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Comparing routine inpatient data and death records as a means of identifying children and young people with life limiting conditions

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ABSTRACT

Background

Recent estimates of the number of children and young people with life limiting conditions derived from routine inpatient data are higher than earlier estimates using death record data.

Aim

To compare routine inpatient data and death records as means of identifying life limiting conditions in children and young people.

Design

Two national cohorts of children and young people with a life limiting condition (primary cohort from England with a comparator cohort from Scotland) were identified using linked routinely collected healthcare and administrative data.

Participants

37563 children and young people with a life limiting condition in England who died between 1 April 2001 and 30 March 2015 and 2249 children and young people with a life limiting condition in Scotland who died between 1 April 2003 and 30 March 2014.

Results

In England, 16642 (57%) non-neonatal cohort members had a life limiting condition recorded as the underlying cause of death; 3364 (12%) had a life limiting condition -related condition recorded as the underlying cause and 3435 (12%) had life limiting conditions recorded only among contributing causes. 5651 (19%) non-neonates and 3443 (41%) neonates had no indication of a life limiting condition recorded in their death records. Similar results were seen in Scotland (overall, 16% had no indication of life limiting conditions). In both cohorts, the recording of life limiting condition was highest amongst those with haematology or oncology diagnoses and lowest for genitourinary and gastrointestinal diagnoses.

Conclusions

Using death record data alone to identify children and young people with life limiting condition - and therefore those who would require palliative care services - would underestimate the numbers. This underestimation varies by age, deprivation, ethnicity and diagnostic group.

What is already known about the topic?

- Children and young people with life limiting conditions have complex healthcare needs—often with repeated hospital admissions, particularly at end of life.
- Recent estimates of prevalence of life limiting conditions among children and young people using routinely collected inpatient data are much higher than earlier estimates using death records.

What this paper adds?

- Compares identification of life limiting conditions in children and young people using death records and inpatient data in the same population for the first time.
- Shows where the differences occur (by diagnostic group, age group, ethnic group and deprivation).
- Identifies shortcomings in use of death records to identify life limiting conditions in children and young people.

Implications for practice, theory or policy?

- Inpatient-based estimates of life limiting conditions prevalence among children and young people should be used for service planning.
- Epidemiological studies based on life limiting conditions identification from death record data may be biased.
- Use of death records for life limiting conditions identification is particularly limited in countries that only record the underlying cause of death.

Keywords:

Palliative medicine

Child

Inpatients

Cause of death

BACKGROUND

Children and young people with Life-limiting conditions (encompassing both life-limiting conditions that will lead to premature death, e.g. Duchenne muscular dystrophy, and life threatening conditions that may lead to premature death, but may be cured, e.g. cancer) typically have complex healthcare needs, often with repeated hospital admissions, particularly towards the end of life.^{1, 2} Planning paediatric palliative care services, where there is evidence that demand outstrips provision,³ requires accurate estimates of prevalence (in addition to estimates of future survival times).

Recent prevalence estimates³⁻⁵ of children and young people with a life limiting condition in England and Scotland, which used an International Classification of Diseases Version 10⁶ coding framework to identify life limiting conditions within routine inpatient hospital data, were much higher than earlier estimates.⁷⁻⁹ This may be due to previous estimates being based on death record data, about which there are concerns of quality and completeness,⁹ but no work has been previously published comparing the two methods within the same population to quantify these differences. Previous research has found that there are discrepancies between recorded cause of death and conditions recorded during life for a population sample¹⁰ and for children with chronic conditions,¹¹ but these differ from the present study in not focusing on children (first study) and not limiting analyses to life limiting conditions (both studies). In the latter study it was noted that the discrepancies may be due to chronic conditions not being related to the cause of death, but for children with life limiting conditions most deaths are expected to be related to the condition.

This study compares methods of identifying children and young people with a life limiting condition by analysing recorded cause of death for children and young people identified with a life limiting condition from routinely collected English and Scottish inpatient hospital data.

METHODS

Definition of Life-Limiting Conditions

Individuals with a life limiting condition were identified using a refined version⁵ of a previously developed International Classification of Diseases Version 10 coding framework.⁴

Data used and cohort identification

Two national cohorts were identified: a primary cohort from England and a comparison cohort from Scotland (the latter was used to assess whether any differences found were unique to England and is described in the supplement).

For England, linked inpatient Hospital Episode Statistics data and Office for National Statistics death records were used. Data access was granted by the Health and Social Care Information Centre and Office for National Statistics microdata release panel (ref: NIC-379681-D6L7G). Children and young people with a life limiting condition were identified by matching recorded diagnostic codes in inpatient records against the coding framework, for individuals aged 0-25 in the English study period (1 April 2001 to 30 March 2015). The cohort was restricted to individuals with a death record with a date of death in the respective study period.

Data Management

The English datasets were linked by the Health and Social Care Information Centre based on National Health Service number, gender, date of birth and postcode.¹² Management of Scottish datasets is described in the supplement.

Date of birth was assigned as the most commonly recorded date in the inpatient data. Dates of death came from death records. Individuals with invalid dates of death (more than one day before the beginning of an inpatient record) were excluded from the cohort.

Individuals who had died were assigned an age group at death: neonates (0-27 days), postneonatal infants (28-364 days), 1-5, 6-10, 11-15, 16-20, 21-25 and over 25 years. Age at death was determined by subtracting date of birth from date of death. Only month and year of birth was provided, so postneonatal ages of death were approximated setting all birth dates to the 15th day of the month. Neonates were separately identified from the presence of neonate-specific cause of death fields in the death records.

Published populations¹³ and Index of Multiple deprivation 2004¹⁴ rankings¹⁵ for the Lower Super Output Areas provided in the data were used to assign each individual to one of five Index of Multiple deprivation categories, with approximately 20% of the population of England in each category. The Index of Multiple

deprivation 2004 is an area based measure of deprivation under seven domains.¹⁶ Individuals were assigned the last recorded category before death.

Recorded ethnic categories were collapsed to seven groups: White, Indian, Pakistani, Bangladeshi, Black, Mixed and Other. The most commonly recorded ethnicity (from the collapsed groupings) was assigned to each individual.

Life limiting condition diagnoses were categorized into 11 groups based on the main International Classification of Diseases Version 10 chapters: neurology, haematology, oncology, metabolic, respiratory, circulatory, gastrointestinal, genitourinary, perinatal, congenital, and 'other'. Individuals may be assigned more than one diagnostic group if they had more than one life limiting condition recorded in the inpatient hospital data. A primary diagnostic group, the most common diagnostic group across all inpatient records, was assigned to each individual. Where there was more than one most common diagnostic group, later diagnoses were prioritised (diagnoses from the earliest records were progressively removed until the tie was broken).

Analyses

Cause of death

The English death records contained underlying cause of death (the condition that initiated the chain of events that led to death, not necessarily the proximate cause of death) and other causes of death for non-neonates. It was expected that most deaths in the cohort (all of whom were known to have a life limiting condition) would have a life limiting condition as the underlying cause. The proximate cause of death may differ – for example, it may be an infection – but with a life limiting condition as the underlying cause. The proximate cause of death may differ – for example, it may be an infection – but with a life limiting condition as the underlying cause in most cases, making infection either more likely or more severe. A small number of individuals were also expected to die from trauma (e.g. car or other accidents) not related to the life limiting condition. Whether the underlying cause of death was a life limiting condition was checked using the coding framework. If not a life limiting condition, underlying cause was assessed to see whether it was related to a life limiting condition identified in the inpatient data. For example, nonspecific cerebral palsy as cause of death was considered related to quadriplegic cerebral palsy; unspecified congenital malformations of heart to tetralogy of Fallot. Finally, for those with underlying cause neither a life limiting condition nor life limiting condition-related, contributing causes of death were checked against the life limiting condition coding framework. Where life limiting condition was recorded as a

contributing cause, trauma-related underlying causes were determined (all codes starting S; T0; T1; T2; T30; T31; T32; T5; T6; T7; T9; V; W; X; Y1; Y2; Y3) as it was expected that most deaths in the cohort not due to life limiting conditions would be due to trauma.

For English neonate death records, underlying cause of death was not specified. All causes of death were checked for life limiting conditions or being life limiting condition-related (only presence of absence of life limiting conditions among causes of death could be determined, subdivision into underlying, related or contributory causes was not possible for this age group).

The analyses were split by age group (at death), by ethnic group, by deprivation category, by diagnostic category and by financial year of death. Neonates were excluded from analyses by ethnic group, deprivation category, diagnostic category and year as they could not be categorised as having underlying, related or contributory cause of death as a life limiting condition.

Statistical modelling

Predictors of life limiting conditions being present in death records were explored. A binary outcome variable was defined indicating presence of life limiting conditions in a death record, set to 1 if the underlying cause was a life limiting condition or was related to a life limiting condition or a contributing cause of death was a life limiting condition and to 0 if there was no indication of life limiting conditions. Candidate predictor variables were: age group at death, primary diagnostic group, deprivation category and ethnic group. Multivariable logistic regression models were fitted, with candidate predictors added in turn and retained if their odds-ratios were significantly (p < 0.05) different to 1 or inclusion reduced Schwarz's Bayesian Information Criterion for the model by more than 2.^{17, 18} Interactions between deprivation and ethnic group were also considered (using the same inclusion criteria). Models stratified between neonates and non neonates and between oncology and non-oncology primary diagnoses were also developed as it was observed that levels of life limiting condition recording in death records varied greatly between these groups. Individuals with data missing for any included predictors were excluded from the corresponding models.

RESULTS

England

Cohort size

411154 children and young people with a life limiting condition were identified between 1 April 2001 and 31 March 2015 while aged 0-25; 37784 had death records with a date of death in the period. 221 death records (0.6%) were considered invalid as there were one or more inpatient admissions after their recorded date of death and were excluded, leaving 37563 individuals in the final cohort. There were 73 individuals (0.2% of final cohort) with conflicting dates of birth between records: in each case the more commonly recorded date of birth was used. Numbers of deaths in each year and cohort demographics are shown in Table 1.

Missing data

6% of non-neonates had unknown ethnicity, although this figure reduced to 3% for individuals with a date of death on or after 1 April 2009. Including neonates, 10% missed ethnicity information (5% excluding deaths before 1 April 2009, Table 1). 3% of non-neonates had an unknown deprivation category, rising to 10% including neonates. There were no missing data for age group at death or diagnostic category. To test for effects of missing data, a sensitivity analyses was undertaken in the statistical modelling, generating models for the whole time period and also for only 1 April 2009 onwards and with and without neonates.

Cause of death

Among non-neonates, 16642 (57%) had a life limiting condition recorded as underlying cause of death (Table 2); 3364 (12%) had a life limiting condition-related underlying cause and 3435 (12%) had life limiting conditions only among contributing causes, of which 116 had a trauma-related underlying cause (Table 2). 5651 (19%) had no indication of life limiting conditions in their death records. Among neonates, 5028 (59%) had a life limiting condition-related condition among their causes of death; 3443 (41%) had no indication of life limiting conditions among causes of death.

Table 1: Demographics, r	nissing data and cause	e of death by year of de	ath for the English cohort.

		Financial year of dea	th
	2001/02 – 2004/05	2005/06-2009/10	2010/11-2014/15
Deaths in period	9,055	13,984	14,524
Deaths by age group			
Neonate	1,781	3,118	3,572
Post-neonatal infant	1,476	2,282	2,161
1-5 years	1,334	1,793	1,723
6-10 years	713	928	891
11-15 years	882	1,120	886
16-20 years	1,233	1,697	1,465
21-25 years	1,324	2,021	2,009
Over 25 years	312	1,025	1,817
Deaths by ethnic group			
White	5,837	9,505	10,154
Indian	199	359	381
Pakistani	478	995	1,089
Bangladeshi	102	223	195
Black	348	739	851
Mixed	84	300	383
Other	358	641	730
Not known	1,649	1,222	741
Not known (excl nnates)	964	622	299
Deaths by deprivation category			
1 (most deprived)	2,496	3,912	4,157
2	1,716	2,802	2,969
3	1,507	2,123	2,309
4	1,325	1,889	2,003
5 (least deprived)	1,200	1,757	1,704
Not known	811	1,501	1,382
Not known (excl nnates)	221	390	310
Deaths by diagnostic category			
Neurology	1,530	2,698	2,993
Haematology	1,486	2,258	2,417
Oncology	2,670	3,431	3,433
Respiratory	1,263	2,268	2,975
Circulatory	675	1,267	1,242
Gastrointestinal	496	931	1,272
Genitourinary	993	1,907	2,493
Perinatal	1,447	2,673	3,328
Congenital	2,286	3,830	4,190

		112	704	0.45	
Metabolic		413	791	945	
Other		205	336	429	
Life limiting of	condition recording	excluding neonates)			
As underlying	Matched*	4,496 (62%)	6,204 (57%)	5,942 (54%)	
cause	Related*	759 (10%)	1,275 (12%)	1,330 (12%)	
As contributi	ng cause	780 (11%)	1,319 (12%)	1,336 (12%)	
With trauma-related underlying cause		26	46	44	
Not recorded		1,239 (17%)	2,068 (19%)	2,344 (21%)	
All non-neon	ate deaths	7,274	10,866	10,952	

*Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by financial year of death

There was only minor variation in recording of life limiting conditions across financial year of death (Table 1).

The proportion of deaths reporting a life limiting condition as underlying cause varied from 53% to 62% while

the proportion with no indication of life limiting conditions varied from 16% to 22%. There was no clear trend

Ago at death - English data

over time.

Table 2: Recorded cause of death for cohort members, split by age group at death.

	Age at death – English data									
Life limiting recording	condition	Neonate	Post- neonatal Infant	1-5 years	6-10 years	11-15 years	16-20 years	21-25 years	> 25 years	All non- neonates
As	Matched		2463 (42%)	2598 (54%)	1601 (63%)	1865 (65%)	2808 (64%)	3378 (63%)	1929 (61%)	16642 (57%)
underlying cause	Related		1231 (21%)	673 (14%)	261 (10%)	302 (10%)	386 (9%)	300 (6%)	211 (7%)	3364 (12%)
As contribut	a-related		903 (15%) <i><10</i>	631 (13%) <i><10</i>	278 (11%) <10	323 (11%) <i>14</i>	433 (10%) <i>21</i>	522 (10%) <i>43</i>	345 (11%) <i>20</i>	3435 (12%) <i>116</i>
Not recorded		3443 (41%)	1322 (22%)	948 (20%)	392 (15%)	398 (14%)	768 (17%)	1154 (22%)	669 (21%)	5651 (19%)
All deaths in	n age group	8471	5919	4850	2532	2888	4395	5354	3154	29092

* 'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

** Neonate cause of death could not be split between underlying, related or contributory life limiting condition in the English data – all causes of death that were a life limiting condition or related to a life limiting condition were counted

Cause of death by age at death

Neonates were significantly more likely to have no indication of life limiting conditions in death records than non neonates (40.6%, 95% CI 39.6-41.7% compared to 19.4%, 95%CI 19.0-19.9%) (Table 2). Recording of a life limiting condition as underlying cause of death was lowest (2463, 42%) amongst postneonatal infants, but they had the highest percentage of life limiting condition -related deaths (1231, 21%). The youngest and eldest were most likely among the age groups to have no indication of life limiting condition in death records.

Cause of death by ethnic group

Children and young peopleof Bangladeshi or Black ethnicity were less likely than White children and young

people to have a life limiting condition as underlying cause of death (Bangladeshi: 48.5%, 95%CI 43.5-53.4%;

Black: 49.4, 95% CI 46.9-52.0%; White: 58.9%, 95% CI 58.2-59.6%) although they had higher levels of life

limiting condition-related underlying or contributory causes of death (Table 3). Black children and young people were significantly more likely than White children and young people to have no indication of life limiting conditions in their death records (Black: 24.0%, 95% Cl 21.8-26.2%; White: 18.8%, 95% Cl 18.3-19.3%).

Table 3: Recorded cause of death for England cohort members, split by ethnic group.

Life limiting recording	condition	White	Indian	Pakistani	Bangladeshi	Black	Mixed	Other	Unknown	All groups
As	Matched*	12176	415	1104	191	729	299	749	979	16642
underlying		(59%)	(57%)	(53%)	(48%)	(49%)	(54%)	(56%)	(52%)	(57%)
cause	Related*	2325	69	263	50	201	74	170	212	3364
cause		(11%)	(9%)	(13%)	(13%)	(14%)	(13%)	(13%)	(11%)	(12%)
		2284	100	324	74	191	68	186	208	3435
As contribu	ting cause	(11%)	(14%)	(16%)	(19%)	(13%)	(12%)	(14%)	(11%)	(12%)
With traum underlying c		92	≤10	≤10	≤10	≤10	≤10	≤10	≤10	116
Not recorded		3883	144	376	79	354	108	221	486	5651
		(19%)	(20%)	(18%)	(20%)	(24%)	(20%)	(17%)	(26%)	(19%)
All deaths a with ethnic		20668	728	2067	394	1475	549	1326	1885	29092

Ethnic group – English data

*'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by deprivation category

Individuals in the most deprived categories were less likely than those in the least deprived category to have a

life limiting condition recorded as the underlying cause of death (category 1: 53.7%, 95% CI 52.6-54.7%;

category 5: 62.6%, 95% CI 61.1-64.1%) and more likely to have no indication of life limiting conditions

(category 1: 22.3%, 95% CI 21.5-23.2%; category 5: 15.7%, 95% CI 14.5-16.8%) (Table 4).

Table 4: Recorded cause of death for cohort members, excluding English neonates, split by deprivation category of last recorded deprivation score. The categories are population weighted so that 20% of the general population is in each category.

Life limiting condition recording						
		1 (most deprived)	2	3	4	5 (least deprived)
A .c.	Matched*	4639	3436	3012	2576	2494
As underlying cause	watched.	(54%)	(56%)	(59%)	(60%)	(63%)
	-	948	727	605	510	446
	Related*	(11%)	(12%)	(12%)	(12%)	(11%)
A		1125	767	563	449	418
As contributi	ng cause	(13%)	(12%)	(11%)	(11%)	(10%)
	auma-related erlying cause	34	30	21	18	11
		1930	1259	906	737	624
Not recorded		(22%)	(20%)	(18%)	(17%)	(16%)
All deaths ass with IMD sco		8642	6189	5086	4272	3982

*'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by diagnostic group

94% of individuals with an Oncology diagnosis had a life limiting condition as underlying cause of death; only

3% had no indication of life limiting conditions in their death records (Table 5). Only 28% of patients with a

perinatal diagnosis had a life limiting condition as the underlying cause of death, while those with

genitourinary diagnoses were most likely (31%) to have no life limiting condition among any cause of death.

Table 5: Recorded cause of death for cohort members, excluding English neonates, split by diagnostic group.

Life limiting recording	condition	Neurology	Haematology	Oncology	Respiratory	Circulatory	Gastrointestina	Genitourinary	Perinatal	Congenital	Metabolic	Other
As	Matched*	3143	5105	8862	3188	1457	1242	2167	652	2766	1288	700
underlying		(45%)	(83%)	(94%)	(49%)	(52%)	50%)	(43%)	(28%)	(41%)	(65%)	(72%)
		1433	108	22	100	788	443	174	411	656	1638	25
cause	Related*	(20%)	(2%)	(0%)	(12%)	(16%)	(7%)	(8%)	(28%)	(24%)	(5%)	(2%)
A a a a a t uib	•!	946	457	292	1046	431	425	900	400	1088	333	155
As contribu	ting cause	(13%)	(7%)	(3%)	(16%)	(15%)	(17%)	(18%)	(17%)	(16%)	(17%)	(16%)
	uma-related rlying cause	33	<10	11	18	<10	16	32	<10	29	<10	<10
Not recorded		1516	455	299	1448	455	650	1571	609	1334	1983	91
		(21%)	(7%)	(3%)	(22%)	(16%)	(26%)	(31%)	(26%)	(20%)	(13%)	(9%)
All deaths a with diagno		7028	6125	9474	6467	2782	2491	5046	2319	6823	1983	970

Diagnostic group – English data

*'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Multivariable model

The final model (Table 6) showed some differences to the univariable analyses. Neonates were least likely to have a life limiting condition recorded, but 21-25 year olds, and 1-5 and 6-10 year olds were next least likely to have a life limiting condition recorded (odds ratio compared to post neonatal infants: neonate 0.54, 95% CI 0.49-0.60; 1-5 year olds 0.74, 95% CI 0.67-0.83; 6-10 year olds 0.74, 95% CI 0.64-0.86; 21-25 year olds 0.62, 95% CI 0.55-0.69). Variations by ethnic group were also different in the multivariable model, with minority ethnic groups either not significantly different to White children and young people in likelihood of having a life limiting condition recorded or more likely (Pakistani: 1.40, 95% CI 1.25-1.57 times more likely than White children and young people; Other ethnicity 1.25, 95% 1.08-1.43 times more likely than White children and young people). Children and young people in less deprived categories were more likely than children and young people in more deprived categories to have a life limiting condition recorded (odds ratio for least deprived category 1.19, 95% CI 1.08-1.32 compared to most deprived category). However, there were only small differences between the three least deprived categories. Primary diagnostic group showed similar patterns to those seen in the univariable analyses: haematology and oncology diagnoses were most likely to be associated with life limiting condition recording in death records and genitourinary, gastrointestinal and perinatal diagnoses least likely.

There were only minor differences between models stratified between neonates and non-neonates or between individuals with and without oncology as primary diagnostic group. Neither was there significant evidence for interaction between deprivation and ethnic group, so the interaction was not included in the final model. The sensitivity analyses produced models that were not significantly different to the main model (Tables S7 and S8, in the supplement). Table 6: Logistic regression model, using the English data, for the odds ratio of a death record containing any indication of life limiting condition (underlying, related or contributing) depending on age group at death, ethnic group, deprivation category and primary diagnostic group.

	Odds Ratio for life limiting condition in death record		nfidence erval	P value
Age group at death				
Neonate	0.54	0.49	0.60	< 0.01
Post neonatal infant	1 (ref)			
1-5	0.74	0.67	0.83	< 0.01
6-10	0.74	0.64	0.86	< 0.01
11-15	0.98	0.85	1.13	0.77
16-20	0.80	0.71	0.90	< 0.01
21-25	0.62	0.55	0.69	< 0.01
> 25	0.74	0.65	0.84	< 0.01
Ethnic group				
White	1 (ref)			
Indian	1.13	0.94	1.34	0.19
Pakistani	1.40	1.25	1.57	< 0.01
Bangladeshi	1.15	0.91	1.45	0.24
Black	0.99	0.87	1.11	0.81
Mixed	1.10	0.90	1.33	0.35
Other	1.25	1.08	1.43	< 0.01
Last recorded deprivation of	category			
1 (most deprived)	1 (ref)			
2	1.11	1.02	1.20	0.01
3	1.17	1.07	1.28	< 0.01
4	1.21	1.10	1.33	< 0.01
5 (least deprived)	1.19	1.08	1.32	< 0.01
Primary diagnostic group				
Neurology	0.09	0.08	0.10	< 0.01
Haematology	0.06	0.05	0.08	< 0.01
Oncology	1 (ref)			
Respiratory	0.06	0.05	0.07	< 0.01
Circulatory	0.09	0.07	0.11	< 0.01
Gastrointestinal	0.03	0.03	0.04	< 0.01
Genitourinary	0.03	0.02	0.03	< 0.01
Perinatal	0.04	0.03	0.05	< 0.01
Congenital	0.10	0.08	0.11	< 0.01
Metabolic	0.18	0.14	0.22	< 0.01
Other	0.08	0.05	0.12	< 0.01
Model characteristics				
Log likelihood	-13744			
BIC	27778			
Degrees of freedom	28			

Scotland

Full results are presented in the supplement. Of 2249 individuals in the final cohort, 57% had a life limiting condition recorded as underlying cause of death, 14% had a life limiting condition-related underlying cause, 12% had a life limiting condition only among contributing causes and 16% had no indication of life limiting conditions. Under 1 and over 20 year olds were most likely to have no life limiting condition recorded. Those in the most deprived category were more likely (20%) to have no life limiting condition recorded than those in the least deprived category (15%). Only primary diagnostic group showed significant associations with life limiting condition recording in the multivariable model, with life limiting condition recording most likely for individuals in the haematology and oncology primary diagnostic group and least likely for those in the genitourinary group (odds ratio 0.02, 95%CI 0.01-0.05 compared to haematology and oncology).

DISCUSSION

This study has shown that using death certificate data alone to identify the numbers of children and young people with a life limiting condition would have resulted in underestimation of approximately 24% when compared to those identified via inpatient hospital data (for neonates and non-neonates, counting life limiting conditions in all cause of death fields in the English data). The Scottish data provided similar results (see supplement), although recording of life limiting conditions was higher (only 16% had no indication of life limiting conditions in any cause of death field). Readily available death register data in many countries only include the underlying cause of death and using these alone would underestimate the number of children and young people with a life limiting condition further (in the English data, 31% of non-neonate cohort members had neither a life limiting condition nor life limiting condition-related underlying cause of death recorded; for Scotland this figure was 28% for all ages). This may explain the differences in life limiting condition prevalence estimates between studies using routine inpatient hospital data³⁻⁵ and studies using death records.⁷⁻⁹

There are grounds to favour estimates from the inpatient data over estimates from the death records. The number of deaths each year among children identified with a life limiting condition from the inpatient data (e.g. 1766 among 0-14 year olds in 2013) is close to a previous estimate of 50% of child deaths being due to life limiting conditions, ¹⁹ (3631 0-14 year olds died in England in 2013).²⁰ It would be expected that most deaths in individuals identified as having a life limiting condition should either have the life limiting condition as underlying cause or be due to trauma (as broadly defined here), but only 116 of the 3435 deaths with life

limiting conditions recorded as a contributing cause were trauma-related. This suggests there are quality or completeness issues with the death records. The cause of death data are produced by automated analysis of death certificates²¹ and the ONS have noted issues and changes in the way that underlying cause of death is determined in recent years,²² which may result in more maternal conditions as underlying cause of death being recorded as perinatal conditions.²³ This study was concerned with whether death records for children and young people known (from routine hospital data) to have had a life limiting condition would also record the life limiting condition , whether as underlying or any cause of death as this affects the reliability of life limiting condition prevalence estimates based on these data. It is immaterial, when using these data to estimate prevalence, whether any errors in recording are in the manual completion of death certificates or their later automated analysis.

As paediatric palliative care is recommended to start at the point of diagnosis (or recognition) of a life limiting condition rather than just end of life care, only counting the number of children who have died from death records cannot provide a useful estimate of paediatric palliative care need. As treatments improve and survival times for many life limiting conditions increase, death rates are likely to lag behind prevalence increases and underestimate current life limiting condition prevalence.⁹ There is no indication that the gap between life limiting conditions recorded on death records and indicated in inpatient data is decreasing over time, for either the English or Scottish data.

The multivariable model for England is broadly consistent with the univariable descriptive analyses – similar variations are seen for age, deprivation category and diagnostic group. Variation by ethnic group however appears reversed. After controlling for age, primary diagnostic group and deprivation category, individuals of Pakistani and Other ethnicity appear more likely than White individuals to have life limiting condition recorded on their death records; there were no significant differences between White individuals and those from Indian, Bangladeshi, Black or Mixed ethnic groups. This may suggest that the previously observed variation by ethnicity was due to factors such as deprivation or diagnoses (both known to vary by ethnic group²⁴⁻²⁷). The decreased likelihood of life limiting condition recording in death certificates for individuals from more deprived areas may suggest geographical variations in cause of death recording, possibly due to differences in resource provision and quality of recording. Lower likelihood of life limiting condition recording in death records for the very young may be due to greater uncertainty about the exact causes of death; for the older groups, reduced

recording may be due to death following a longer and more complex chain of events from underlying life limiting condition to proximate cause of death, with the underlying life limiting condition not always being recorded. Differences in life limiting condition recording in different diagnostic categories may also be linked to the directness or otherwise of the life limiting condition leading to death. Differences in conditions and the clarity of any link between the life limiting condition and death in those conditions may influence levels of recording among ethnic groups. Further work is needed to investigate these issues. Similar results were seen for the multivariable model for the Scottish data (see Table S6, supplement), although most of the variations – except by primary diagnostic group – were not statistically significant, possibly due to the smaller sample size.

Underestimation of life limiting condition prevalence from death records particularly affects some diagnostic groups (e.g. genitourinary diagnoses), the more deprived and (perhaps as a consequence of diagnostic and deprivation variations) some minority ethnic groups. This has implications for service planning where it could lead to under-provision for these groups or incorrect prioritisation of other groups that appear to have comparatively higher demand and for epidemiological studies, such as those looking at levels of particular life limiting conditions in populations, where bias may be introduced, underestimating prevalence of some conditions more than others. For example, LF's earlier study⁴ on prevalence of life limiting conditions among children and young people in England would be expected to produce different estimates with different relative levels across diagnostic and age groups if using death records. Estimation of demand for service planning should be based on routine hospital data as this is both more up to date and more complete than the death records.

This study used high quality national healthcare data and compared data independently collected in England and Scotland to verify that the variations seen were not unique to one country. The cohorts were identified using an objectively applied coding framework. However, decisions over what constituted a life limiting condition-related underlying cause of death were subjective. The number of cohort members with unknown ethnicity and deprivation category is a concern for the robustness of the analyses with regard to ethnicity and deprivation. The missing data could not be imputed from other fields, but a sensitivity analysis using only data from 2009/10 onwards (with more complete ethnicity data, Table S7, supplement) and excluding neonates (providing more complete ethnicity and deprivation data, Table S8, supplement) supports the findings using the whole study period and all age groups.

CONCLUSION

Using death record data to estimate need for paediatric palliative care services should be undertaken with caution as 19% of non-neonates (31% if using only underlying cause of death) and 41% of neonates identified using the life limiting condition coding framework as having a life limiting condition would have been missed. The most deprived, the youngest and oldest, Black individuals and those with genitourinary, gastrointestinal or perinatal diagnoses were most likely to be missed using death records alone.

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Supplement Part 1: Scotland

METHODS

Data used and cohort identification

For Scotland, the inpatient dataset (SMR01), Scottish Birth Records and General Records Office death record data were used. Data access was approved by the Privacy Advisory Committee (ref: XRB14010). The Scottish data were analysed within the NSS Electronic Data Research and Innovation Service safe haven¹ and results underwent disclosure control.²

Children and young people with a life limiting condition were identified by matching recorded diagnostic codes in inpatient records and birth records against the International Classification of Diseases Version 10 coding framework, for individuals aged 0-25 in the study period (1 April 2003 to 30 March 2014).

Data Management

The Scottish datasets were linked by NHS National Services Scotland using the Community Health Index number.^{3,4}

Date of birth was assigned as the most commonly recorded date in the inpatient and birth data. Dates of death came from death records. Individuals with invalid dates of death (more than one day before the beginning of an inpatient record) were excluded from the cohort.

Individuals who had died were assigned an age group at death: under 1 year old (in Scotland, neonates and post-neonatal infants could not be separated), 1-5, 6-10, 11-15, 16-20, 21-25 and over 25 years. Age at death was determined by subtracting date of birth from date of death.

Provided within the data was a population-weighted Scottish Index of Multiple Deprivation 2009 category, with 10% of the population in each category. These ten categories were collapsed to five for analyses. Individuals were assigned the last deprivation category recorded before death.

The available ethnic categories were collapsed to three, due to small numbers: White, South Asian (Indian, Pakistani, Bangladeshi) and Other. Ethnic group was determined as the most commonly recorded ethnicity (from these collapsed groupings) within the data.

Life limiting condition diagnoses were categorized into 9 groups based on the main diagnostic chapters: neurology, haematology and oncology, metabolic and other, respiratory, circulatory, gastrointestinal, genitourinary, perinatal and congenital (for the Scottish data, small numbers in some groups meant that haematology and oncology were combined into one group, as were metabolic and other). Individuals may be assigned more than one diagnostic group if they had more than one life limiting condition recorded in the inpatient hospital data. A primary diagnostic group, the most common diagnostic group across all inpatient records, was assigned to each individual. Where there was more than one most common diagnostic group, later diagnoses were prioritised (diagnoses from earliest records were progressively removed until the tie was broken).

Analyses

Cause of death

The Scottish death records contained underlying and other causes of death. Whether underlying cause of death was a life limiting condition was checked using the coding framework. If not a life limiting condition, underlying cause was assessed to see whether it was related to a life limiting condition identified in the

inpatient data. For example, nonspecific cerebral palsy as cause of death was considered related to quadriplegic cerebral palsy; unspecified congenital malformations of heart to tetralogy of Fallot. Finally, for those with underlying cause neither a life limiting condition nor life limiting condition-related, contributing causes of death were checked against the life limiting condition coding framework. Where life limiting condition was recorded as a contributing cause, trauma-related underlying causes were determined (all codes starting S; T0; T1; T2; T30; T31; T32; T5; T6; T7; T9; V; W; X; Y1; Y2; Y3).

The analyses were split by age group (at death), by ethnic group, by deprivation category, by diagnostic category and by financial year of death.

Statistical modelling

Predictors of life limiting condition being present in death records were explored. A binary outcome variable was defined indicating presence of life limiting condition in a death record, set to 1 if the underlying cause was a life limiting condition or was related to a life limiting condition or a contributing cause of death was a life limiting condition and to 0 if there was no indication of life limiting conditions. Candidate predictor variables were: age group at death, primary diagnostic group, deprivation category and ethnic group. Multivariable logistic regression models were fitted, with candidate predictors added in turn and retained if their odds-ratios were significantly (p < 0.05) different to 1 or inclusion reduced Schwarz's Bayesian Information Criterion for the model by more than 2.^{5, 6} Interactions between deprivation and ethnic group were also considered (using the same inclusion criteria). Individuals with data missing for any included predictors were excluded from the corresponding models.

RESULTS

Cohort size

In Scotland, 20436 children and young people with a life limiting condition were identified between 1 April 2009 and 31 March 2014 in the Scottish inpatient and birth records. Of these, 2320 had death records in the period. 71 death records were excluded as there were one or more inpatient admissions after the recorded date of death, leaving 2249 individuals in the final cohort. Numbers of deaths in each year and cohort demographics are shown in Table S1.

Missing data

There were large numbers of missing ethnicity data in the Scottish cohort, over half for those dying in the first few years of the study (Table S1). There were no missing data for age group or diagnostic group and missing data for deprivation were at or below 1%.

		Financial year of death	ı	
	2003/04	2004/05	2005/06	
Deaths in year	598	843	808	
Age group				
Under 1 year	186	217	183	
1-5 years	121	97	89	
6-10 years	≤50**	66	58	
11-15 years	68	79	64	
16-20 years	79	155	144	
21-25 years	92	171	154	
Over 25 years	≤10	58	116	
Ethnic group				
White	209	381	580	
South Asian	≤20**	≤20**	≤20**	
Other	≤10	≤10	≤10	
Not known	385	441	188	
Deprivation category				
1 (most deprived)	159	252	208	
2	117	194	192	
3	115	148	152	
4	107	126	133	
5 (least deprived)	96	121	118	
Not known	≤10	≤10	≤10	
Diagnostic category				
Neurology	115	175	187	
Haematology &				
Oncology	193	286	255	
Respiratory	98	196	181	
Circulatory	48	60	64	
Gastrointestinal	30	42	66	
Genitourinary	54	105	132	
Perinatal	93	97	76	
Congenital	151	204	200	
Metabolic & Other	69	85	72	
Life limiting condition				
recording				
As Matched*	362 (61%)	479 (57%)	450 (56%)	
underlying				
cause Related*	93 (16%)	118 (14%)	108 (13%)	
As contributing cause	57 (10%)	113 (13%)	98 (12%)	
With trauma-related				
underlying cause	≤10	≤10	≤10	
Not recorded	86 (14%)	133 (16%)	152 (19%)	
	00 (17/0)	133 (10/0)	152 (1570)	

* Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

** larger values censored to prevent reconstruction of smaller censored values

Cause of death

In Scotland, 1291 cohort members (57%) had a life limiting condition recorded as underlying cause of death (Table S2); 319 (14%) had a life limiting condition-related underlying cause and 268 (12%) had life limiting

conditions only among contributing causes, of which 10 had a trauma-related underlying cause (Table S2). 371 (16%) had no indication of life limiting conditions in their death records.

Cause of death by financial year of death

There was only minor variation in recording of life limiting conditions across financial year of death (Table S1), with no clear trend over time.

Cause of death by age at death

In Scotland, recording of a life limiting condition as underlying cause of death was lowest (239, 41%) amongst under 1 year olds, but they had the highest percentage of life limiting condition-related deaths (150, 26%) (Table S2). Under 1 and over 20 year olds were most likely to have no life limiting condition among any cause of death (under 1s: 19%; 21-25 year olds: 21%; over 25 year olds: 19%).

		Age at death							
Life limiting	; condition	Under 1	1-5	6-10	11-15	16-20	21-25	> 25	All
recording		year	years	years	years	years	years	years	ages
٨c	Matched*	239	185	114	147	236	264	106	1,291
As Matched* underlying	(41%)	(60%)	(67%)	(70%)	(62%)	(63%)	(59%)	(57%)	
cause	Related*	150	51	20	28	36	23	11	319
cause	Relateu	(26%)	(17%)	(12%)	(13%)	(10%)	(6%)	(6%)	(14%)
As contribu	ting course	84	30	15	18	49	44	28	268
AS CONTINU	ting cause	(14%)	(10%)	(9%)	(9%)	(13%)	(11%)	(16%)	(12%)
With traum underlying d	u leiuteu	≤5	≤5	≤5	≤5	≤5	7	≤5	10
Not recorded		113	41	22	18	57	86	34	371
		(19%)	(13%)	(13%)	(9%)	(15%)	(21%)	(19%)	(16%)
All deaths i	n age group	586	307	171	211	378	417	179	2,249

Table S2: Recorded cause of death for cohort members, split by age group at death.

*'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by ethnic group

In Scotland, White individuals were more likely to have life limiting conditions recorded as underlying cause of death (675, 58%) than those in the South Asian (21, 49%) or Other (\leq 10, <50%) groups. Disclosure control prevented release of most values for non-White groups.

Life limiting	condition		Ethnic group		
recording		White	South Asian	Other	
As	Matched*	675	21	≤10	
	watcheu	(58%)	(49%)	510	
underlying		177	<10	<10	
cause	Related*	(15%)	210	≥10	
Ac contribut		134	<10	<10	
As contribut	ling cause	(11%)	210	510	
With trauma	a-related	≤10	<10	≤10	
underlying c	ause	210	210	210	
Not recorde	d	184	<10	<10	
Notrecorded		(16%)	210	210	
All deaths as	ssociated	1170	43	18	
with ethnic	group	11/0	40	10	

Table S3: Recorded cause of death for Scotland cohort members, split by ethnic group.

with ethnic group

*'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by deprivation category

In Scotland, recording of a life limiting condition as the underlying cause of death was less likely for those in the more deprived categories than in the least deprived categories (category 1: 50.4%, 95% CI 46.5-54.3%; category 5: 61.2%, 95% CI 56.0-66.4%).

Table S4: Recorded cause of death for Scotland cohort members, split by deprivation category of last recorded deprivation score. The categories are population weighted so that 20% of the general population is in each category.

Life limiting condition recording		Deprivation category					
		1 (most deprived)	2	3	4	5 (least deprived)	
A. Matahad		312	286	245	236	205	
As	Matched	(50%)	(57%)	(59%)	(64%)	(61%)	
underlying cause		101	70	57	42	49	
	Related	(16%)	(14%)	(14%)	(11%)	(15%)	
As contributing cause With trauma-related underlying cause		81	61	54	39	30	
		(13%)	(12%)	(13%)	(11%)	(9%)	
		≤10	≤10	≤10	≤10	≤10	
Not recorded		125	86	59	49	51	
		(20%)	(17%)	(14%)	(13%)	(15%)	
All deaths ass with deprivat		619	503	415	366	335	

* 'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Cause of death by diagnostic group

In Scotland, 94% of individuals with a haematology or oncology diagnosis had a life limiting condition as underlying cause of death; only 2% had no indication of life limiting conditions in their death records (Table S5). Only 36% of patients with a genitourinary diagnosis had a life limiting condition as the underlying cause of death and 33% no life limiting condition among any cause of death.

Table S5: Recorded cause of death for Scotland cohort members split by diagnostic group.

Life limiting recording	condition	Neurology	Haematology & Oncology	Respiratory	Circulatory	Gastrointestinal	Genitourinary	Perinatal	Congenital	Metabolic & Other
٨٠	Matched*	222	900	265	93	59	104	91	254	183
As underlying	Matcheu	(47%)	(94%)	(56%)	(54%)	(43%)	(36%)	(34%)	(46%)	(79%)
	Related*	106	8	60	25	17	28	72	155	10
cause		(22%)	(1%)	(13%)	(15%)	(12%)	(10%)	(27%)	(28%)	(4%)
Ac contribu	ting course	75	30	75	28	35	63	50	86	17
As contribu	ting cause	(16%)	(3%)	(16%)	(16%)	(25%)	(22%)	(19%)	(15%)	(7%)
	auma-related lerlying cause	≤5	≤5	≤5	≤5	7	≤5	≤5	≤5	≤5
Not record	- d	74	23	75	26	27	96	53	60	21
Not recorded		(16%)	(2%)	(16%)	(15%)	(20%)	(33%)	(20%)	(11%)	(9%)
All deaths a with diagno		477	961	475	172	138	291	266	555	231

Diagnostic group – Scottish data

* 'Matched' underlying causes are those that matched diagnostic codes within the life limiting condition coding framework. 'Related' causes were considered to be related to framework diagnoses present for the individual in the inpatient data.

Multivariable model

The multivariable model for the Scottish data (Table S6) showed no significant variations by age. Neither were there significant variations by ethnic group or deprivation category (although for deprivation category there was an apparent trend towards increased likelihood of life limiting condition recording for individuals from less deprived areas, albeit not significant). Primary diagnostic group showed similar patterns to those seen in the univariable analyses: haematology and oncology diagnoses were most likely to be associated with life limiting condition recording in death records and genitourinary, gastrointestinal and perinatal diagnoses least likely.

 Table S6: Logistic regression model, using the Scottish data, for the odds ratio of a death record containing

 any indication of life limiting conditions (underlying, related or contributing) depending on age group at

death, ethnic group, deprivation category and primary diagnostic group.

	Odds Ratio for life limiting conditions in death record	95% confidence interval		P value
Age group				
Under 1	1 (ref)			
1-5	0.95	0.53	1.68	0.85
6-10	0.84	0.42	1.70	0.63
11-15	1.11	0.52	2.36	0.79
16-20	0.88	0.49	1.57	0.67
21-25	0.58	0.32	1.05	0.07
> 25	0.73	0.36	1.48	0.38
Ethnic group				
White	1 (ref)			
South Asian	1.44	0.57	3.66	0.45
Other	0.63	0.21	1.89	0.41
Last recorded deprivation catego	ory			
1 (most deprived)	1 (ref)			
2	1.06	0.67	1.65	0.81
3	1.15	0.71	1.86	0.58
4	1.27	0.74	2.19	0.39
5 (least deprived)	1.40	0.79	2.49	0.26
Primary diagnostic group				
Neurology	0.09	0.04	0.19	< 0.01
Haematology and oncology	1 (ref)			
Respiratory	0.06	0.03	0.14	< 0.01
Circulatory	0.08	0.03	0.20	< 0.01
Gastrointestinal	0.05	0.02	0.13	< 0.01
Genitourinary	0.02	0.01	0.05	< 0.01
Perinatal	0.06	0.02	0.18	< 0.01
Congenital	0.17	0.08	0.39	< 0.01
Metabolic and Other	0.14	0.05	0.43	< 0.01
Model characteristics				
Log likelihood	-452			
BIC	1053			
Degrees of freedom	21			

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Supplement Part 2: Sensitivity analyses for England

Table S7: Logistic regression model, using the English data, for the odds ratio of a death record containing any indication of life limiting conditions (underlying, related or contributing) depending on age group at death, ethnic group, deprivation category and primary diagnostic group. Sensitivity analysis for deaths from 1 April 2009 onwards when there are fewer missing ethnicity data.

Post neonatal infant 1 (ref) 1-5 0.71 0.60 0.83 < 0.01 6-10 0.67 0.55 0.83 < 0.01 11-15 1.03 0.83 1.29 0.76 16-20 0.69 0.58 0.82 < 0.01 21-25 0.55 0.47 0.64 < 0.01 > 25 0.71 0.60 0.84 < 0.01 21-25 0.55 0.47 0.64 < 0.01 > 25 0.71 0.60 0.84 < 0.01 Pakistani 1.11 0.87 1.41 0.39 Pakistani 1.51 1.28 1.78 < 0.01 Bangladeshi 1.34 0.96 1.86 0.09 Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75 < 0.01 4 1.19 1.04 1.36 0.01		Odds Ratio for life limiting conditions in death record	95% confidence interval		P value
Post neonatal infant 1 (ref) 1-5 0.71 0.60 0.83 < 0.01 6-10 0.67 0.55 0.83 < 0.01 11-15 1.03 0.83 1.29 0.76 16-20 0.69 0.58 0.82 < 0.01 21-25 0.55 0.47 0.64 < 0.01 > 25 0.71 0.60 0.84 < 0.01 21-25 0.55 0.47 0.64 < 0.01 > 25 0.71 0.60 0.84 < 0.01 Pakistani 1.11 0.87 1.41 0.39 Pakistani 1.51 1.28 1.78 < 0.01 Bangladeshi 1.34 0.96 1.86 0.09 Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75<<<0.01 4 1 1.07 1.36 0.01 5	Age group at death				
1-5 0.71 0.60 0.83 <0.01	Neonate	0.55	0.48	0.63	< 0.01
6-10 0.67 0.55 0.83 <0.01	Post neonatal infant	1 (ref)			
11-15 1.03 0.83 1.29 0.76 16-20 0.69 0.58 0.82 < 0.01	1-5	0.71	0.60	0.83	< 0.01
16-20 0.69 0.58 0.82 < 0.01 21-25 0.55 0.47 0.64 < 0.01 > 25 0.71 0.60 0.84 < 0.01 Ethnic groupWhite $1 (ref)$ Indian 1.11 0.87 1.41 0.39 Pakistani 1.51 1.28 1.78 < 0.01 Bangladeshi 1.34 0.96 1.86 0.09 Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75 < 0.01 Last recorded deprivation category1 (most deprived) $1 (ref)$ 2 1.11 1.00 1.24 0.06 3 1.21 1.07 1.36 < 0.01 5 0.04 0.01 < 0.01 Primary diagnostic groupNeurology 0.09 0.07 0.11 < 0.01 Haematology 0.05 0.04 0.07 < 0.01 Gastrointestinal 0.03 0.02 0.04 < 0.01	6-10	0.67	0.55	0.83	< 0.01
21-25 0.55 0.47 0.64 < 0.01	11-15	1.03	0.83	1.29	0.76
> 25 0.71 0.60 0.84 < 0.01 Ethnic group	16-20	0.69	0.58	0.82	< 0.01
Ethnic group White 1 (ref) Indian 1.11 0.87 1.41 0.39 Pakistani 1.51 1.28 1.78 <0.01	21-25	0.55	0.47	0.64	< 0.01
White1 (ref)Indian1.11 0.87 1.41 0.39 Pakistani1.51 1.28 1.78 < 0.01 Bangladeshi 1.34 0.96 1.86 0.09 Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75 < 0.01 Last recorded deprivation category 1 1.11 1.00 1.24 0.06 3 1.21 1.07 1.36 < 0.01 4 1.19 1.04 1.36 0.01 5 (least deprived) 1.20 1.05 1.38 < 0.01 Primary diagnostic groupNeurology 0.09 0.07 0.11 < 0.01 Haematology 0.05 0.04 0.07 < 0.01 Circulatory 0.09 0.07 0.12 < 0.01 Gastrointestinal 0.03 0.02 0.04 < 0.01	> 25	0.71	0.60	0.84	< 0.01
Indian1.110.871.410.39Pakistani1.511.281.78< 0.01	Ethnic group				
Pakistani 1.51 1.28 1.78 < 0.01	White	1 (ref)			
Bangladeshi 1.34 0.96 1.86 0.09 Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75 < 0.01	Indian	1.11	0.87	1.41	0.39
Black 1.04 0.88 1.23 0.64 Mixed 1.13 0.89 1.45 0.31 Other 1.44 1.18 1.75 < 0.01	Pakistani	1.51	1.28	1.78	< 0.01
Mixed1.130.891.450.31Other1.441.181.75< 0.01Last recorded deprivation category1(ref)1 (most deprived)1 (ref)21.111.001.240.0631.211.071.36< 0.01	Bangladeshi	1.34	0.96	1.86	0.09
Other 1.44 1.18 1.75 < 0.01 Last recorded deprivation category 1 (ref) <	Black	1.04	0.88	1.23	0.64
Last recorded deprivation category 1 (most deprived) 1 (ref) 2 1.11 1.00 1.24 0.06 3 1.21 1.07 1.36 < 0.01	Mixed	1.13	0.89	1.45	0.31
1 (most deprived) 1 (ref) 2 1.11 1.00 1.24 0.06 3 1.21 1.07 1.36 < 0.01	Other	1.44	1.18	1.75	< 0.01
1.11 1.00 1.24 0.06 3 1.21 1.07 1.36 < 0.01	Last recorded deprivatio	n category			
3 1.21 1.07 1.36 < 0.01	1 (most deprived)	1 (ref)			
41.191.041.360.015 (least deprived)1.201.051.38< 0.01	2	1.11	1.00	1.24	0.06
5 (least deprived) 1.20 1.05 1.38 < 0.01	3	1.21	1.07	1.36	< 0.01
Primary diagnostic group Neurology 0.09 0.07 0.11 < 0.01	4	1.19	1.04	1.36	0.01
Neurology 0.09 0.07 0.11 < 0.01 Haematology 0.06 0.05 0.08 < 0.01	5 (least deprived)	1.20	1.05	1.38	< 0.01
Haematology 0.06 0.05 0.08 < 0.01 Oncology 1 (ref) <t< td=""><td>Primary diagnostic group</td><td>)</td><td></td><td></td><td></td></t<>	Primary diagnostic group)			
Oncology 1 (ref) Respiratory 0.05 0.04 0.07 < 0.01	Neurology	0.09	0.07	0.11	< 0.01
Respiratory0.050.040.07< 0.01Circulatory0.090.070.12< 0.01	Haematology	0.06	0.05	0.08	< 0.01
Circulatory 0.09 0.07 0.12 < 0.01 Gastrointestinal 0.03 0.02 0.04 < 0.01	Oncology	1 (ref)			
Gastrointestinal 0.03 0.02 0.04 < 0.01 Genitourinary 0.03 0.02 0.03 < 0.01	Respiratory	0.05	0.04	0.07	< 0.01
Genitourinary 0.03 0.02 0.03 < 0.01	Circulatory	0.09	0.07	0.12	< 0.01
•	Gastrointestinal	0.03	0.02	0.04	< 0.01
Perinatal 0.05 0.04 0.06 < 0.01	Genitourinary	0.03	0.02	0.03	< 0.01
	Perinatal	0.05	0.04	0.06	< 0.01
Congenital 0.10 0.08 0.12 < 0.01	Congenital	0.10	0.08	0.12	< 0.01
Metabolic 0.18 0.13 0.24 < 0.01	Metabolic	0.18	0.13	0.24	< 0.01
Other 0.06 0.03 0.12 < 0.01	Other	0.06	0.03	0.12	< 0.01
Model characteristics	Model characteristics				
Log likelihood -7048	Log likelihood	-7048			
BIC 14366		14366			
Degrees of freedom 28	Degrees of freedom	28			

Table S8: Logistic regression model, using the English data, for the odds ratio of a death record containing any indication of life limiting condition(underlying, related or contributing) depending on age group at death, ethnic group, deprivation category and primary diagnostic group. Sensitivity analysis excluding neonates.

	Odds Ratio for a life limiting condition in death record	95% confidence interval		P value	
Age group at death					
Post neonatal infant	1 (ref)				
1-5	0.76	0.68	0.85	< 0.01	
6-10	0.76	0.66	0.88	< 0.01	
11-15	1.00	0.86	1.16	0.98	
16-20	0.82	0.72	0.93	< 0.01	
21-25	0.63	0.56	0.71	< 0.01	
> 25	0.76	0.67	0.87	< 0.01	
Ethnic group					
White	1 (ref)				
Indian	1.07	0.88	1.31	0.49	
Pakistani	1.36	1.19	1.54	< 0.01	
Bangladeshi	1.14	0.87	1.49	0.34	
Black	0.92	0.81	1.06	0.27	
Mixed	1.10	0.87	1.39	0.42	
Other	1.20	1.02	1.41	0.03	
Last recorded deprivation (1 (most deprived)	category 1 (ref)				
2	1.09	1.00	1.20	0.06	
3	1.20	1.08	1.32	< 0.01	
4	1.23	1.10	1.37	< 0.01	
5 (least deprived)	1.27	1.13	1.42	< 0.01	
Primary diagnostic group					
Neurology	0.09	0.08	0.10	< 0.01	
Haematology	0.06	0.05	0.08	< 0.01	
Oncology	1 (ref)				
Respiratory	0.06	0.05	0.07	< 0.01	
Circulatory	0.10	0.08	0.12	< 0.01	
Gastrointestinal	0.03	0.02	0.04	< 0.01	
Genitourinary	0.03	0.02	0.03	< 0.01	
Perinatal	0.05	0.04	0.06	< 0.01	
Congenital	0.10	0.08	0.11	< 0.01	
Metabolic	0.19	0.15	0.23	< 0.01	
Other	0.08	0.05	0.13	< 0.01	
Model characteristics Log likelihood	-10801				
BIC	21876				
Degrees of freedom	27				