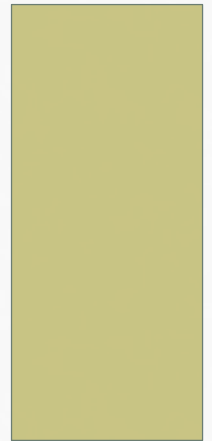


NATIONAL THROMBOSIS WEEK

IMPROVING PREVENTION AND BEST MANAGEMENT OF VTE-
REDUCING THE RISK IN PREGNANCY

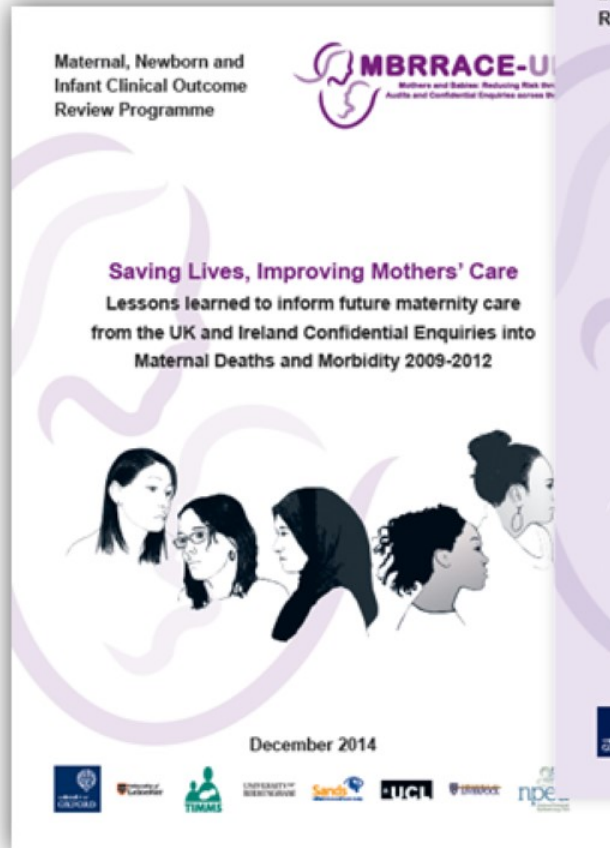


THROMBOEMBOLISM IN PREGNANCY

- **Pregnancy** increases the risk of venous **thromboembolism (VTE)** 4- to 5-fold over that in the nonpregnant state. The 2 manifestations of **VTE** are deep venous thrombosis (DVT) and pulmonary embolus (PE).

Maternal Mortality in the UK


2010-12




Maternal, Newborn and Infant Clinical Outcome Review Programme

MBRRACE-UK
Mothers and Babies: Reducing Risk Through Audits and Confidential Enquiries across the UK

Saving Lives, Improving Mothers' Care
Lessons learned to inform future maternity care from the UK and Ireland Confidential Enquiries into Maternal Deaths and Morbidity 2009-2012

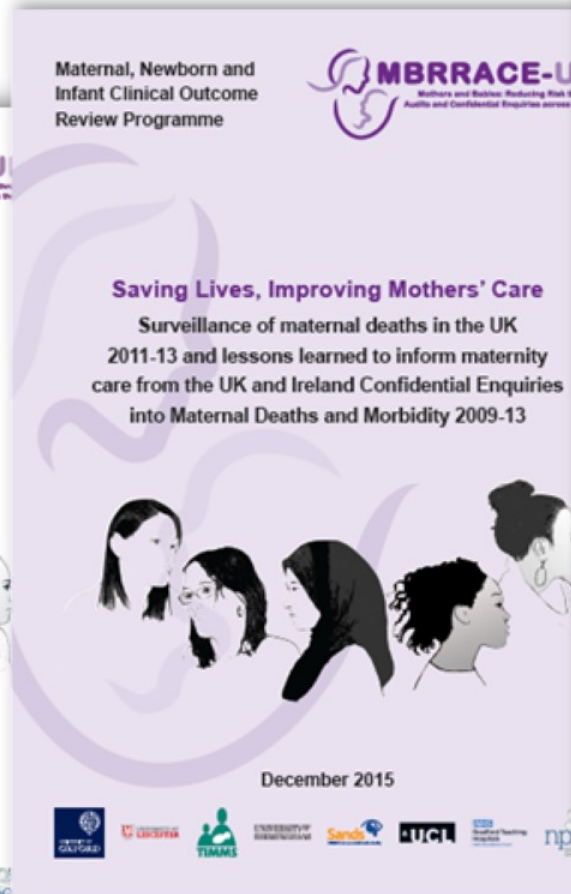


December 2014



10 per 100,000 maternities


2011-13




Maternal, Newborn and Infant Clinical Outcome Review Programme

MBRRACE-UK
Mothers and Babies: Reducing Risk Through Audits and Confidential Enquiries across the UK

Saving Lives, Improving Mothers' Care
Surveillance of maternal deaths in the UK 2011-13 and lessons learned to inform maternity care from the UK and Ireland Confidential Enquiries into Maternal Deaths and Morbidity 2009-13



December 2015



9 per 100,000 maternities

2012-14



Maternal, Newborn and Infant Clinical Outcome Review Programme

MBRRACE-UK
Mothers and Babies: Reducing Risk Through Audits and Confidential Enquiries across the UK

Saving Lives, Improving Mothers' Care
Surveillance of maternal deaths in the UK 2012-14 and lessons learned to inform maternity care from the UK and Ireland Confidential Enquiries into Maternal Deaths and Morbidity 2009-14



December 2016



8.5 per 100,000 maternities

BACKGROUND

- Maternal deaths have decreased, but there are still lessons we can learn
- Two thirds of women die from medical and mental health problems and one third from direct complications of pregnancy
- Three quarters of the women who died had medical or mental health problems before they become pregnant

CAUSES OF MATERNAL DEATH

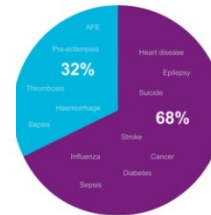
Key messages from the report



Maternal deaths have decreased

from **11** to **10** per 100,000 women giving birth
2008-09 2010-12

Causes of mothers' deaths



Women with pre-existing medical and mental health problems need:

- Pre-pregnancy advice
- Joint specialist and maternity care

Think Sepsis



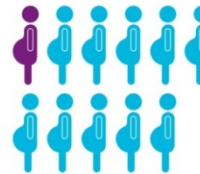
Almost a quarter of women who died had **Sepsis** (severe infection).

Women with sepsis need:

- Early diagnosis
- Rapid antibiotics
- Review by senior doctors and midwives

Prompt treatment and action can make the difference between life and death

Prevent Flu



1 in 11 of the women died from **Flu**
More than half of these women's deaths could have been prevented by a flu jab.

Flu vaccination will save mothers' and babies' lives

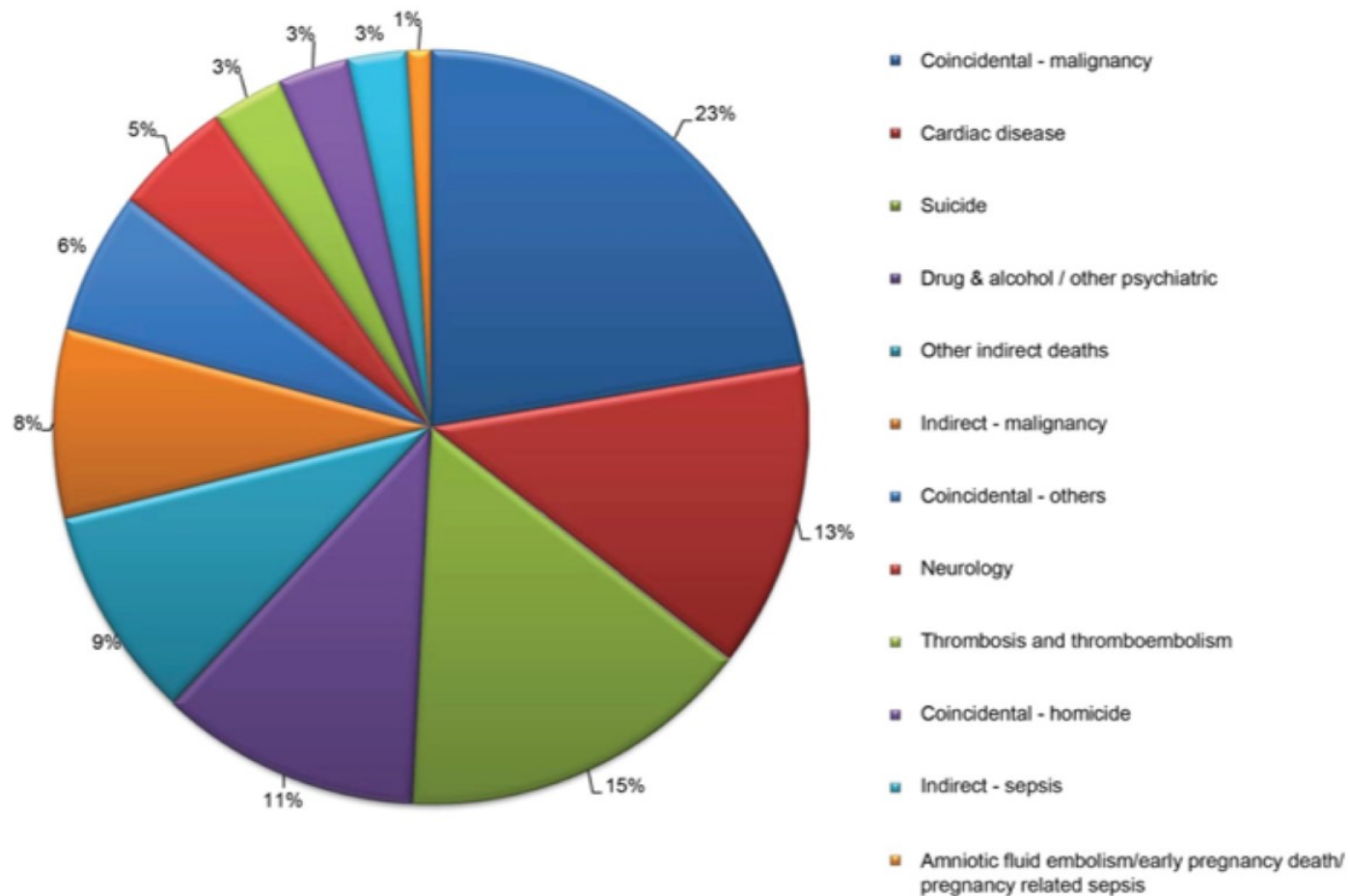
Causes of Death in women < 24 weeks UK & Ireland 2009-2014

| Cause of death | Number of women | Percentage of women |
|--------------------------------|-----------------|---------------------|
| Amniotic Fluid Embolism | 1 | 0.5 |
| Anaesthetic deaths | 1 | 0.5 |
| Pre-eclampsia and eclampsia | 1 | 0.5 |
| Sepsis | 19 | 10.0 |
| Thrombosis and thromboembolism | 22 | 11.5 |
| Cardiac disease | 24 | 12.6 |
| Mental health problems | 24 | 12.6 |
| Early pregnancy-related causes | 12 | 6.3 |
| Ectopic pregnancy | 9 | 4.8 |
| Legal termination of pregnancy | 2 | 1.0 |
| Self-attempted abortion | 1 | 0.5 |
| Haemorrhage | 2 | 1.1 |
| Neurology | 22 | 11.5 |
| Indirect deaths | 29 | 15.1 |
| Unascertained | 1 | 0.5 |
| Coincidental deaths | 33 | 17.3 |
| Total | 191 | 100 |

Conditions resulting in death

- **12 Deaths discussed here**
 - 9 women died as a result of ectopic pregnancies
 - 2 women died following legal termination of pregnancy
 - 1 woman died following a self-attempted abortion
- **7 Early deaths in other chapters**
 - 1 HELLP secondary to a molar pregnancy
 - 1 Cardiac death post termination
 - 2 Pulmonary Embolism after second trimester miscarriage
 - 3 Sepsis associated with miscarriage

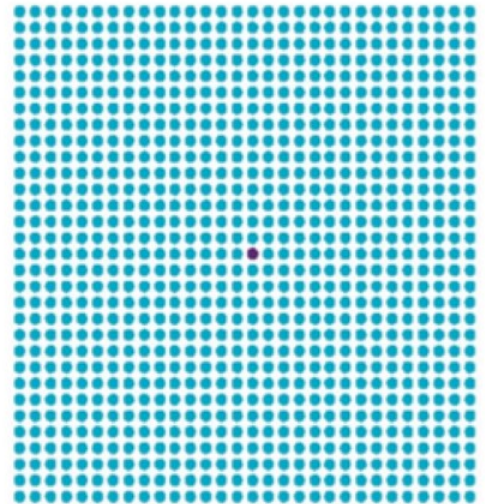
Late maternal deaths UK 2012-14



Direct Maternal Deaths 2012-14

- Thrombosis and thromboembolism the leading direct cause of death
 - 0.85 per 100,000 maternities
- Good care makes a difference

Less than **1 woman in every million** who gives birth now dies from **pre-eclampsia**



Maternal Morbidity and Mortality Annual Report Topic Cycle

- **2014:** Surveillance of maternal deaths in the UK 2009-12. Confidential enquiries on **sepsis morbidity** and deaths, haemorrhage, AFE, anaesthetic, neurological, respiratory, endocrine and other indirect deaths in the UK and Ireland.
- **2015:** UK surveillance 2011-13. Lessons for care from confidential enquiries of maternal deaths due to psychiatric, thrombosis, malignancy, late and coincidental deaths.
- **2016 (This report):** UK Surveillance 2012-14. Confidential enquiries on pre-eclampsia and eclampsia, **cardiac morbidity** and mortality, early pregnancy mortality, lessons for critical care.
- **2017:** UK Surveillance 2013-15. Confidential enquiries on sepsis, haemorrhage, AFE, anaesthetic, neurological, respiratory, endocrine and other indirect deaths, **morbidity from severe uncontrolled epilepsy and psychiatric morbidity.**
- **2018:** UK surveillance 2014-16. Lessons for care from confidential enquiries of maternal deaths due to psychiatric, thrombosis, malignancy, late and coincidental deaths, **morbidity from severe haemorrhage.**

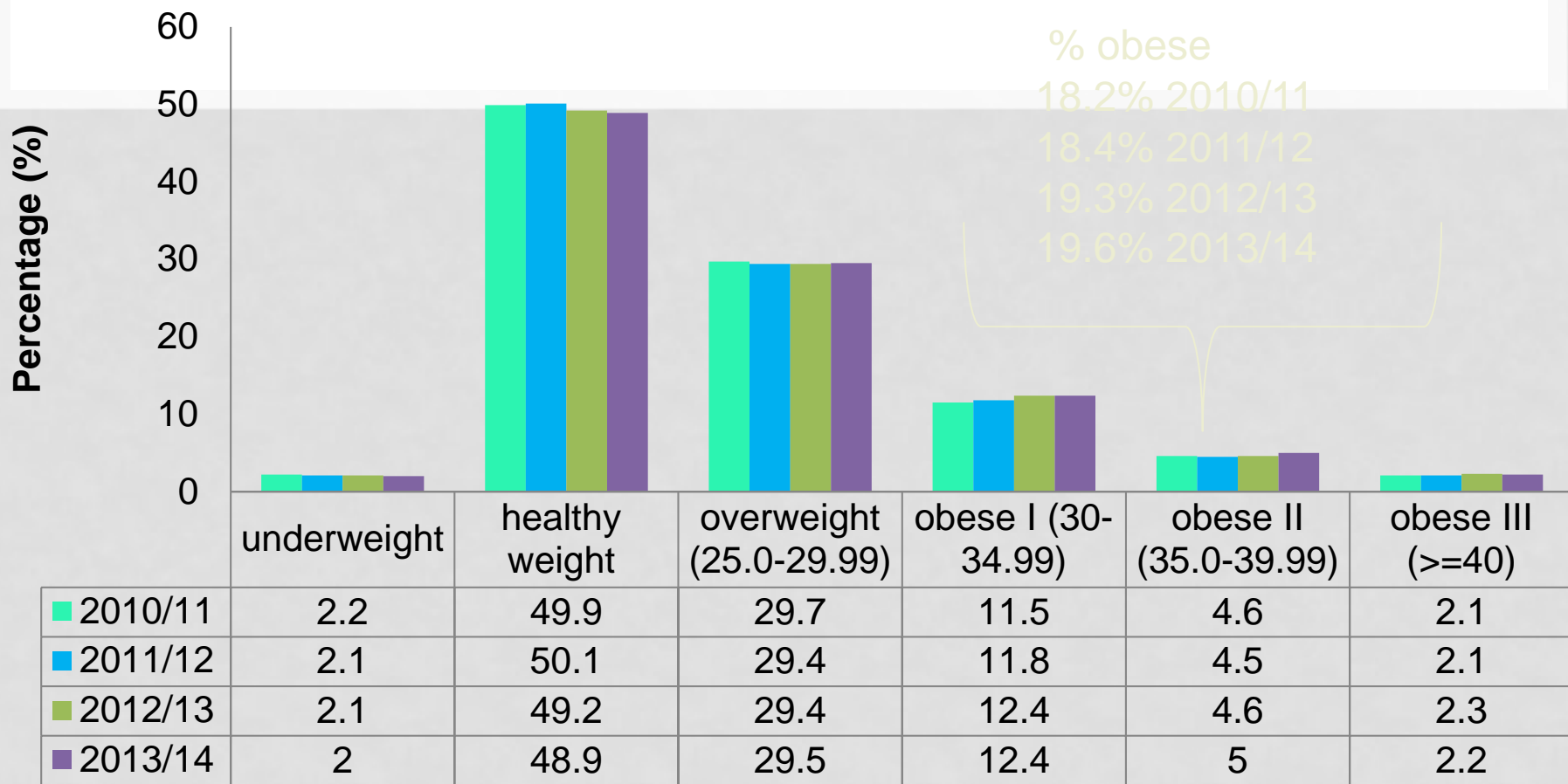
*Call for topics for 2019 morbidity enquiry open until 31/12/2016

CO-EXISTING MEDICAL COMPLICATIONS

- Nearly three quarters of women who died had a co-existing medical complication
- There has been no significant change in the rate of indirect maternal death of the last ten years, when the rate of deaths from direct causes has halved
- The rate of indirect maternal deaths (6.87 per 100,000 maternities) is now twice that of direct deaths (3.25 per 100,000 maternities)

Actions are urgently needed to address deaths from indirect causes

*WEIGHT STATUS IN PREGNANCY, 2010/11- 2013/14



Source: Results published in Children's health in Northern Ireland. Public Health Agency 2015.

LEARNING LESSONS TO IMPROVE CARE

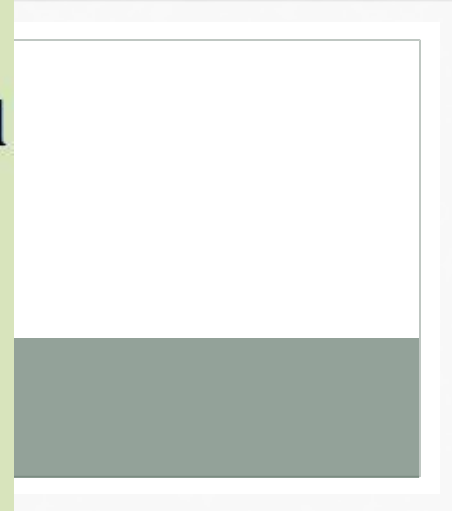
- We owe it to those left behind to learn from the death of their mother, partner, daughter or friend and to make changes for the future to prevent other women from dying



Northern Ireland's Regional Maternity Hand Held Record

Operational Guidance

Dr Briega M Lagan with
Ms Brenda Devine and
Ms Verena Wallace



Antenatal VTE Risk Assessment – Booking (Risk assessment to be completed at booking)

Name:

ID No.:

Antenatal assessment and management (to be assessed at booking and repeated if admitted)

Any previous VTE except a single event related to major surgery

Hospital admission
 Single previous VTE related to major surgery
 High risk thrombophilia - no VTE
 Medical comorbidities e.g. cancer, heart failure, active SLE, IBD or inflammatory polyarthropathy, nephrotic syndrome, type I DM with nephropathy, sickle cell disease, current IVU
 Any surgical procedure e.g. appendectomy
 OHS (first trimester only)

Obesity (BMI > 30 kg/m²)
 Age > 35
 Parity > 3
 Smoker
 Gross varicose veins
 Current pre-eclampsia
 Immobility, e.g. paraplegia, PGP
 Family history of unprovoked or estrogen-provoked VTE in first-degree relative
 Low risk thrombophilia
 Multiple pregnancy
 IVT/ART
 Transient risk factors: Dehydration/hypertension, current systemic infection, long distance travel

HIGH RISK
 Requires antenatal prophylaxis with LMWH
 Refer to trust-nominated thrombosis in pregnancy expert/ team

INTERMEDIATE RISK
 Consider antenatal prophylaxis with LMWH

Four or more risk factors:
 prophylaxis from first trimester
 Three risk factors:
 prophylaxis from 28 weeks

Fewer than three risk factors

LOWER RISK
 Mobilisation and avoidance of dehydration

| Weight | AN and PN prophylactic dose of LMWH |
|--------------|-------------------------------------|
| < 50 kg | 20 mg OD |
| 50 – 90 kg | 40 mg OD |
| 91 – 130 kg | 60 mg OD |
| 131 – 170 kg | 80 mg OD (or 40 mg BD) |
| > 170 kg | 0.6 mg / kg / day |

Total Risk Factor Score: _____

Comment: _____

Signature and Profession: _____ Date: _____ Time: _____

Prophylaxis dose of LMWH provided

Yes No

Antenatal VTE Risk Assessment – Booking (Risk assessment to be completed at booking)

Risk assessment for venous thromboembolism (VTE)

- If total score > 4 antenatally, consider thromboprophylaxis from the first trimester
- If total score > 2 antenatally, consider thromboprophylaxis from 28 weeks
- If total score > 2 postnatally, consider thromboprophylaxis for at least 6 days
- If admitted to hospital antenatally consider thromboprophylaxis
- If prolonged admission (> 3 days) or readmission to hospital within the puerperium consider thromboprophylaxis

For patients with an identified bleeding risk, the balance of risks of bleeding and thrombosis should be discussed in consultation with a haematologist with expertise in thrombosis and bleeding in pregnancy

Risk factors for VTE

| Pre-existing risk factors | Risk | Score |
|---|------|---------|
| Previous VTE (except a single event related to major surgery) | | 4 |
| Previous VTE provoked by major surgery | | 1 |
| Known high-risk thrombophilia | | 1 |
| Medical comorbidities e.g. cancer, heart failure, active systemic lupus erythematosus, inflammatory polyarthropathy or inflammatory bowel disease, nephrotic syndrome, type I diabetes mellitus with nephropathy, sickle cell disease, current intravenous drug use | | 1 |
| Family history of unprovoked or estrogen-related VTE in first-degree relative | | 1 |
| Known low-risk thrombophilia (no VTE) | | 0* |
| Age < 35 years | | 1 |
| Obesity | | 1 or 2* |
| Parity > 3 | | 1 |
| Smoker | | 1 |
| Gross varicose veins | | 1 |
| Gynaecologic risk factors | | |
| Pre-eclampsia in current pregnancy | | 1 |
| ART/IVF (current only) | | 1 |
| Multiple pregnancy | | 1 |
| Obstetric practice in labour | | 1 |
| Elective caesarean section | | 1 |
| Mid-early or rotational operative delivery | | 1 |
| Prolonged labour (> 24 hours) | | 1 |
| PPH (> 1000 ml or transfusion) | | 1 |
| Protein loss > 3000 mg in current pregnancy | | 1 |
| Mitochondria in current pregnancy | | 1 |
| Transient risk factors | | |
| Any surgical procedure in pregnancy or puerperium except obstetrical repair of the perineum, e.g. episiotomy, haemorrhoidectomy, perineorrhaphy | | 1 |
| Hypertension | | 1 |
| OHS (first trimester only) | | 4 |
| Current systemic infection | | 1 |
| Immobility, dehydration | | 1 |
| TOTAL | | |

Abbreviations: ART assisted reproductive technology, IVF in vitro fertilisation, OHS ovarian hyperstimulation syndrome, VTE venous thromboembolism.

*If the known low risk thrombophilia is in a woman with a family history of VTE in a first-degree relative postpartum thromboprophylaxis should be continued for 6 weeks.

*BMI < 30 = 1, BMI > 30 = 2

Antenatal VTE Risk Assessment – Hospitalised

(Risk assessment to be completed on EVERY admission)

Name:

ID No:

Antenatal assessment and management (to be assessed at booking and repeated if admitted)

Any previous VTE except a single event related to major surgery

HIGH RISK
Requires antenatal prophylaxis with LMWH
Refer to trust-nominated thrombosis in pregnancy expert/team

Hospital admission

Single previous VTE related to major surgery

High-risk thrombophilia + no VTE

Medical comorbidities e.g. cancer, heart failure, active SLE, IBD or inflammatory polyarthropathy, nephrotic syndrome, type 1 DM with nephropathy, sickle cell disease, current IVDU

Any surgical procedure e.g. appendicectomy

OHSS (first trimester only)

INTERMEDIATE RISK
Consider antenatal prophylaxis with LMWH

Obesity (BMI > 30 kg/m²)

Age > 35

Parity ≥ 3

Smoker

Gross varicose veins

Current pre-eclampsia

Immobility, e.g. paraplegia, PGP

Family history of unprovoked or estrogen-provoked VTE in first-degree relative

Low-risk thrombophilia

Multiple pregnancy

IVF/ART

Transient risk factors: Dehydration/hyperemesis; current systemic infection; long-distance travel

Four or more risk factors:
prophylaxis from first trimester

Three risk factors:
prophylaxis from 28 weeks

Fewer than three risk factors

LOWER RISK
Mobilisation and avoidance of dehydration

| Weight | AN and PN prophylactic dose of LMWH |
|--------------|-------------------------------------|
| < 50 kg | 20 mg OD |
| 50 – 80 kg | 40 mg OD |
| 81 – 100 kg | 60 mg OD |
| 101 – 170 kg | 80 mg OD (or 40 mg BD) |
| > 170 kg | 0.6 mg / kg / day |

| Prophylaxis dose of LMWH prescribed | |
|-------------------------------------|-----------------------------|
| <input type="checkbox"/> Yes | <input type="checkbox"/> No |

| | |
|--------------------------|------|
| Total Risk factor Score: | |
| Comment: | |
| Signature and Profession | Date |
| | Time |

Antenatal VTE Risk Assessment – Hospitalised

(Risk assessment to be completed on EVERY admission)

Risk assessment for venous thromboembolism (VTE)

- If total score ≥ 4 antenatally, consider thromboprophylaxis from the first trimester.
- If total score ≥ 3 antenatally, consider thromboprophylaxis from 28 weeks.
- If total score ≥ 2 postnatally, consider thromboprophylaxis for at least 10 days.
- If admitted to hospital antenatally consider thromboprophylaxis.
- If prolonged admission (> 3 days) or readmission to hospital within the puerperium consider thromboprophylaxis.

For patients with an identified bleeding risk, the balance of risks of bleeding and thrombosis should be discussed in consultation with a haematologist with expertise in thrombosis and bleeding in pregnancy.

Risk factors for VTE

| Pre-existing risk factors | Tick | Score |
|---|------|---------------------|
| Previous VTE (except a single event related to major surgery) | | 4 |
| Previous VTE provoked by major surgery | | 3 |
| Known high-risk thrombophilia | | 3 |
| Medical comorbidities e.g. cancer, heart failure, active systemic lupus erythematosus, inflammatory polyarthropathy or inflammatory bowel disease, nephrotic syndrome, type 1 diabetes mellitus with nephropathy, sickle cell disease, current intravenous drug use | | 3 |
| Family history of unprovoked or estrogen-related VTE in first-degree relative | | 1 |
| Known low-risk thrombophilia (no VTE) | | 1* |
| Age (> 35 years) | | 1 |
| Obesity | | 1 or 2 [†] |
| Parity ≥ 3 | | 1 |
| Smoker | | 1 |
| Gross varicose veins | | 1 |
| Obstetric risk factors | | |
| Pre-eclampsia in current pregnancy | | 1 |
| ART/IVF (antenatal only) | | 1 |
| Multiple pregnancy | | 1 |
| Caesarean section in labour | | 2 |
| Elective caesarean section | | 1 |
| Mid-cavity or rotational operative delivery | | 1 |
| Prolonged labour (> 24 hours) | | 1 |
| PPH (> 1 litre or transfusion) | | 1 |
| Preterm birth < 32 nd weeks in current pregnancy | | 1 |
| Stillbirth in current pregnancy | | 1 |
| Transient risk factors | | |
| Any surgical procedure in pregnancy or puerperium except immediate repair of the perineum, e.g. appendicectomy, postpartum sterilisation | | 3 |
| Hyperemesis | | 3 |
| OHSS (first trimester only) | | 4 |
| Current systemic infection | | 1 |
| Immobility, dehydration | | 1 |
| TOTAL | | |

Abbreviations: ART assisted reproductive technology; IVF in vitro fertilisation; OHSS ovarian hyperstimulation syndrome; VTE venous thromboembolism.

*If the known low-risk thrombophilia is in a woman with a family history of VTE in a first-degree relative postpartum thromboprophylaxis should be continued for 6 weeks.

[†]BMI ≥ 30 = 2; BMI ≥ 40 = 3



THANK YOU

QUESTIONS?

