for study project

Name	Academic/clinical role	Institution			
Professor Jennifer J Kurinczuk	Professor Of Perinatal	National Perinatal Epidemiology Unit (NPEU),			
Troicssor seminer's Raintezak	Epidemiology & Director	University of Oxford			
Dr Jonathan Cusack	Consultant Neonatologist	University Hospitals of Leicester			
	Consultant in Paediatric				
Dr Patrick Davies	Intensive Care, Honorary	Nottingham University Hospitals, University of			
DI Fattick Davies	Assistant Professor of	Nottingham			
	Paediatrics				
	Clinical Senior Lecturer,	Neonatal Data Analysis Unit - Imperial College			
Dr Cheryl Battersby	Honorary Consultant	London, Chelsea and Westminster NHS			
	Neonatologist	Foundation Trust			
	Consultant Paediatric	Bristol Royal Hospital for Children, University			
Dr Peter Davis	Intensivist	Hospitals Bristol and Weston NHS Foundation			
	ea.svise	Trust			
	Associate Professor in	Department of Population Health Sciences,			
Dr Nicola Mackintosh	Social Science applied to	University of Leicester			
	Health				
	HEE NIHR ICA Clinical				
	Lecturer, Clinical Associate				
	Professor in CYP &	Nottingham Children's Hospital and			
	Families Nursing Charge	Neonatology, Nottingham University Hospitals			
Dr Joseph C Manning MBE	Nurse, Paediatric CCOT,	NHS Trust, School of Health Sciences, University			
	Associate Professor and	of Nottingham			
	Deputy Director - Centre				
	for Children & Young				
	People's Health Research				
	Professor of Neonatal	Faculty of Medicine, School of Public Health,			
Professor Chris Gale	Medicine, Honorary	Imperial College London			
	Consultant Neonatologist				

Supplementary Table 2 - Logistic regression analysis for unplanned PICU admission for children discharged home from neonatal care, using gestation as grouped categorical variable (n=40,290)

		Adjusted odds ratio (95% confidence interval)	р
Gestation at birth (weeks)	<25	Reference	-
	25-27	0.73 (0.61 to 0.87)	<0.001
	28-31	0.53 (0.44 to 0.64)	<0.001
Sex	Male	Reference	-
	Female	0.80 (0.72 to 0.87)	0.096
Small for gestational age	Present	1.15 (0.98 to 1.36)	<0.001
BPD	Present	1.47 (1.31 to 1.65)	0.012
Severe NEC	Present	1.42 (1.08 to 1.86)	<0.001
Brain injury	Present	1.45 (1.23 to 1.70)	<0.001

BPD: bronchopulmonary dysplasia requiring oxygen at 36 weeks corrected gestational age

Severe NEC: necrotising enterocolitis requiring surgery

Supplementary Table 3 - Logistic regression analysis for unplanned PICU admission for children discharged home from neonatal care, using gestation as categorical variable (n=40,290)

		Adjusted odds ratio (95% confidence interval)	р
Contains at high (seeds)	22	2.24 /4 (55) = 2.20)	-0.004
Gestation at birth (weeks)	23	2.31 (1.66 to 3.20)	<0.001
	24	2.29 (1.81 to 2.90)	<0.001
	25	1.96 (1.58 to 2.43)	<0.001
	26	1.80 (1.48 to 2.19)	<0.001
	27	1.31 (1.08 to 1.59)	0.006
	28	1.28 (1.07 to 1.52)	0.006
	29	1.27 (1.08 to 1.50)	0.005
	30	1.17 (1.00 to 1.36)	0.056
	31	Reference	-
Sex	Male	Reference	-
	Female	0.79 (0.72 to 0.87)	<0.001
Small for gestational age	Present	1.18 (1.00 to 1.39)	0.051
BPD	Present	1.37 (1.22 to 1.54)	<0.001
Severe NEC	Present	1.37 (1.04 to 1.80)	0.023
Brain injury	Present	1.41 (1.20 to 1.65)	<0.001

BPD: bronchopulmonary dysplasia requiring oxygen at 36 weeks corrected gestational age

Severe NEC: necrotising enterocolitis requiring surgery

Supplementary Table 4 - Logistic regression analysis for unplanned PICU admission for children discharged home from neonatal care, using gestation as grouped categorical variable, after excluding children with any congenital anomaly (n=39,552)

Variables		Adjusted odds ratio (95% confidence interval)	р
Gestation at birth	(weeks)	0.90 (0.87 to 0.92)	<0.001
Sex	Male	Reference	-
	Female	0.78 (0.71 to 0.86)	<0.001
Small for gestational age	Present	1.15 (0.97 to 1.36)	0.112
BPD	Present	1.35 (1.19 to 1.52)	<0.001
Severe NEC	Present	1.39 (1.05 to 1.83)	0.022
Brain injury	Present	1.43 (1.22 to 1.68)	<0.001

BPD: bronchopulmonary dysplasia requiring oxygen at 36 weeks corrected gestational age

Severe NEC: necrotising enterocolitis requiring surgery

Supplementary Table 5 – Tests of model collinearity, predictive ability, and fit, during logistic regression model development

Variables	N	Mean VIF	Variables with highest VIF	AIC	Brier score	Pseudo R²	C- statisti c	HL test (p)	Link test (p)
Primary model: Gestation (weeks, linear), sex, SGA, BPD, NEC, brain injury	40,290	1.19	Gestation 1.52 BPD 1.47 Brain injury 1.07	14,858	0.042	2.1%	0.614	0.205	0.853
Sensitivity analysis: Gestation (weeks, grouped categorical), sex, SGA, BPD, NEC, brain injury	40,290	2.15	Gestation 28-31wk 5.14 Gestation 25-27wk 4.41 BPD 1.38	14,877	0.042	2.0%	0.609	0.488	0.424
Sensitivity analysis: Gestation (weeks, categorical), sex, SGA, BPD, NEC, brain injury	40,290	1.25	BPD 1.48 Gestation 30wk 1.41 Gestation 28wk 1.36	14,864	0.042	2.1%	0.615	0.432	0.533
Sensitivity analysis (exclusion of any congenital anomaly) Gestation (weeks, linear), sex, SGA, BPD, NEC, brain injury	39,552	1.19	Gestation 1.52 BPD 1.47 Brain injury 1.06	14,440	0.042	2.1%	0.614	0.539	0.781

VIF: Variance inflation factor AIC: Akaike Information Criterion HL test: Hosmer-Lemeshow test SGA: Small for gestational age

BPD: Bronchopulmonary dysplasia requiring oxygen at 36 weeks

NEC: Severe necrotising enterocolitis requiring surgery

Supplementary Figure 1 – Predicted percentages of unplanned PICU admission for children discharged from neonatal care by gestational age at birth, comparing unadjusted observed data with model predictions by neonatal morbidity

