

Severe Maternal Morbidity Audit Clinical Reference Manual

Queries to:

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Please return all completed forms to the above addressee at:

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Overview



Background

- Severe maternal morbidity (SMM) has been acknowledged internationally as an important indicator of quality of maternal care and welfare, particularly in developed countries where maternal death rates are relatively low.
- For each woman who loses her life due to causes related to pregnancy, many more experience lifethreatening complications or longterm morbidities.
- In this context, the NPEC in collaboration with the NPEC Severe Maternal Morbidity Advisory Group has collected and analysed data on SMM from Irish maternity units since 2011.

Methodology

To allow for international comparison, the NPEC adapted the validated methodology of the Scottish
 Confidential Audit of Severe Maternal Morbidity (SCASMM) to evaluate severe maternal morbidity (SMM) in Ireland.
 This methodology utilises organ dysfunction criteria described by Mantel et al,¹ with modifications used by SCASMM to include intervention-based criteria.²

Objective

 The fundamental aim of the audit is to provide a national review of clearly defined SMMs, to identify quality improvement initiatives and make recommendations for the improvement of maternity care for women in Ireland.

Inclusion criteria and reportable morbidities



Inclusion criteria for the SMM audit

All pregnant or recently pregnant women (up to and including 42 days following delivery, miscarriage, ectopic pregnancy or termination of pregnancy) who experienced a SMM as defined in this audit.

Reportable severe maternal morbidities

In this audit there are fourteen, clearly defined, organ dysfunction morbidities and two management proxies for maternal morbidity (ICU/CCU admission and interventional radiology). Definitions for all reportable SMM events are detailed in Table 1 on the next slide.

Please indicate all SMM events experienced by the woman in the current / most recent pregnancy
 Please note more than one SMM in the pregnancy may apply to a woman and should be reported on <u>one</u> SMM notification form.

✤ In the event of an eclampsia case, the NPEC kindly request you complete the detailed Eclampsia Audit dataset.

Table 1: SMM Definitions



		Estimated blood loss ≥ 2500ml and/or transfused 5 or more units of blood. Also includes				
		Estimated blood loss ≥ 2500ml and/or transfused 5 or more units of blood. Also includes miscarriage, ectopic pregnancy or termination of pregnancy meeting these criteria.				
1	Major obstetric haemorrhage	(Please record as well whether treatment for coagulopathy was received).				
-		A complete separation of the wall of the pregnant uterus, with or without expulsion of the fetus,				
		involving rupture of membranes at the site of the uterine rupture or extension into uterine				
		muscle separate from any previous scar, and endangering the life of the mother or fetus.				
2	Uterine rupture	Excluded: any asymptomatic palpable or visualised defect (e.g. dehiscence noted incidentally at				
		caesarean delivery)				
3	Peripartum hysterectomy	Peripartum hysterectomy				
4	Eclampsia	Seizure associated with antepartum, intrapartum or postpartum symptoms and signs of pre-				
	Renal or liver dysfunction	eclampsia				
5	Acute onset of biochemical disturbance, urea >15mmol/l, creatinine>400mmol/l,					
		AST/ALT >200u/I				
c		Clinically diagnosed pulmonary oedema associated with acute breathlessness and O2				
6	Pulmonary oedema	saturation <95%, requiring O2, diuretics or ventilation				
7	Acute respiratory	Requiring intubation or ventilation for >60 minutes (not including				
	dysfunction	duration of general anaesthetic)				
		Increased respiratory rate (>20/min), tachycardia, hypotension.				
8	Pulmonary embolism	Diagnosed as "high" probability on V/Q scan or positive spiral chest CT scan. Treated by heparin,				
		thrombolysis or embolectomy				
9	Cardiac arrest	No detectable major pulse				
10	Coma	Including diabetic coma. Unconscious for >12 hours				
11	Cerebro-vascular event	Stroke, cerebral/cerebellar haemorrhage or infarction, subarachnoid haemorrhage, dural venous sinus thrombosis				
12	Status epilepticus	Constant or near constant state of having seizures that last 30mins or more				
12		-				
		Sepsis induced tissue hypoperfusion or hypotension persisting after resuscitation with 30mls/kg intravenous isotonic crystalloid fluid as evidenced by:				
13		 Systolic blood pressure < 90 mmHg or MAP < 65 mmHg 				
13	Septicaemic shock	 Decrease in systolic blood pressure by 40mmHg from baseline and/or 				
14	A no osthotic problem	 Lactate > 4 mmol/l. Aspiration, failed intubation, high spinal or epidural anaesthetic 				
14	Anaesthetic problem					
12	ICU/CCU admission	Unit equipped to ventilate adults. Admission for one of the above				
		problems or for any other reason. Includes CCU admissions				

Received planned (a) or unplanned (b) interventional radiology

16

Interventional radiology

Please ensure reported SMM events meet the defined criteria as outlined in the Table,

1. Eclampsia is defined as: "Seizure associated with antepartum, intrapartum or postpartum symptoms and signs of preeclampsia". This definition does NOT include cases of severe PET without seizures or epileptic seizures.

2. Acute respiratory Dysfunction: is defined as acute respiratory dysfunction "Requiring intubation or ventilation for >60 minutes (not including duration of general anaesthetic)". This definition does NOT include cases of respiratory dysfunction which does not require mechanical ventilation in the management of the woman e.g. Pneumonia treated with oxygen therapy via facial mask.

e.g:

Data Submission



- It is recommended that cases be submitted to the NPEC on a monthly basis, if at all possible. An annual submission date for complete data will be within five months of the reporting year end.
- > Audit data can be submitted online via the NPEC secure database or alternatively in paper format.
- > An operational manual and training video for the NPEC online database is available on the NPEC website

Manual and training video

Calculating SMM rates

Severe Maternal Morbidity cases are included in a maternity unit's rate if the woman was delivered in that maternity unit. If a woman experiencing a SMM is transferred to another unit following delivery, (or is admitted during the postnatal period to another maternity unit), the SMM should be reported by the maternity unit where the delivery took place.

The NPEC can assist in communications between unit co-ordinators if required/requested. This will help validate complete case ascertainment at national level.



Most questions should be self-explanatory, but the following notes give guidance to specific questions within the sections of the data collection form. Of note, the online form has been developed to show questions that are relevant to the reported SMM category. For example, if MOH is selected, you will then be asked a series of MOH specific questions. See examples below:

- In the event of a Major Obstetric Haemorrhage (MOH), please indicate:
 - How the criteria was met (i.e. EBL \geq 2,500mls and/or transfused with \geq 5 units of blood)
 - What was the primary cause of the MOH. The following list will be available for you to select from:

Abruption
Retained placenta/membranes
Uterine rupture
Bleeding from uterine incision
Placenta praevia
Vaginal laceration/haematoma
Uterine inversion
Placenta Accreta Spectrum (PAS), previously known as Morbidly adherent placenta (e.g. placenta accreta or per accreta)
Uterine atony
Cervical laceration
Broad ligament haematoma
Other

- In the event of Peripartum hysterectomy (PH), please indicate:
 - The indication for the PH. A list will be available for you to choose from including: Placenta accreta spectrum (PAS), Ο placenta praevia, major obstetric haemorrhage, uterine atony, uterine or cervical cancer, or other
 - Was the PH a planned elective surgery (i.e. planned PH prior to delivery)
 - Place of surgery (maternity unit or general hospital) Ο

Please specify indication for PH		 Placenta Accreta Spectrum (PAS), e.g. placenta percreta, and placenta accreta Placenta praevia Major obstetric haemorrhage Uterine or cervical cancer Other 		
Was this a planned elective PH surgery?	~ M	○ Yes ○ No		
Place of surgery	@ 19	O Maternity hospital O General hospital		



- In the event of Septic Shock, please indicate the underlying cause of infection, e.g. •
 - If the cause was pregnancy-related, e.g. chorioamnionitis, wound infection following caesarean section etc. Or non-pregnancy related, e.g. pneumonia, COVID-19 infection etc..

Septicaemic shock	 Sepsis induced tissue hypoperfusion or hypotension persisting after resuscitation with 30mls/kg intravenous isotonic crystalloid fluid as evidenced by: Systolic blood pressure < 90 mmHg or MAP < 65 mmHg Decrease in systolic blood pressure by 40mmHg from baseline and/or Lactate > 4 mmol/l. 	Please select type of septicaemic shoc O Pregnancy-related O Non-Pregnancy related O Unknown if pregnancy related
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- In the event of an ICU/CCU admission
 - In cases where ICU admission was the ONLY reportable event, and the woman did not experience another SMM as defined by this audit, you will be asked to report the indication for admission to ICU/CCU. A list of clinical events (not exhaustive) to choose from is available:
 - Later in the form you will be asked how long the woman's ICU stay was in days.
 Please enter a number only, e.g. 1.5
- In the event of Interventional Radiology (IR), please indicate:

 $\circ~$ If it was IR was planned or unplanned

- Alone, no other associated SMM as defined in this audit
- With associated SMM as defined is this audit

If no other morbidity **as defined in this audit** was present, please select the indication for ICU admission:

- Obstetric haemorrhage < 2,500mls - monitoring
- O Pre-eclampsia monitoring
- Pre-eclampsia MgSO4 infusion / antihypertensive treatment
- Sepsis not septic shock -pregnancy related
- Sepsis not septic shock -not pregnancy related
- \bigcirc Cardiac condition please describe
- Other medical condition, please specify
- Other surgical condition, please specify



- Woman's details:
 - Q. Place of delivery

Please indicate the hospital the woman gave birth in, or in relevant cases if the women gave birth in a general hospital, had a homebirth or was a BBA.

Q. What was the antenatal care pathway assigned to this woman **PRIOR** to the SMM event?
 This refers to the pathway of care based on the National Maternity Strategy Model of Care as outlined below and in Appendix A.

- Supported Care Pathway; Midwifery led and delivered care.
- > Assisted Care Pathway; Obstetric led, Midwifery and Obstetric delivered care.
- Specialised Care Pathway; Obstetric led, Obstetric and Midwifery delivered care

Please note once a woman experiences one or more SMM, her pathway of care will be escalated to medium or high risk.

Woman's details:

Booking: Some data sought by the NPEC relate to the time of booking. Booking in this regard relates to the mother's first antenatal visit at the maternity unit.

Parity: The NPEC refer to parity prior to the current pregnancy or most recent pregnancy if delivered. The number of completed pregnancies, whether live birth or stillbirth, of at least 24 weeks gestation or with a birthweight \geq 500g. Number of pregnancy losses (less than 24 weeks of gestation)

Hospital transfer:

If the woman was transferred FROM or TO another hospital, please indicate the hospital and the timing of pregnancy in relation to the pregnancy status:

- Woman transferred with fetus in utero.
- Woman transferred following delivery of baby.



• <u>Obstetric history, pregnancy and delivery details:</u> *Definitions of baby outcome:*

Alive. to the complete expulsion or extraction from its mother of a product of conception, irrespective of the duration of the pregnancy, which, after such separation, breathes or shows any other evidence of life - e.g. beating of the heart, pulsation of the umbilical cord or definite movement of voluntary muscles - whether or not the umbilical cord has been cut or the placenta is attached. Each product of such a birth is considered live born.³

Early miscarriage. The spontaneous expulsion of a fetus from the womb before 13 weeks gestation.

Ectopic pregnancy. An extra uterine pregnancy which occurs when a fertilized egg implants and grows outside the main cavity of the uterus, most commonly in the fallopian tube.

Late miscarriage. The spontaneous expulsion of a fetus from the womb between 13 weeks to 24 weeks of gestation.

Termination of pregnancy. The medical or surgical termination of pregnancy with the expected outcome of fetal or early neonatal death. (See further detail below)

Stillbirth. Baby delivered without signs of life from 24 weeks gestation or with a birthweight ≥500g. 4

Early neonatal death. Death of a live born baby occurring within 7 completed days of birth.

Late neonatal death. Death of a live born infant occurring after the 7th day and within 28 days of birth.

• Obstetric history, pregnancy and delivery details:

If the case is associated with a termination of pregnancy (TOP), please select under which Section of the Health (Regulation of Termination of Pregnancy) Act 2018 the TOP was carried out:

- Section 9. Risk to life or health
 There is a risk to the life, or of serious harm to the health, of the pregnant woman and the foetus has not reached viability
- Section 10. Risk to life or health in emergency
 There is an immediate risk to the life, or of serious harm to the health, of the pregnant woman
- Section 11. Condition likely to lead to death of foetus

There is present a condition affecting the foetus that is likely to lead to the death of the foetus either before, or within 28 days of, birth

Section 12. Early pregnancy
 Whereby the pregnancy concerned has not exceeded 12 weeks of pregnancy



• Obstetric history, pregnancy and delivery details:

If Caesarean section (CS) is indicated as the mode of birth, please indicate:

The category of CS, as per the NICE⁵ classification or urgency,

- Category 1. Immediate threat to the life of the woman or fetus (for example, suspected uterine rupture, major placental abruption, cord prolapse, fetal hypoxia or persistent fetal bradycardia).
- o Category 2. Maternal or fetal compromise which is not immediately life-threatening.
- Category 3. No maternal or fetal compromise but needs early birth.
- Category 4. Birth timed to suit woman or healthcare provider.

In the case that CS was selected with MOH, you will be asked the following:

- $\circ~$ If the woman was fully dilated before the CS
- $\circ~$ If it was a failed instrumental delivery





Obstetric history, pregnancy and delivery details:

If Caesarean section (CS) is indicated as the mode of birth, following a spontaneous or induced onset of labour, please select the relevant indication from the list below:

Indication for caesarean section - In Labour/After Induction

	O EUA - Cephalopelvic disproportion	Exa
	O EUA - Persistent malposition	EUA
Ð	🔿 Fetal reason (no oxytocin)	EUA
M	\bigcirc IUA - Inability to treat fetal intolerance	Feta
	○ IUA - Poor response	IUA

EUA: Efficient uterine activity, IUA: Inefficient uterine activity

mp	les*:	

EUA - Cephalopelvic disproportion: Failure to progress

EUA - Persistent malposition: Occipito-posterior position

Fetal reason (no oxytocin)

- IUA Inability to treat fetal intolerance: Non-reassuring CTG
- IUA Poor response: Failed induction/ poor response to augmentation

*EUA: Efficient uterine activity, IUA: Inefficient uterine activity

If Caesarean section (CS) is indicated as the mode of birth, where the woman was never in labour, please select the relevant indication from the list below:

	○ Fetal reason
	O Maternal medical reason/pains
	O Non medical reason/patient request
aesarean section: never in labour	O PET/Hypertension
	Previous caesarean section
	Other
	reset

Location of Care:

Please indicate the location of care <u>DURING</u> the SMM event, please tick all that apply.

Level of Care:

Within the term 'critical care', care is subdivided into four levels, dependent on organ support and the level of monitoring required independent of clinical diagnosis. ⁶ Please indicate the <u>HIGHEST</u> level of care during the SMM clinical event. . Examples of Level of Care are outlined in Appendix B.

Level of care	Definition	Please tick one box
Level 0: Normal ward care	Care of low risk pregnant women	
Level 1: Additional monitoring or intervention, or step down from higher level of care	Patients at risk of their condition deteriorating and needing a higher level of observation or those recently relocated from higher levels of care	
Level 2: Single Organ Support**	Patients requiring invasive monitoring/ intervention* including support for a single failing organ system (excluding advanced respiratory support).	
Level 3: Advanced respiratory support alone, or support of two or more organ systems**	Patients requiring advanced respiratory support (mechanical ventilation) alone or basic respiratory support along with support of at least one additional organ.	



 On the ward Delivery Suite Theatre High Dependency Unit

Redcap specific detail on SMM form



- Please note: **Redcap** is the secure online platform used to collect data for this audit. More detail regarding how to get set up and use Redcap will be available from our website or by contacting a member of the NPEC staff.
- There are some features to be aware of that are specific to the SMM audit, as follows:
 - On your unit's list of cases under "Record status dashboard", "DCE" refers to the Date of Clinical Event, i.e. the date the morbidity occurred.

 If you wish to review your data using the "Stats & Charts" function, you will be asked to select an 'instrument', this is referring to which data collection form you would like to review in this format.

rd",	NPEC reference number				Severe Maternal Morbidity form	Eclampsia	
5	<u>1</u> (Local	2023)			0		
	<u>2</u> (Local	Ref. no. Test2)	(DCE 10-05-	2023)		۲	
	<u>3</u> (Local	Ref. no. Test3)	(DCE 10-05-	2023)		۲	
	4 (Local	Ref. no. Test4)	(DCE 10-05-	2023)		۲	
evere Mate	ernal Mor	bidity Audit (SI	MM) PID 29				
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+ Create Net	w Report	Hy Reports & Expo	orts 🕑 Other I	Export Options	E Sta	ats & Charts: A	ll data (all records
l umber of resul otal number of r All data (all	records querie	d: 26	Q View Reput	Dort Export Da		Print Page	s Group]
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Viewing option	s: Show plo	ots & stats Show p	olots only Sho	w stats only			



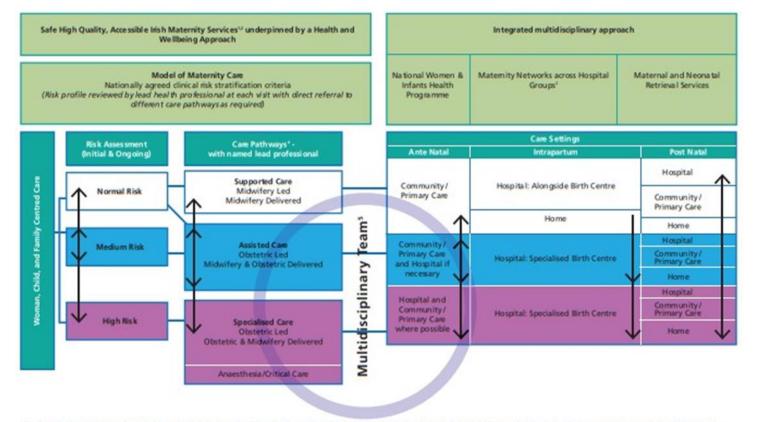
Thank you!

THANK YOU FOR COMPLETING THIS SMM AUDIT DATASET.

YOUR CONTRIBUTION IS GREATLY APPRECIATED AND VALUED.



Appendix A: National Maternity Strategy Model of Care



1 In line with the overriding safety principle, a risk based approach will be utilised. Practice will be evidence based using national clinical guidelines and audit, and quality improvement approaches will be adopted.

- 2. Each birth centre should have access to an immediate emergency team response for dinical deterioration.
- 3 For high risk and complex women or babies specialist services outside the network may be required.
- 4 Within each of these care pathways, women can also avail of a shared model of care with the GP as provided for by the Matemity and Infant Care Scheme.

5 Spanning the acute, primary and community sectors, modern maternity services are multi-disciplinary in nature, and as such require the involvement of a range of health professionals. The input of the wider multi-disciplinary team will be co-ordinated by the lead healthcare professional.



Appendix B: Defining Level of Care

Levels of care

National and International guidelines have recommended that the terms high dependency and intensive care be replaced by the term critical care. Within the term critical care, care is subdivided into four levels, dependent on organ support and the level of monitoring required independent of clinical diagnosis.

Level of care	Definition	Please tick one box
Level 0: Normal ward care	Care of low risk pregnant women	
Level 1: Additional monitoring or intervention, or step down from higher level of care	Patients at risk of their condition deteriorating and needing a higher level of observation or those recently relocated from higher levels of care	
Level 2: Single Organ Support**	Patients requiring invasive monitoring/ intervention* including support for a single failing organ system (excluding advanced respiratory support).	
Level 3: Advanced respiratory support alone, or support of two or more organ systems**	Patients requiring advanced respiratory support (mechanical ventilation) alone or basic respiratory support along with support of at least one additional organ.	

* invasive monitoring/intervention includes the use of arterial and CVP lines

**Examples of level 2 and 3 care in the critically ill pregnant or recently pregnant woman are outlined below

Level 2 examples

Basic Respiratory Support (BRS): 50% or more oxygen via face-mask to maintain oxygen saturation; Continuous Positive Airway Pressure (CPAP), Bi- Level Positive Airway Pressure (BIPAP)

Basic Cardiovascular Support (BCVS): Intravenous anti-hypertensive, to control blood pressure in pre-eclampsia; Arterial line used for pressure monitoring or sampling; CVP line used for fluid management and CVP monitoring to guide therapy

Advanced Cardiovascular Support (ACVS): Simultaneous use of at least two intravenous, anti-arrhythmic/anti-hypertensive/vasoactive drugs, one of which must be a vasoactive drug; Need to measure and treat cardiac output

Neurological Support: Magnesium infusion to control seizures / prophylaxis of eclampsia in severe PET

Hepatic Support: Management of acute fulminant hepatic failure, e.g. from HELLP syndrome or acute fatty liver, such that transplantation is being considered

Level 3 examples

Advanced Respiratory Support: Invasive mechanical ventilation

Support of two or more organ systems: Renal support and BRS; BRS/BCVS and an additional organ supported; Intracranial pressure monitoring



Reference: Saravanakumar K, Davies L, Lewis M, Cooper GM.. High dependency care in an obstetric setting in the UK. Anaesthesia 2008:63, 1081–6.



- 1. Mantel G et al. Severe Acute maternal morbidity: a pilot study of a definition for a near-miss. BJOG 1998; 105: 985-90
- 2. Scottish Confidential Audit of Severe Maternal Morbidity: 10th Annual Report (2014). Available from:http://www. healthcareimprovementscotland.org/our_work/reproductive,_maternal child/programme_resources/scasmm.aspx
- 3. World Health Organisation. Available at: <u>http://www.who.int/healthinfo/statistics/indmaternalmortality/en/</u>
- 4. Stillbirths Registration Act, 1994
- 5. National Institute for Health and Care Excellence (2021) Caesarean birth. NICE Guidelines.
- 6. Clinical Practice Guideline No 30 (2014). Guideline for the Critically ill Woman in Obstetrics: Institute of Obstetricians and Gynaecologists, Royal College of Physicians of Ireland and Directorate of Strategy and Clinical Programmes, Health Service Executive



SEVERE MATERNAL MORBIDITY in Ireland