Original Article

Severe maternal morbidity in Scotland

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Summary

Using a cohort study design, we analysed 17 diagnoses and 9 interventions (including critical care admission) as a composite measure of severe maternal morbidity for pregnancies recorded over 14 years in Scotland. There were 762,918 pregnancies, of which 7947 (10 in 1000 pregnancies) recorded 9345 severe maternal morbidity events, 2802 episodes of puerperal sepsis being the most common (30%). Severe maternal morbidity incidence increased from 9 in 1000 pregnancies in 2012 to 17 in 1000 pregnancies in 2018, due in part to puerperal sepsis recording. The odds ratio (95%CI) for severe maternal morbidity was higher for: older women, for instance 1.22 (1.13-1.33) for women aged 35–39 years and 1.44 (1.27-1.63) for women aged > 40 years compared with those aged 25-29 years; obese women, for instance 1.13 (1.06-1.21) for BMI 30-40 kg.m⁻² and 1.32 (1.15-1.51) for BMI > 40 kg.m⁻² compared with BMI 18.5–24.9 kg.m⁻²; multiple pregnancy, 2.39 (2.09–2.74); and previous caesarean delivery, 1.52 (1.40–1.65). The median (IQR [range]) hospital stay was 3 (2–5 [1–8]) days with severe maternal morbidity and 2 (1-3 [1-5]) days without. Forty-one women died during pregnancy or up to 42 days after delivery, representing mortality rates per 100,000 pregnancies of about 365 with severe maternal morbidity and 1.6 without. There were 1449 women admitted to critical care, 807 (58%) for mechanical ventilation or support of at least two organs. We recorded an incidence of severe maternal morbidity higher than previously published, possibly because sepsis was coded inaccurately in our databases. Further research may determine the value of this composite measure of severe maternal morbidity.

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Introduction

The number of women dying during pregnancy and childbirth in the UK is at an all-time low [1]. Consequently, enquiries into maternal death have limited power to

improve maternal care, whereas measures of severe – but usually non-lethal – events during pregnancy and the puerperium may continue to inform care in the UK [2–4]. A composite of individual morbidities is up to 100 times more frequent than maternal death and therefore affords opportunities to intervene to reduce morbidity and mortality [5, 6].

The rate of severe maternal morbidity may be increasing in some high-income countries, which is associated with modifiable factors such as obesity, and other factors that, whilst not modifiable, might trigger intervention, such as age and previous caesarean delivery [7, 8]. The increasing prevalence of these factors in UK mothers might reverse the historical reductions in mortality and will increase healthcare during pregnancy and childbirth [9].

The UK Obstetric Surveillance System (UKOSS) reports specific conditions or healthcare themes of interest, including morbidities [10]. In contrast, semi-automated collection of healthcare data may provide the opportunity to monitor rates of maternal morbidities routinely. Linked electronic data resources could help identify women more likely to experience severe maternal morbidity and may improve our understanding of how severe maternal morbidities affect women, children and healthcare services.

Accordingly, we aimed to use routinely collected data sources to generate a composite measure of severe maternal morbidity, report its incidence and the variables associated with its occurrence.

Methods

Using a cohort design, we studied women with a live birth, stillbirth or late second trimester loss (defined as 20 weeks to `23 weeks and 6 days' gestation) from 1 January 2005 to 31 December 2018, using data from the Scottish Morbidity Records, the National Records of Scotland and the Scottish Intensive Care Society Audit Group (please see online Supporting Information, Appendices S1 and S2 for a detailed account of methods)[11–14].

The primary outcome was any severe maternal morbidity recorded from conception to 42 days postpartum, defined by English Maternal Morbidity Indicator codes for 17 diagnoses, for instance puerperal sepsis or status epilepticus, and nine interventions, for instance repair of bladder or evacuation of haematoma, with corresponding disease (ICD-10) and procedural (OPCS) codes (online Supporting Information, Table S1) [15]. We added intensive care unit admission as an indicator for severe maternal morbidity and, in combination with other procedural codes related to managing haemorrhage, we consequently excluded the ICD-10 code for major obstetric haemorrhage (O72), which is substantially overestimated using ICD coding [16]. We report the first event for women who experienced multiple morbidities.

We analysed duration of hospital admission and maternal mortality as well as the duration and level of critical care, mechanical ventilation, renal replacement therapy and cardiovascular support (online Supporting Information, Appendices S1 and S2)[17].

We analysed the associations of severe maternal morbidity with sociodemographic measures, prepregnancy maternal health status, obstetric history and current pregnancy information (online Supporting Information, Appendices S1 and S2). We searched for ICDcodes recorded in the five years preceding the estimated date of conception to define comorbidities (online Supporting Information, Appendix S1 and Table S2) [18–20]. As a sensitivity analysis, the comorbidity variable was replaced with a new count which included those comorbidities coded before and during a pregnancy.

We used Stata version 14.1 for analyses (Stata Corp, College Station, TX, USA). We used logistic regression to analyse the associations of severe maternal morbidity with variables present before hospital admission or early in pregnancy (model 1), and variables present later in pregnancy and during delivery (model 2) (online Supporting Information, Appendices S1 and S2). We used robust variance estimation to account for clustering due to multiple pregnancies. We used chi-squared and Mann– Whitney-U tests for categorical and continuous variables, respectively. Patient information was pseudonymised and all analyses were conducted within NHS Scotland's safe haven environment. Access to the data was granted following approval by the Public Benefit and Privacy Panel.

Results

We analysed 762,918 pregnancies, during which 9345 severe maternal morbidity events were recorded relating to 7947 pregnancies (10.4 in 1000 pregnancies) (Fig. 1,



Figure 1 Cohort flow diagram.

	Severe maternal morbidity		
	No	Yes	
Characteristic	n = /54,9/1	n = /94/	p value
Maternal age; y	29 (25–34)	30 (25–34)	< 0.001
< 20	42,888 (6%)	451 (6%)	
20-24	132,218 (18%)	1279(16%)	
25-29	204,106(27%)	2013 (25%)	
30–34	222,061 (29%)	2316 (29%)	
35–39	126,108(17%)	1479 (19%)	
> 39	27,590(4%)	409(5%)	
Mother's region of birth		(() () () () () () () () () (0.004
British Isles	645,889(86%)	6621 (83%)	< 0.001
Rest of Europe	45,946(6%)	4/6(6%)	
Middle East or Asia	29,610(4%)	408 (5%)	
Africa	15,443 (2%)	229 (3%)	
North America	4781 (1%)	58 (1%)	
Missing	8660(1%)	96(1%)	
Oceania	2662	32	
South America	1980	27	
Scottish index of multiple deprivation			
1 (most deprived)	104,057 (14%)	1058 (13%)	
2	91,041 (12%)	982 (12%)	
3	81,898(11%)	957 (12%)	
4	76,649(10%)	833 (10%)	
5	73,976(10%)	775 (10%)	
Missing	935	19	
Urban rural indicator			
Large urban area	309,478(41%)	3223 (41%)	0.014
Other urban area	235,541 (31%)	2565 (32%)	
Accessible small town	61,754(8%)	658(8%)	
Remote small towns	25,956 (3%)	267 (3%)	
Accessible rural	74,770(10%)	743 (9%)	
Remote rural	35,093 (5%)	332 (4%)	
Missing	12,379 (2%)	159 (2%)	
Sole parental registration			
No	705,309(93%)	7303 (92%)	< 0.001
Yes	37,961 (5%)	505 (6%)	
Missing	11,701 (2%)	139(2%)	
Pre-pregnancy BMI; kg.m ⁻²	25.0 (22.1–29.1)	25.5 (22.3–30.0)	< 0.001
Smoking status			
None	482,925(64%)	5028(63%)	< 0.001
Current	135,757 (18%)	1426(18%)	
Former	84,042 (11%)	1011 (13%)	
Missing	52,247 (7%)	482 (6%)	

Table 1Maternal characteristics and pregnancy outcomes, stratified by whether a severe maternal morbidity was recorded for
the pregnancy. Values are median (IQR) or number (proportion).

(continued)

Table 1 (continued)

No Yes Characteristic n = 754,971 n = 7947 pvalue Diabetes
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Diabetes No 659,140(87%) 6985(88%) < 0.01
No 659,140(87%) 6985(88%) < 0.001 Gestational 12,324(2%) 208(3%) Pre-gestational 4374(1%) 106(1%) Hypertension before admission < 0.001
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Pre-gestational 4374 (1%) 106 (1%) Hypertension before admission < 0.001
Hypertension before admission < 0.001 No 733,485 (97%) 7701 (97%) Yes 1782 43 Previous pregnancy loss < 0.001
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2007 55,510(99.3%) 403(0.7%)
2008 57,457 (99.2%) 458 (0.8%)
2009 56,610(99.2%) 467(0.8%)
2010 56,545 (99.1%) 512 (0.9%)
2011 56,268 (99.1%) 503 (0.9%)
2012 55,598(99.1%) 506(0.9%)
2013 53,685 (98.9%) 591 (1.1%)
2014 54,339 (98.8%) 636 (1.2%)
2015 52,702 (98.7%) 718 (1.3%)
2016 52,082 (98.7%) 705 (1.3%)
2017 50,563 (98.5%) 794 (1.5%)
2018 49,105(98.3%) 848(1.7%)
Multiple gestation
No 741,507 (98%) 7642 (96%) < 0.001
Yes 11,301(2%) 270(3%)
Missing 2163 35
Estimated gestation (Weeks) 40 (38–40) 39 (38–40) < 0.001
Very preterm (< 28 weeks) 5152 (1%) 175 (2%)
Preterm (28 < 37 weeks) 49,355 (7%) 1308 (16%)

(continued)

Table 1 (continued)

	Severe maternal morbidity			
	Νο	Yes		
Characteristic	n = 754,971	n = 7947	p value	
Term(37 < 41 weeks)	528,812(70%)	4976 (63%)		
Overdue (> 41 weeks)	171,652(23%)	1488(19%)		
Mode of delivery				
Unassisted vaginal	445,102(59%)	2695 (34%)	< 0.001	
Emergency caesarean section	118,652(16%)	3051 (38%)		
Assisted vaginal	95,597 (13%)	1034(13%)		
Elective caesarean section	93,392 (12%)	1131(14%)		
Missing	2228	36		
Antenatal steroids				
No	664,385(88%)	6527 (82%)	< 0.001	
Yes	26,973 (4%)	810(10%)		
Missing	63,613(8%)	610(8%)		
Analgesia during pregnancy or labour				
Yes	651,916(86%)	7165 (90%)	< 0.001	
No	33,166(4%)	184 (2%)		
Missing	69,889(9%)	598 (8%)		
Induction of labour				
No	547,456(73%)	5426 (68%)	< 0.001	
Yes	199,574(26%)	2409 (30%)		
Missing	7941 (1%)	112(1%)		
Obstetric admissions before delivery				
0	466,734(62%)	4260 (54%)	< 0.001	
1	141,192(19%)	1598 (20%)		
> 1	142,400(19%)	2011 (25%)		
Missing	4645(1%)	78 (1%)		
Stay after delivery; days	2(1–3)	3 (2–5)	< 0.001	
Maternal death to 42 days	12 (1.6 per 100,000)	29 (365 per 100,000)	< 0.001	
Outcome of birth				
Live birth	740,537 (98%)	7676(7%)	< 0.001	
Stillbirth	3001 (0.4%)	138 (2%)		
Late second trimester loss	1232 (0.2%)	22 (0.3%)		
Missing	10,201 (1%)	111(1%)		

Tables 1 and 2 and online Supporting Information, Tables S2–S4). One severe maternal morbidity was recorded for 6891 (87%) women, two for 806 (10%) women and three or more for 250 women (3%). Puerperal sepsis or admission to critical care were the most common indicators of severe maternal morbidity (Fig. 2 and online Supporting Information, Figures S1 and S2).

Severe maternal morbidity increased in the years after 2012 due to puerperal sepsis, from 9 in 1000 pregnancies in 2012 to 17 in 1000 pregnancies in 2018 (online Supporting Information, Figure S3). Severe maternal morbidity was

independently associated with maternal age; BMI; preexisting morbidity; previous smoking; previous caesarean section; multiple pregnancy; and maternal birth in Africa or the Middle East (model 1, online Supporting Information Table S5; Fig. 3a–c and online Supporting Information, Figures S4–S6).

Hospital admission during the current pregnancy was associated with severe maternal morbidity, as were antenatal steroids, early or late delivery, induction of labour, assisted delivery (in particular emergency caesarean section) and analgesia (model 2, online Supporting Information, **Table 2** Severe maternal morbidities recorded relative to the day of labour and birth. We tabulated only the first severe maternal morbidity recorded for a woman. The frequency of the morbidities sums to more than the number of pregnancies (n) as a woman may have had multiple different severe maternal morbidity events during a time period. Values are number (proportion) or number.

	Before, during or after labour and birth				
Morbidity	Before n = 2839	During n = 2226	After n = 2882	Total n = 7947	p value
Sepsis	728 (26%)	520(19%)	1554 (55%)	2802	< 0.001
ICU admission	389 (27%)	688(47%)	372 (26%)	1449	< 0.001
Caesarean dehiscence	129 (26%)	178(36%)	188 (38%)	495	< 0.001
Curettage with anaesthetic	87(18%)	91 (19%)	307 (63%)	485	< 0.001
Eclampsia	257 (54%)	146(30%)	76(16%)	479	< 0.001
Cardiac event	173 (38%)	138(30%)	146 (32%)	457	< 0.001
Acute abdomen	279(69%)	7 (2%)	116 (29%)	402	< 0.001
Embolism	205 (56%)	60(16%)	102 (28%)	367	< 0.001
Evacuation of haematoma	121 (36%)	138(41%)	74 (22%)	333	< 0.001
Acute renal failure	160(51%)	90 (29%)	65 (21%)	315	< 0.001
Hysterectomy	107 (43%)	108(43%)	35(14%)	250	< 0.001
Assisted ventilation	86(37%)	105 (45%)	40(17%)	231	< 0.001
Cystotomy or repair	*	129(57%)	*	227	< 0.001
Shock	87(41%)	107 (50%)	18(8%)	212	< 0.001
Uterine rupture	*	119(60%)	*	200	< 0.001
Anaesthetic complication	62(48%)	63 (48%)	5 (4%)	130	< 0.001
Stroke	59(51%)	16(14%)	41 (35%)	116	< 0.001
Intestine repair	41 (43%)	15(16%)	39(41%)	95	< 0.001
Status asthmaticus	71(84%)	8(9%)	6(7%)	85	< 0.001
Uterine artery intervention	25	11	12	48	< 0.001
Acute psychosis	17	6	8	31	< 0.001
Cerebral oedema or coma	22	*	*	31	< 0.001
DIC	*	15	*	24	< 0.001
Sickle cell crisis	19	*	*	23	< 0.001
Cerebral vein thrombosis	*	*	10	20	< 0.001
Dialysis	7	6	6	19	< 0.001
Status epilepticus	16	*	*	19	< 0.001
Total number of events				9345	

Cardiac event, arrest or failure or infarction; DIC, disseminated intravascular coagulopathy. *Output suppressed due to disclosure risk.

Table S5). The associations in model 2 should be interpreted with caution as they may have been the consequence of morbidity, rather than its cause. The inclusion in the model of comprehidition coded before and during programs, had little

comorbidities coded before and during pregnancy had little effect on these associations in sensitivity analyses (online Supporting Information, Tables S6 and S7).

Women with severe maternal morbidity were significantly more likely to require a longer hospital stay following delivery (11.1% vs. 3.0% staying > 7 days, p < 0.001) and have a stillbirth (1.7% vs. 0.4%, p < 0.001) (Table 1). More women with severe morbidity died during pregnancy or up to 42 days postpartum than without, with

mortality rates of 36.5 in 10,000 vs. 0.16 in 10,000, p < 0.001.

Online Supporting Information (Table S8) details the care of the 1449 women (0.2%) admitted to intensive care, of whom 807 (56%) were ventilated or had more than one organ supported.

Discussion

We found that severe maternal morbidity was recorded for about 10 in 1000 (1%) pregnant women in Scotland.

A national audit in Scotland reported a lower incidence of severe maternal morbidity of 6 in 1000, the most common



Figure 2 Pareto chart of the rates of 26 severe maternal morbidities per 1000 pregnancies, conditions (red) and procedures (blue). The black line is the cumulative proportion for all severe morbidities. DIC, disseminated intravascular coagulation.

of which was post-partum haemorrhage, a morbidity we excluded [16]. We think that inaccurate coding in our study databases may have overestimated the rate of morbidity. The rate of morbidity in our study would be 6 in 1000 if we excluded puerperal sepsis, the coding of which increased substantially after 2012. Several studies suggest that ICD coding for sepsis is unreliable [21, 22]. The reliability of puerperal sepsis coding could be improved by requiring a simultaneous code for organ dysfunction derived from critical care databases [23].

Other studies have reported similar associations of variables with severe maternal morbidity [3, 8, 24, 25]. The variables associated with morbidity did not necessarily cause morbidity, directly or indirectly, particularly variables recorded during labour and delivery. Such variables are more likely to be the consequence of morbidity, for instance emergency caesarean section. Some studies have used a causal framework and undertaken mediation analyses to better explore causation [25]. Interventions early in pregnancy, such as ensuring ready access to maternity services and pre-emptive individualised support, might improve maternal and foetal outcomes for women who are

older, comorbid, obese, born in non-European countries or who have had previous caesarean sections.

Women who had a severe morbidity were more than 2000 times more likely to die than women who did not, which is greater than risk ratios of 100–500 reported by other studies [26, 27]. Unsurprisingly, women with morbidity stayed in hospital longer than women without, although the difference was less than in a Canadian study [27]. The disparity could be due to differences in health service organisation, or because of the incorrect coding that we suspect of our study.

The use of multiple databases increased our ability to detect associations with an uncommon outcome and increased the external validity of our study. We included women cared for in obstetric high dependency units, who are usually missed by current national reports of critically ill pregnant women. We think that admissions to obstetric high dependency units might be incorporated as a component of severe maternal morbidity, perhaps in combination with coding for particular conditions or procedures. However, as guidance increasingly supports pre-emptive admission to critical care for those at higher



Figure 3 (a–c) Odds ratios (95%CI) of characteristics associated with severe maternal mortality after model 1 multivariable adjustment. Reference categories have an odds ratio of 1.

risk of adverse outcomes, it may be that unplanned critical care admission would be a more suitable morbidity to analyse in the future [28].

There are also limitations to this study. Some variables, such as BMI, were missing many values. We intentionally analysed a composite outcome, which may misrepresent the associations of individual component morbidities with variables, as there is already an established body of research on individual morbidities, for instance post-partum haemorrhage and maternal sepsis [29–33].

In conclusion, severe maternal morbidity accompanied 10 in 1000 (1%) pregnancies in Scotland. Morbidity was independently associated with maternal age; BMI; preexisting morbidity; previous smoking; previous caesarean section; multiple pregnancy; and maternal birth in Africa or the Middle East. Morbidity was associated with delayed hospital discharge, stillbirths and maternal deaths. Our composite measure of severe maternal morbidity might be tested with other administrative healthcare databases.

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Supporting Information

Additional supporting information may be found online via the journal website.

Appendix S1. Detailed methods.

Appendix S2. Supplementary methods.

Table S1. Components of severe maternal morbidityderived from Scottish datasets modified from the EnglishMaternal Morbidity Indicator.

Table S2. Prevalence of comorbidities in women for whom severe maternal morbidity was or was not reported, for comorbidities coded in databases during the 5 y before conception only and combined with comorbidities coded during pregnancy.

Table S3. Incidence of severe maternal morbidity.

Table S4. The proportions of severe maternalmorbidity that were conditions or interventions.

Table S5. Unadjusted associations of characteristics with severe maternal morbidity and adjusted associations in model 1 and model 2.

Table S6. Associations of 1, 2 or 3 comorbidities with

 severe maternal morbidity.

Table S7. A sensitivity analysis of Table S5, with comorbidities recorded during pregnancy added to models 1 and 2.

 Table S8. Intensive care variables for women with severe maternal morbidity.

Figure S1. A sensitivity plot of Fig. 2, with the addition of admission to critical care – intensive care or high dependency care – as a component of severe maternal morbidity.

Figure S2. A sensitivity plot of Fig. 2 and online Supporting Information (Figure S2), with the addition of admission to obstetric high dependency as a component of severe maternal morbidity.

Figure S3. Annual incidence of severe maternal morbidity 2004–18.

Figure S4. Predicted rate of severe maternal morbidity vs. maternal age.

Figure S5. Predicted rate of severe maternal morbidity vs. maternal body mass index.

Figure S6. Predicted rate of severe maternal morbidity vs. date.