

For NPEC Office use only: CASE NUMBER

PLACE OF DEATH:

PERINATAL DEATH NOTIFICATION FORM 2020

CHOOSE Type of Case (TICK)
STILLBIRTH: A baby delivered without signs of life from 24 weeks' gestation and/or with a birth weight of ≥ 500g.
*If the birth occurred unattended and there was no lung aeration seen at Post Mortem (PM) and no other circumstantial evidence of life at birth, it should be assumed that the baby was stillborn.
OR
EARLY NEONATAL DEATH: Death of a live born baby occurring before 7 completed days after birth.
OR
LATE NEONATAL DEATH: Death of a live born baby occurring from the 7 th day and before 28 completed days after birth.
* For the purpose of reporting, a 'live born' baby is defined as any baby born with evidence of life such as breathing movements, presence of a heart beat, pulsation of the cord or definite movement of voluntary muscles.
If a baby born at <22 completed weeks is being registered as a neonatal death, please report same to NPEC.

The National Perinatal Epidemiology Centre is sincerely grateful for your contribution to this audit.

Guidance for completing this form, with specific reference to Sections 11, 12 and 13 on Cause of Death, is outlined in the accompanying reference manual.

The National Perinatal Epidemiology Centre also acknowledges with thanks the Centre for Maternal and Child Enquiry (CMACE) UK for permission to modify and use its Perinatal Mortality Notification Proforma for use in the Irish context.

CTION 1. WOMANS' DETAILS			
1.1. Mother's age			
1.2. Ethnic group:			
White - Irish	☐ Irish Traveller		
Any other White background	Please specify count	ry of origin	
Asian or Asian Irish	Black or Black Irish		
Other including mixed ethnic b	ackgrounds: Please specify ₋		
Not recorded			
1.3. Marital status:	d Never marrie	ed Separated/Divorced	I ☐ Widowed ☐ Unknown
1.4. Living with partner / spous	se? □Yes	□No	Unknown
1.5. Woman's employment star	tus at booking?		
☐Employed or self-employed (fu	ıll or part time)	☐ Unemployed (looking for w	vork)
_ ` ` ` ` _ ` `	me maker	Permanently sick/disabled	,
Other	Unknown		
1.7. Height at booking (round u	p to the nearest cm):		
1.8. Weight at booking (round u	p to the nearest kg):		
If weight is unavailable, was ther	e evidence that the woman w	as too heavy for hospital scale	es? Yes No
1.9. Body Mass Index at bookin	g (BMI):		
4.40 a Biddhawanan amala da		:£	
1.10.a. Did the woman smoke at		ify quantity smoked per day	
	∐ No	Unknown	
1.10.b. Did she give up smoking	g during pregnancy?	☐ Yes ☐ No	☐ Unknown ☐ N/A
1.11. Is there documented histo	ry of alcohol abuse?		
None recorded	Prior to this pregnancy	During this pre	gnancy
1.12. Is there documented histo			
None recorded	Prior to this pregnancy	During this pre	gnancy
	2		

SECTION 2. PREVIOUS PREGNANCIES	
2.1. Did the woman have any previous pregnancies? If yes, please complete	ete questions 2.2-2.4 Yes No
2.2. No. of completed pregnancies ≥24 weeks and or with a birth weight	ht ≥ 500g (all live and stillbirths): ☐ ☐
2.3. No. of pregnancies <24 weeks and with a birth weight < 500g:	
2.4. Were there any previous pregnancy problems? If yes, please tick all that	at apply below Yes No
☐ Three or more miscarriages ☐ Pre-term birth or mid trimester loss	Stillbirth, please specify number
☐ Infant requiring intensive care ☐ Baby with congenital anomaly	Neonatal death, please specify number
Previous caesarean section Placenta praevia	Placental abruption
Pre-eclampsia (hypertension & proteinuria)	Post-partum haemorrhage requiring transfusion
☐ Other, please specify	∐Unknown
SECTION 3. PREVIOUS MEDICAL HISTORY	
3.1. Were there any pre-existing medical problems? If yes, please tick all the	at apply below Yes No Unknown
☐ Cardiac disease (congenital or acquired) ☐ Epilep	osy
☐ Endocrine disorders e.g. hypo or hyperthyroidism ☐ Renal	I disease
	niatric disorders
	rtension
	, please specify
 4.1. Final Estimated Date of Delivery (EDD):	Unknown a 40 week gestation, or the final date agreed Yes No
4.3. Was this pregnancy a result of infertility treatment?	☐ Yes ☐ No ☐ Unknown
4.4 Gestation at first booking appointment: weeks + days 4.5 Intended place of delivery at booking: Name of unit	☐ Not booked ☐ Unknown
Please specify the type of unit	
Obstetric Unit Alongside Midwifery Unit Home	Unbooked
4.6 What was the intended type of delivery care at booking?	
Obstetric-Led Care Midwifery-Led Care Self-Emp	ployed Community Midwife
Home c/o Hospital DOMINO Scheme	

4.8 a Did the woman undergo an anatomy scan? If yes please answer question 4.8 b 4.8 b Gestation at time of anatomy scan: CTION 5. DELIVERY 5.1. Onset of labour: Spontaneous Induced Never in labour Please specify the type of unit Obstetric Unit Alongside Midwifery Unit Home 5.3. What was the intended type of care at onset of labour? Obstetric-Led Care Midwifery-Led Care Self-Employed Community Midwife Home c/o Hospital DOMINO Scheme	atomy scan? 8 b scan: weeks + days we	If yes please answer question 4.7 b
If yes please answer question 4.8 b 4.8 b Gestation at time of anatomy scan:	scan: weeks + days Induced Never in labour Induced Yes No Induced Never in labour Induced Never in	4.7b Gestation at time of in-utero transfer:
5.1. Onset of labour: Spontaneous Induced Never in labour Spontaneous Induced Never in labour 5.2. Intended place of delivery at onset of labour: Name of unit Please specify the type of unit Home 5.3. What was the intended type of care at onset of labour? Obstetric-Led Care Midwiftery-Led Care Self-Employed Community Midwife Home c/o Hospital DOMINO Scheme 5.4. Was the intended mode of delivery a planned caesarean section? Yes New of unit Please specify the type of unit Other, please specify Obstetric-Led Care Midwiftery-Led Care Born Before Arrival (BBA) - Unattended Self-Employed Community Midwife Home c/o Hospital DOMINO Scheme 5.7. Date and time of delivery/birth: Date: // // Time: : 5.8. What was the presentation at delivery? Longitudinal Oblique Transverse 5.9. What was the presentation at delivery?	nduced Never in labour Name of unit	• ,
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Please specify the type of unit Obstetric Unit	care at onset of labour? Midwifery-Led Care	☐ Spontaneous ☐ Induced ☐ Never in labour
Obstetric Unit	Care at onset of labour? Midwifery-Led Care	5.2. Intended place of delivery at onset of labour: Name of unit
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5.8. What was the lie of the fetus <u>at delivery</u> ? Longitudinal Oblique Transverse 5.9. What was the presentation <u>at delivery</u> ?	delivery? Oblique Transverse elivery? Compound (includes transverse and shoulder presentations) Brow Face (? (Please tick all that apply) Ventouse Forceps Assisted Breech delivery	Self-Employed Community Midwife Home c/o Hospital DOMINO Scheme
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Longitudinal Oblique Transverse 5.9. What was the presentation <u>at delivery</u> ?	Oblique Transverse clivery? Compound (includes transverse and shoulder presentations) Brow Face (? (Please tick all that apply) Ventouse Forceps Assisted Breech delivery	5.8. What was the lie of the fetus at delivery?
5.9. What was the presentation <u>at delivery</u> ?	Compound (includes transverse and shoulder presentations) Brow Face (? (Please tick all that apply) Ventouse Forceps Assisted Breech delivery	
Vertex Breech Compound (includes transverse and shoulder presentations) Brow	Y? (Please tick all that apply) Ventouse	
	Ventouse	
5.10. What was the mode of delivery? (Please tick all that apply)		5.9. What was the presentation <u>at delivery</u> ?
☐ Vaginal cephalic delivery ☐ Ventouse ☐ Forceps ☐ Assisted Breech delivery		5.9. What was the presentation <u>at delivery</u> ? Uvertex Breech Compound (includes transverse and shoulder presentations) Brow Face
☐ Vaginal Breech delivery ☐ Pre-Labour Caesarean Section ☐ Caesarean Section After Onset of Lab	Pre-Labour Caesarean Section	5.9. What was the presentation <u>at delivery?</u> Vertex Breech Compound (includes transverse and shoulder presentations) Brow Face 5.10. What was the mode of delivery? (Please tick all that apply)

CAESAREAN SECTIONS ONLY	
5.11. What was the type of or indication for Caesarean Section?	
Elective - At a time to suit woman or maternity team Urgent - Maternal or fetal compron	nise which is not immediately life threatening
Emergency - Immediate threat to life of woman or fetus Failed instrumental deli	very
SECTION 6. ALL BABY OUTCOME	
6.1. Sex of fetus/baby:	Female Indeterminate
6.2. Number of fetuses/babies in this delivery: (all identifiable including papyraceous Birth order of this fetus/baby:	s)
Singleton	
☐ Twin 1 ☐ Twin 2	
☐ Triplet 1 ☐ Triplet 2 ☐ Triplet 3	
Other multiple birth pregnancy, please specify Birth Order	
6.3. If from a multiple delivery, what was the chorionicity? Please tick all that ap	pply
☐ Dichorionic diamniotic ☐ Monochorionic diamniotic ☐ Monochorionic monoam	niotic Trichorionic
☐ Singleton ☐ Not known	
6.4. Birth weight (kg):	
6.5. Gestation at delivery: □ □ weeks + □ days	Unknown
6.6. Was this a termination of pregnancy? Please refer to the reference manual	☐ Yes ☐ No
6.7. Was a local hospital review of this case undertaken? Please refer to the reference manual	☐ Yes ☐ No
SECTION 7. MATERNAL OUTCOME	
7.1. Admission to HDU:	☐ Yes ☐ No
7.2. Admission to ICU:	☐ Yes ☐ No
7.3. Maternal Death:	☐ Yes ☐ No
SECTION 8. STILLBIRTH (If not a stillbirth, please go to Section 9)	
8.1. At what gestation was death confirmed to have occurred?	□ □ weeks + □ days
If known, what date was death confirmed?	
8.2. Was the baby alive at <u>onset of care</u> in labour?	
	attended Unknown
5	

SECTION 9. NEONATAL DEATH ONLY	
9.1. Was spontaneous respiratory activity absent or ineffective at 5 minutes?	☐ Yes ☐ No
If a baby is receiving any artificial ventilation at 5 minutes, the assumption is absent/ineffective activity: absent activity.	a 0 Apgar score indicates
9.2. Was the heart rate persistently <100bpm? (i.e. heart rate never rose above 100bpm b	efore death)
Persistently <100bpm	Rose above 100bpm
9.3. Was the baby offered *active resuscitation in the delivery room? (*active resuscitation includes BMV, PPV, intubation, cardiac massage)	☐ Yes ☐ No
9.4. Was the baby admitted to a neonatal unit? (Includes SCBU and ICU)	☐ Yes ☐ No
9.5a. Was the baby transferred to another unit after birth? If yes please answer 9.5 b	☐ Yes ☐ No
9.5 b. Date and Time of Transfer to other unit <u>after birth</u> : Date \(\Boxed{\omega} \subseteq \subsete \Boxed{\omega} \subsete \Boxed{\omega} \subsete \Boxed{\omega}	Time : .
9.6. Date and Time of Death:	Time
9.7. Place of Death*: Labour Ward Neonatal Unit Ward	☐ Theatre
☐ In Transit ☐ Paediatric Centre ☐ Home	
Name of unit:	
*This question refers to where the baby actually died, e.g. 'ICU, 'at home' or 'in transit'. Babies are deemed to have died 'at home' if there are no signs of life documented in the home even if resuscitation A baby is deemed to have died 'in transit' if signs of life are documented prior to transfer but the baby was either do the hospital or showed no subsequent signs of life in the hospital, despite attempted resuscitation	
SECTION 10. POST-MORTEM INVESTIGATIONS	
10.1. Was this a coroner's case? If yes, please complete question 10.2.	☐ Yes ☐ No
10.2. Has the post-mortem report been received from the coroner's office?	☐ Yes ☐ No
10.4. Was a post-mortem performed? Yes No If no, please complete question 10.5.	
10.5. Was a post-mortem offered?	☐ Yes ☐ No
10.6. Were any of the following procedures carried out after death? Please tick all that apply	
☐ MRI ☐ X-Ray ☐ CT ☐ External Examination ☐	Genetic testing
10.7. Was the placenta sent for histology?	☐ Yes ☐ No
6	

SECTION 11. CAUSE OF DEATH AND ASSOCIATED FACTORS - STILLBIRTH & NEONATAL DEATH 11. Please TICK ALL the maternal or fetal conditions that were present during pregnancy or were associated with the death. PLEASE REFER TO THE REFERENCE MANUAL. 11.1.1. MAJOR CONGENITAL ANOMALY: Central nervous system ☐ Cardiovascular system Respiratory system Gastro-intestinal system Musculo-skeletal anomalies Multiple anomalies Urinary tract Metabolic diseases Other major congenital anomaly, please specify _____ Chromosomal disorder*, please specify _____ * In the event of a chromosomal disorder how was the diagnosis made? Clinically Genetic analysis * Ultrasound *See reference manual 11.1.1 (b) Was the diagnosis of major congenital anomaly confirmed/suspected before delivery by a Consultant Fetal Yes, in your unit **Medicine Specialist?** Yes, in another unit, please specify name of unit 11.1.2. HYPERTENSIVE DISORDERS OF PREGNANCY: Pregnancy induced hypertension Pre-eclampsia HELLP syndrome Eclampsia 11.1.3. ANTEPARTUM or INTRAPARTUM HAEMORRHAGE: Praevia Abruption Other, please specify _____ 11.1.4. MECHANICAL: Cord around neck Prolapse cord Other cord entanglement or knot Cord compression: ☐ During labour Before labour Uterine rupture: Breech Face Compound Mal-presentation: Transverse Other, please specify __ Shoulder dystocia: 11.1.5. MATERNAL DISORDER: Pre-existing hypertensive disease Diabetes Other endocrine conditions (excluding diabetes) Obstetric cholestasis ☐ Thrombophilias Uterine anomalies Connective tissue disorders, please specify_____ Other, please specify___ 11.1.6. INFECTION: (confirmed by microbiology/placental histology) Bacterial Syphilis ☐ Viral diseases Maternal infection: Group B Streptococcus Protozoal Other, please specify organism ___ Ascending infection: Chorioamnionitis Other, please specify ___ 11.1.7. SPECIFIC FETAL CONDITIONS: Twin-twin transfusion Feto-maternal haemorrhage Non-immune hydrops Iso-immunisation Other, please specify_____ 7

11.1.8. SPECIFIC PLACENTAL CONDITIONS:

PLEASE NOTE THERE IS NO REQUIREMENT TO COMPLETE THIS SECTION SHOULD YOU WISH TO SUMIT AN ANONYMISED COPY OF THE PLACENTAL HISTOLOGY REPORT AS AN ATTACHMENT TO THIS FORM.

Please refer to the reference manual, page	10, for guidance on completing	this section.	
☐No abnormal histology reported			
<u>Chorioamnionitis</u> → Mi	d	Severe	
Fetal vasculitis → □Ar	terial	Both	
Maternal vascular malperfusion (utero	placental insufficiency)		
Distal villous hypoplasia	Placental hypoplasia		
Accelerated villous maturation	Ischaemic villous crowding		
☐ Placental infarction →	_		
☐ Placental Intarction →	Please specify approximate per	centage involved	
Retroplacental haemorrhage	Please specify approximate per	ercentage of maternal surface involved _	
Fetal vascular malperfusion:			
Please specify pathology			
Patchy hypoperfusion	Scattered avascular villi	☐ Thrombosis in fetal circulation	Fetal thrombotic vasculopathy
Cord pathology as sole finding Please specify pathology Hypercoiled cord Vasa praevia	☐ Hypocoiled cord	☐ Meconium associated va ☐ Other , please specify_	
— vasa praevia	i veramentous coru	Curer, prease specify_	
Cord pathology associated with dis			
please specify associated distal dis	ease: Thrombosis in fetal of	circulation	
☐ <u>Delayed Villous maturation defect</u>	_(distal villous immaturity/ delay	ed villous maturation)	
\square <u>Villitis</u> \rightarrow \square Low grade	☐High grade	☐With stem vessel oblitera	ition
Other, please specify			

I1.1.9. INTRA-UTERINE GROWTH RESTRIC	TION DIAGNOSIS MADE: YES
What was this based on? Please tick all that a	pply
Suspected antenatally Observed a	at delivery Observed at post-mortem
11.1.10. ASSOCIATED OBSTETRIC FACTOR	RS: Please tick all that apply
Birth trauma ☐ Intracranial haemorrhage	☐ Subgaleal haematoma
Fracture, please specify	
Other, please specify	
Intrapartum fetal blood sample result < 7.25	☐ Yes ☐ No
Polyhydramnios	Premature rupture of membranes
Prolonged rupture of membranes (> 24hours)	Amniocentesis
1	
Spontaneous premature labour	Other, please specify
	OR ASSOCIATED OBSTETRIC FACTORS PRESENT? YES D NO D
11.1.11. WERE THERE ANY ANTECEDENT Of the second of the se	egory as sparingly as possible
11.1.12. UNCLASSIFIED: Please use this cate CTION 12. MAIN CAUSE OF DEATH: STI 12.1. Which condition, indicated in Sectionsing or associated with the death. Please	egory as sparingly as possible
11.1.12. UNCLASSIFIED: Please use this cate CTION 12. MAIN CAUSE OF DEATH: STI 12.1. Which condition, indicated in Sections or associated with the death. Please (NB "non-MAIN" conditions are best described.	egory as sparingly as possible LL BIRTH & NEONATAL DEATHS on 11 as being present, was the MAIN condition or sentinel event se refer to the post-mortem and placental histology reports.
11.1.12. UNCLASSIFIED: Please use this cate CTION 12. MAIN CAUSE OF DEATH: STI 12.1. Which condition, indicated in Sections or associated with the death. Please (NB "non-MAIN" conditions are best described.	egory as sparingly as possible LL BIRTH & NEONATAL DEATHS on 11 as being present, was the MAIN condition or sentinel event se refer to the post-mortem and placental histology reports.
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2. Sources of information used to determ	LL BIRTH & NEONATAL DEATHS on 11 as being present, was the MAIN condition or sentinel event se refer to the post-mortem and placental histology reports. It as the "Other clinically relevant maternal or fetal conditions/ factors that were associated."
2. Sources of information used to determ	LL BIRTH & NEONATAL DEATHS on 11 as being present, was the MAIN condition or sentinel event se refer to the post-mortem and placental histology reports. It as the "Other clinically relevant maternal or fetal conditions/ factors that were associated as the cause of death?
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SECTION 13. NEONATAL DEATH ONLY: NEONATAL CONDITIONS ASSOCIATED WITH THE DEATH 13.1. Please TICK ALL the neonatal conditions causing and associated with the death. PLEASE REFER TO THE REFERENCE MANUAL. 13.1.1. MAJOR CONGENITAL ANOMALY: Central nervous system Cardiovascular system Respiratory system Gastro-intestinal system Musculo-skeletal anomalies ☐ Urinary tract Metabolic diseases Multiple anomalies Other major malformation, please specify Chromosomal disorder*, please specify _____ * In the event of a chromosomal disorder how was the diagnosis made? Ultrasound ☐ Clinically Genetic analysis * *See reference manual 13.1.1 (b) Was the diagnosis of major congenital anomaly confirmed/suspected before delivery by a Consultant **Fetal Medicine Specialist?** ∐No. \square Yes, in another unit, please specify name of unit $_$ П 13.1.2. PRE-VIABLE: (less than 22 weeks) 13.1.3. RESPIRATORY DISORDERS: Severe pulmonary immaturity Surfactant deficiency lung disease Pulmonary hypoplasia Meconium aspiration syndrome Chronic lung disease / Bronchopulmonary dysplasia (BPD) Primary persistent pulm. hypertension Uther (includes pulmonary haemorrhage), please specify_ 13.1.4. GASTRO-INTESTINAL DISEASE: Necrotising enterocolitis (NEC) Other, please specify 13.1.5. NEUROLOGICAL DISORDER: Hypoxic-ischaemic encephalopathy (HIE) *Intraventricular / Periventricular haemorrhage, please specify highest grade (0 – 4) * Hydrocephalus*, please tick all that apply: Acquired Communicating Obstructive * Congenital Other Other, please specify_ 13.1.6. INFECTION: Generalised (sepsis) Pneumonia Meningitis Please specify specific organism Other, specify _____ 10

13.1.7. INJURY / TRAUMA: (Postnatal)
Please specify
13.1.8. OTHER SPECIFIC CAUSES:
Malignancies / Tumours In-born errors of metabolism, please specify
Specific conditions, please specify
13.1.9. SUDDEN UNEXPECTED DEATHS:
Sudden Infant Death Syndrome (SIDS)
13.1.10. UNCLASSIFIED: (Use this category as sparingly as possible) \Box
13.2. Which condition, indicated in Section 13.1 as being present, was the MAIN condition causing or associated with the death. Please refer to the post-mortem report. In the absence of a post-mortem report, please refer to the death certificate. (NB "non-MAIN" conditions are best described as the "Other clinically relevant maternal or fetal conditions/ factors that were associated with but not necessarily causing the death").
Please tick all that apply Post Mortem Placental Histology Other, please specify
SECTION 14. DETAILS OF REPORTING UNIT (Please print) 14.1. Name of reporting unit:
14.2. Completed by
Name:
Staff Grade:
Work address:
Telephone Number: E-mail Address:
Date of Notification:
Thank you very much for taking the time to complete this form

Please return all completed forms to:
Ms Edel Manning, Project manager perinatal mortality audit, National Perinatal Epidemiology Centre Department of Obstetrics and Gynaecology 5th Floor Cork University Maternity Hospital Wilton Cork
If you have any queries regarding the Perinatal Death Notification Form, please contact us at the National Perinatal
Epidemiology Centre
Tel: (0)21 420 5042 E-mail: npec@ucc.ie

