Evidence for GAIN Guideline for Admission to Midwife-led units in Northern Ireland June 2015

This guideline was developed in accordance with the GAIN Advice for Guideline Development in Northern Ireland and is to assist women and maternity care providers in their decision-making with regard to place of birth for women with a straightforward singleton pregnancy at the point of labour. A team of health professionals, lay representatives and technical experts known as the Guideline Development Group (GDG) with support from GAIN, were established following requests for nominations from main stakeholder organisations. These included maternity service providers, women's and parents' groups, for example: Heads of Midwifery, Midwives, Consultant Obstetricians, Consultant Anaesthetists from the HSC Trusts, a GP, Midwifery Advisor, a representative from the Public Health Agency, NI Practice and Education Council, Royal College of Midwives, Sure start, Parenting NI and Mothers' Voice (a HSC Maternity Service Liaison Committee).

The consensus of the GDG was that if there is any uncertainty regarding admission or care, multidisciplinary discussion is necessary with appropriate documentation.

From the outset, the GDG defined a straightforward pregnancy as:

'a singleton pregnancy, in which the woman does not have any pre-existing condition impacting on her pregnancy, a recurrent complication of pregnancy or a complication in this pregnancy which would require on-going consultant input, has reached 37 weeks gestation and \leq Term +15' (p.3)

The guideline was developed as the result of an in-depth iterative process, which utilised expert professional, experiential knowledge and a range of robust evidence. We undertook a structured search and review of literature on a range of topics areas identified by the GDG as areas where guidance was required, resulting in the published guideline criteria for admission to midwife-led units in Northern Ireland (GAIN, 2016). The search strategy involved a search of CINAHL, Cochrane, PubMed, Medline, Maternity and Infant Care Databases. Back-chaining of reference lists from relevant papers and documents and an online search of departmental strategic and professional resources was also undertaken (See Table 1).

Table 1 Strategic and Professional Resources

American Nurse Midwifery Association (www.midwife.org) Department of Health Social Services and Public Safety (www.dhsspsni.org.uk) Guidelines and Audit Implementation Network (www.gain-ni.org) National Institute for Health and Care Excellence (www.nice.org.uk/) Regulation and Quality Improvement Authority (www.rqia) Royal College of Midwives (www.rcm.org.uk) Royal College of Obstetricians and Gynaecologists (www.rcog.org.uk)

(See evidence tables relating to each criterion for admission to midwife-led units in Northern Ireland)

Inclusions and Exclusions

The Guideline Development Group discussed key evidence utilising the expertise from clinicians and users by experience. The evidence for the guideline was focused on the population of women of childbearing age (16 years +) with straightforward singleton pregnancy at the point of labour who planned to birth in a midwife led unit. In total, there were 20 criteria included in the final version of the guideline (although some other criteria such as Gestational Diabetes were considered for inclusion but following discussion by the GDG were excluded). The final criteria were focused on being as inclusive as possible for pregnant women and the review questions were therefore based on the following:

- 1. Maternal age at booking
- 2. BMI at booking
- 3. Hb level
- 4. Number of previous births
- 5. Assisted conception with Clomifene or similar
- 6. SROM \leq 24hrs & no signs of infection
- 7. Women on Tier 1 and Tier 2 of the Integrated Perinatal Mental Health Care Pathway
- 8. Threatened miscarriage, now resolved
- 9. Threatened preterm labour, now resolved
- 10. Suspected low lying placenta, now resolved
- 11. Medical condition that is not impacting on the pregnancy or the woman's health
- 12. Women who have required social services input and there is no related impact on the pregnancy or the woman's health

- 13. Previous congenital abnormality, with no evidence of reoccurrence
- 14. Non-significant (light) meconium in the absence of any other risk
- 15. Uncomplicated third degree tear/previous extensive vaginal, cervical, or third degree perineal trauma following individual assessment
- 16. Serum antibodies of no clinical significance
- 17. Women who have had previous cervical treatment, now term
- 18. Previous PPH, not requiring blood transfusion or surgical intervention
- 19. Prostaglandin induction resulting in the onset of labour
- 20. Group B Streptococcus positive in this pregnancy with no signs of infection

The GDG judged the evidence against their suitability for inclusion as criteria for Alongside Midwifery Unit (AMU) or a Free standing Midwifery Unit (FMU).

Strengths and Limitations of the Body of Evidence

In recent years, there has been an increase in the number of high quality studies and professional guidelines which support the provision of care for women who plan to birth in Midwife led units rather than obstetric units. The limitations of the current body of evidence does not allow for the GDG to include women with complex pregnancies, e.g. gestational diabetes. Therefore, further high quality research is required and will be included if available in future updates of the guideline.

Consensus Methods Used for Literature and Recommendations

Throughout the process of the review, consultation with a wide range of stakeholders informed the development of the criteria and guidelines. However, each of the 20 criteria were discussed with the Guideline Development Group members and agreement reached through consensus.

Dates Searches Were Undertaken

Searches were undertaken from January 2009 to November 2014 and only those papers written in English were included. In addition, the GDG members were encouraged to bring additional published evidence that came to light during the guideline development process from the national and international links.

Declaration of Interests

At the start of the guideline development process all GDG members' interests were recorded on a standard declaration form that covered consultancies, fee-paid work, share-holdings, fellowships and support from the healthcare industry. At all subsequent GDG meetings, members were required to declare any new or arising conflicts of interest. No conflicts of interest were declared for this guideline.

| Table A. | Maternal age at booking ≥ 16 yrs & ≤40 yrs (FMU & AMU)/<16 or >40 (AMU) |
|----------|---|
| Table B. | BMI at booking ≥ 18kg/m ² & ≤35 kg/m ² (FMU & AMU)/≥35 kg/m ² & ≤40 kg/m ² with good mobility |
| Table C | Last recorded Hb≥100g/I (FMU & AMU)/Hb >85g/I (AMU) |
| Table D | No more than 4 previous births (FMU & AMU)/No more than 5 previous births (AMU) |
| Table E. | Assisted conception with Clomifene or similar (FMU & AMU)/IVF pregnancy at term (excluding ovum donation & maternal |
| | age >40 years) (AMU) |
| Table F | SROM ≤ 24hrs & no signs of infection (FMU & AMU)/ SROM > 24hrs, in established labour & no signs of Infection (AMU) |
| Table G | Women on Tier 1 of the Integrated Perinatal Mental Health Care Pathway (FMU & AMU)/Women on Tier 2 of the Perinatal |
| | Mental Health Care Pathway following individual assessment (AMU) |
| Table H | Threatened miscarriage, now resolved (FMU & AMU) |
| Table I | Threatened preterm labour, now resolved (FMU & AMU) |
| Table J | Suspected low lying placenta, now resolved (FMU & AMU) |
| Table K | Medical condition that is not impacting on the pregnancy or the woman's health (FMU & AMU) |
| Table L | Women who have required social services input and there is no related impact on the pregnancy or the woman's health |
| | (FMU & AMU) |
| Table M | Previous congenital abnormality, with no evidence of reoccurrence (FMU & AMU) |
| Table N | Non-significant (light) meconium in the absence of any other risk (FMU & AMU) |
| Table O | Uncomplicated third degree tear (FMU & AMU)/Previous extensive vaginal, cervical, or third degree perineal trauma |
| | following individual assessment (AMU) |
| Table P | Serum antibodies of no clinical significance (FMU & AMU) |
| Table Q | Women who have had previous cervical treatment, now term (FMU & AMU) |
| Table R | Previous PPH, not requiring blood transfusion or surgical intervention (AMU) |
| Table S | Prostaglandin induction resulting in the onset of labour (AMU) |
| Table T | Group B Streptococcus positive in this pregnancy with no signs of infection (AMU) |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comments/Reference |
|-------------------------|---------------------|-------------|--|--|
| NICE (2014a), Uk | Guideline CG | Age | Indicated individual assessment when planning place of birth | Table 9 |
| | 190 | | if >35 years of age on booking. | |
| RCOG (2010), | Green Top | - | Point up association of advanced maternal age with IUFD; | Fretts (2005); Froen (2001). |
| UK | Guideline No | | age not specified; specific recommendation for care not | |
| | 55 | | provided. | |
| | | | The adjusted odds ratio for age ≥35 years compared with | |
| | | | ≤25 years was 5.1 (95% confidence interval). | |
| MBRRACE | National | - | Increased rates of stillbirth, neonatal mortality and extended | P. 55: Mothers characteristics |
| (2013), UK | Surveillance | | perinatal mortality were seen in both the youngest (<20 | CEMACH (2009) |
| | data Report | | years) and the oldest (>40 years) mothers | |
| RQIA (2012), NI | A Review: | <16 year | Recommended age limits on children moving to adult wards. | PLEASE NOTE: |
| | baseline | (AMU) | Sought assurances that up to date British National Formulary | The Guideline Development Group |
| | assessment to | ≥ 16 (FMU & | for Children is available on adult wards so that staff can | (GDG) advised that regarding |
| | inform the | AMU) | ensure medication decisions are correct. | management of teenage girls in |
| | provision of | | | labour, AMU's are suitable for planned |
| | care & | | | place of birth (although all other risk |
| | admission of | | | factors need to be addressed) - In the |
| | children to adult | | | case of a pregnant teenager who is |
| | wards | | | under 16 requiring intravenous fluids |
| | | | | in labour, the paediatric protocol is to |
| | | | | be followed and care transferred to a |

Table A: BMI at booking ≥ 18kg/m2 & ≤35 kg/m2 (FMU & AMU)/ ≥35 kg/m2& ≤40 kg/m2 with good mobility

| Author/ | Type of | Criteria | Results/Findings | Comments/Reference |
|---------------|--------------|----------|--|---|
| Date/Country | Evidence | | | |
| NICE (2010), | PH Guideline | BMI | 'Lack of evidence on the underlying mechanism linking the | Recommendation 2 Pregnant Women |
| UK | 27 | | gestational weight gain and pregnancy outcome' | Do not weigh women repeatedly during |
| | | | | pregnancy as a matter of routine. Only |
| | | | | weigh again if clinical management can be |
| | | | | influenced or if nutrition is a concern |
| | | | | |
| NICE (2014a), | Clinical | | One of the factors indicating increased risk with suggested | Table 7: Other factors indicating |
| UK | Guideline | | planned birth at an obstetric unit is a 'BMI at booking of greater | increased risk suggesting planned birth |
| | 190 | | than 35 kg/m2 '. | at an obstetric unit (P.19) |
| | | | BMI greater than 35 kg/m2 considered a risk for PPH. | P.74 |
| NICE (2014c) | Clinical | - | Maternal weight and height should be measured at the booking | P 22: Sections 1 5 1 1 |
| | | | anneintment, and the warran's hady mass index should be | |
| UK | Guideline 62 | | appointment, and the woman's body mass index should be | 1.5.1.2 |
| | | | calculated (weight [kg]/height[m]2)'. | |
| | | | Repeated weighing during pregnancy should be confined to | |
| | | | circumstances in which clinical management is likely to be | |
| | | | influenced'. | |
| | | | Women who have a body mass index 30 kg/m2 or above at their | P. 47 Appendix C: Women |
| | | | booking appointment or are underweight (body mass index below | requiring additional care |
| | | | 18 kg/m2 at first contact) require additional care | |
| | | | | |
| RCOG (2010), | Green Top | 1 | Point up association of obesity with IUFD; specific | Arendas (2008);Fretts (2005); Froen |
| UK | Guideline No | | recommendation for care not provided | (2001); Huang (2000) |
| | 55 | | | |

Table B: BMI at booking ≥ 18kg/m2 & ≤35 kg/m2 (FMU & AMU)/ ≥35 kg/m2& ≤40 kg/m2 with good mobility (AMU)

| Hollowell | Systematic | Risks of complications during childbirth go up with increasing P348 |
|---------------|---------------|---|
| (2013), UK | Review | BMI among these healthy women. But the researchers found the |
| | | increase was modest. Overweight, obese or very obese women |
| | | showed a relative increase in risk of 6-12% compared to women |
| | | with a normal BMI. Most of the additional risk of needing |
| | | treatment was accounted for by giving drugs for slow labour |
| | | among those with a high body mass index, though risks of |
| | | serious outcomes for mother and baby were also increased. |
| | | However, whether a woman has had a previous baby plays a |
| | | larger role in influencing the chances of needing hospital care. |
| | | 53% of women of normal weight having their first baby had an |
| | | intervention or a complication at birth. The figure was 21% for |
| | | very obese but otherwise healthy women having a second or |
| | | subsequent baby. |
| | | Otherwise healthy multiparous women obese women may have P 343 |
| | | lower antepartum risks than previously appreciated |
| Aune (2014), | Systematic | Evidence suggests that maternal obesity increases the risk P 1536 |
| UK | Review & | of fetal death, stillbirth, and infant death; however, the optimal |
| | Meta Analysis | body mass index (BMI) for prevention is not known'. |
| | | 'The optimal prepregnancy BMI to prevent fetal and infant |
| | | death has not been established'. |
| NICE (2014c), | Clinical | 'Maternal weight and height should be measured at the booking P22 |
| UK | Guideline 62 | appointment, and the woman's body mass index should be |
| | | calculated (weight [kg]/height[m]2)'. |

| Lawrence et al, | Cochrane | Importance | Upright and ambulant positions versus recumbent positions | P.2 |
|-----------------|----------|-------------|---|-----|
| (2013), | Review | of Mobility | and bed care: women randomised to upright positions in | |
| Australia | | | labour have shorter labours and less likely to have caesarean | |

Table C: Last recorded Hb≥100g/I (FMU/AMU)/ Hb>85g/I (AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comments/Reference |
|-------------------------|---------------------|----------|--|---------------------------------------|
| NICE (2014a), UK | Clinical | Hb | Haemoglobin of 85-105g/litre at onset of labour; | Page 19, Table 8 Medical Conditions |
| | Guideline | | | indicating individual assessment when |
| | 190 | | | planning place of birth |
| | | | Maternal Haemoglobin below 85g/l at onset of labour should | P.74; 1.14.29 |
| | | | deliver in an obstetric unit | |
| | | | Maternal Haemoglobin level below 85 g/litre at onset of labour a risk factor for PPH; advise give birth in Obstetric unit | P.74 |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|------------|---|----------------------------------|
| NICE (2014a), UK | Clinical | 4 Previous | 'Grand multiparity (parity 4 or more) is considered a risk for PPH; | Page 74; Section 1.14.29 The GDG |
| | Guideline | Births | advise give birth in Obstetric unit. | members agreed that the GAIN NI |
| | 190 | | | MLU Guidelines would use the |
| | | | | terminology '5 previous births' |
| | | | | which equates to a Parity of 4. |
| | | | | |
| NICE (2014a), UK | | | Para 4 or more; Other factors indicating individual assessment when | Table 9 |
| | | | planning place of birth | |

Table D: No more than 4 previous births (FMU/AMU)/ No more than 5 previous births (AMU)

Table E: Assisted conception with Clomifene or similar (FMU & AMU)/ IVF pregnancy at term (excluding ovum donation & maternal age >40 years) (AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------------|---------------------|---------------|--|--------------------------------------|
| Okun (2014), | Systematic review - | Assisted | Increased evidence that infertility or subfertility is an | P.64 |
| Canada | Clinical Practice | Conception | increased risk factor for obstetrical complications & | |
| | Guideline | | adverse perinatal outcomes, even without the addition of | |
| El-Chaar | Retrospective | Assisted | Increased risk of birth defects associated with Assisted | P.1557 |
| (2009), Canada | Cohort Study | Human | Human Reproduction (AHR) and the risks are higher with | |
| | | Reproduction | In Vitro Fertilisation (IVF) and (IUI)'. | |
| RCOG (2012), UK | Scientific Impact | In Vitro | Increasing maternal age is a risk factor for almost all | P. 6 |
| | Paper | Fertilisation | pregnancy and perinatal complications. The average age at | The Guideline Development |
| | | | which women attempt to conceive continues to rise and | Group using expert professional, |
| | | | consequently IVF is increasingly used by older women | local and user knowledge |
| | | | who are already predisposed to pregnancy complications. | advocated for inclusion of the 'age' |
| | | | However, even when comparing age matched controls | limit and 'at term' wording within |
| | | | there appears to be an increased risk of complications | this criteria. |
| | | | associated with infertility, with a higher rate of caesarean | |
| | | | section delivery, obstetric haemorrhage, pre-eclampsia, | |
| | | | pregnancy-induced hypertension and gestational diabetes | |
| | | | all noted in older women having IVF'. | |
| Wiggins & Main (2005), USA | Retrospective | Donor egg & | Increased risk of pregnancy induced hypertension | GDG therefore specified the |
| Wall (2003), USA | | | utilising donor egg nowever excellent outcomes can still | |
| | | | | matemai age >40 |

Table F: SROM ≤ 24hrs & no signs of infection (AMU/FMU)/ SROM > 24hrs, in established labour & no signs of infection (AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|----------|---|---------------------------|
| NICE (2014a), UK | Clinical | | If no signs of infection in the woman, do not give antibiotics to | |
| | Guidelines, | | either mother or the baby, even if membranes ruptured >24 | |
| | CG 190a | | hours. | P.882 |
| | | | If evidence of infection, prescribe a full course of broad spectrum | |
| | | | Intravenous antibiotics. | |
| | | | Closely observe and assess any baby born to a woman with | Recommendation 278, P 821 |
| | | | prelabour rupture of membranes that is > 24 before established | |
| | | | labour. | |

Table G: Women on Tier 1 of the Integrated Perinatal Mental Health Care Pathway (AMU/FMU)/Women on Tier 2 of thePerinatal Mental Health Care Pathway following individual assessment (AMU)

| Author/ | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|------------------|---------------------|---------------|--|-------------------|
| Date, oountry | LVIGENCE | | | |
| SE Trust,(2013), | Care Pathway | Mental Health | Advice based on NICE Clinical Guideline 192 which categorises | P 3-4 |
| NI | | | women into Tiers 1-4 with recommendations for management. | |
| NICE (2014b), UK | Clinical | | All healthcare professionals providing assessment and | Section 17.1 |
| | Guideline 192 | | interventions for mental health problems in pregnancy and the | |
| | | | postnatal period should understand the variations in their | |
| | | | presentation and course at these times, how these variations | |
| | | | affect treatment, and the context in which they are assessed and | |
| | | | treated (for example, maternity services, health visiting and | |
| | | | mental health services)'. | |
| | | | | |
| NICE (2014a), UK | Clinical | | Indicate individual assessment when planning place of birth if | Table 9 |
| | Guideline 190 | | under current outpatient psychiatric care. | |
| | | | Indicate increased risk suggesting planned birth at an obstetric | Table 6 |
| | | | unit. | |

Table H: Threatened miscarriage, now resolved (AMU/FMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|-------------|--|--------------------------------|
| NICE (2014a), UK | Clinical | Threatened | Threatened miscarriage (i.e < 24 weeks), can plan to birth in | Table 9; The criteria makes it |
| | Guideline 190 | Miscarriage | MLU Indicate individual assessment when planning place of | clear that it applies to |
| | | | birth if antepartum bleeding of unknown origin (single episode | 'threatened miscarriage; now |
| | | | after 24 weeks gestation). | resolved' (i.e. woman now at |
| | | | Other factors indicating increased risk suggesting planned birth | Table 7 |
| | | | at an obstetric unit-Recurrent antepartum Haemorrhage | |

Table I: Threatened preterm labour, now resolved (AMU/FMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|------------------------------|-------------------------------------|---|---|
| NICE (2014a), UK | Clinical Guideline 190 | <i>Threatened</i> Preterm labour | NICE 2014 CG 190- Preterm labour or preterm prelabour rupture of membranes are factors indicating increased risk Suggesting planned birth at an obstetric unit. However, the criteria in the GAIN MLU guidelines relate to 'threatened preterm labour'- see comments. | Table 7; The GDG members agreed that the GAIN NI MLU Guidelines would include woman who experienced a <i>threatened</i> preterm labour and her pregnancy has now reached term |

Table J: Suspected low lying placenta, now resolved

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|-----------|--|--------------------------------|
| NICE (2014c), UK | Clinical | Low-lying | Because most low-lying placenta detected at the routine anomaly | Section 1.9.4 Placenta Praevia |
| | Guideline 62 | placenta | scan will have resolved by the time the baby is born, only a woman | (low-lying placenta) 1.9.4.1 |
| | | | whose placenta extends over the internal cervical os should be | |
| | | | offered another trans abdominal scan at 32 weeks' | |

| Author/Date/ Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|------------|--|-------------------------------|
| NICE (2014a), UK | Clinical | Medical | Individual assessment for medical conditions needed when | Table 8. P.19 |
| | Guideline 190 | Conditions | planning place of birth. However, the GAIN MLU guideline applies | As there are numerous |
| | | | specifically to a medical condition that is not impacting on the | medical conditions, it is not |
| | | | pregnancy or the woman's health. However, it is also | possible to give specific |
| | | | recommended that –' If any uncertainty, multidisciplinary | advice for each one (many |
| | | | discussion is necessary, with appropriate documentation' | may have no impact on a |
| | | | | pregnancy or a woman's |
| | | | | health); therefore individual |
| | | | | assessment is required to |
| | | | | • |

Table K: Medical condition that is not impacting on the pregnancy or the woman's health (FMU & AMU)

Table L: Women who have required social services input and there is no related impact on the pregnancy or the woman's health (FMU/AMU)

| Author/ T Date/Country Ev | Type of vidence | Criteria | Results/Findings | Comment/Reference |
|------------------------------------|-----------------------------|--------------------|---|--|
| NICE (2010), UK Clir Gui 110 | inical s uideline s 0 | Social Services | Examples of complex social factors in pregnancy include: poverty; homelessness; substance misuse; recent arrival as a migrant; asylum seeker or refugee status; difficulty speaking or understanding English; age under 20; domestic abuse. Complex social factors may vary, in both type and prevalence, across different local populations' (p10). | The Guideline Development Group using expert professional, local and user knowledge advocated for the inclusion of this criteria which focused on those social factors where there was 'no related impact on the pregnancy or the woman's health' in order that women who have 'social' services input are not excluded from planning to birth in an MLU. As with all the criteria in the GAIN MLU Guideline, if any uncertainty, multidisciplinary discussion is necessary. |

| NI Maternity | Maternity | The aim of the strategy is to 'provide high quality, Objective 1 from NI Maternity Strategy |
|---|--|---|
| Strategy (2012), | Strategy | safe sustainable and appropriate maternity services |
| Northern Ireland | | to ensure the best outcome for women and babies'. |
| | | A partnership approach between health and social |
| | | care staff and members of the public is advocated. |
| | | This is because clinical treatment, emotional care |
| | | and social factors are inextricably linked during a |
| | | woman's pregnancy' (p15). How 'to access care |
| | | and the importance of presenting early in |
| | | pregnancy, regardless of previous clinical history or |
| | | social circumstances' (p41) is highlighted- a |
| | | universal partnership approach is advocated. |
| Devane et al (2010), Ireland Hodnett et al (2010), Canada; Sandall et al, | Systematic review; Cochrane review; Systematic | Women who deliver in a MLU are less likely to have unnecessary intervention, have a spontaneous vaginal delivery and are more likely to breastfeed. |
| (2010), UK | Review Lancet | |
| Renfrew et al, | Ĉochrane | The social and health benefits for women and their |
| (2014) ,UK; | Review; | family are also reported |
| Sandall et al, (2013), UK; Tracy (2005), (2013) NZ | RCT | |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|-------------|--|-----------------------------|
| RCOG (2010), UK | Green Top | Previous | Previous Intrauterine Fetal Death (IUFD) related to a known non | Section 10.2 |
| | Guidelines | Congenital | recurrent cause merits individual assessment for place of birth' | |
| | 55 | Abnormality | | |
| NICE, (2014a), UK | Clinical | | Recommends individual assessment when planning place of birth | Table 9. |
| | Guideline | | for woman with previous stillbirth/neonatal death with a known no- | The GAIN GDG |
| | | | recurrent | considered the evidence |
| | 190 | | a known non-recurrent cause | and carefully |
| | | | | considered the evidence |
| | | | | and carefully worded this |
| | | | | criteria to highlight the |
| | | | | importance of 'no evidence |
| | | | | of reoccurrence' in current |
| | | | | pregnancy when planning |

Table M: Previous congenital abnormality, with no evidence of reoccurrence (FMU/AMU)

Table N: Non-significant (light) meconium in the absence of any other risk (FMU/AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|-------------------------|---|-------------------|
| NICE (2014a), UK | Clinical | Non-significant (light) | Significant meconium- defined as dark green or black amniotic | Section 1.5.2 |
| | Guideline 190 | meconium in the | fluid that is thick or tenacious, or any meconium-stained | |
| | | absence of any other | amniotic fluid containing lumps of meconium' | |
| | | risk | 'If significant meconium is present, transfer the woman to | Section 1.5.4 |
| | | | obstetric-led care provided that it is safe to do so and the birth is | |
| | | | unlikely to occur before transfer is completed' | |

Table O: Uncomplicated third degree tear (AMU)/ Previous extensive vaginal, cervical, or third degree perineal traumafollowing individual assessment (AMU)

| Author/ | Type of | Criteria | Results/Findings | Comment/Reference |
|---|---|--------------------------------|--|---|
| Date/Country | Evidence | | | |
| NICE (2014a), UK | Clinical Guideline 190 | Cervical or perineal trauma | Indicate individual assessment when planning place of birth for woman with previous complication of 'extensive vaginal, cervical, or third- (or fourth) degree perineal trauma' | Table 9 The GAIN GDG considered the evidence |
| Edozien (2014), UK Ali et al (2014), Ireland | Cohort study Retrospective data search | Cervical or perineal trauma | 'The relative risk of a repeat tear is a five-fold increase and the absolute risk of a repeat tear is about 7 in 100'. 'that no studies have generated a model that accurately and definitively predicts recurrence antenatally' (p53). 'Only intervention that significantly impacts on Anal sphincter Injury recurrence is elective prelabour caesarean section but obviously | worded this criteria to highlight the importance of <i>uncomplicated</i> third degree tear for a woman planning to birth in |
| | | | this carries its own risks which must be discussed in the context of the patient's other obstetric issues, co- morbidities and short and long-term obstetric plans (p53).' | inclusion of women with a previous extensive vaginal, cervical, or third degree perineal trauma to plan to birth in AMU only after individual antenatal assessment. |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comments/Reference |
|-------------------------|---------------------|------------|---|---------------------------------------|
| NICE (2014a), | Clinical | Serum | Medical Conditions indicating increased risk suggesting | Table 6. |
| UK | Guideline | antibodies | planned birth at an obstetric unit include atypical | The GDG considered this |
| | 190 | | antibodies which carry a risk of haemolytic diseases of | carefully and noted that women |
| | | | the newborn. | with serum antibodies of no clinical |
| | | | | significance are often excluded |
| | | | | from planning to both in a MLU. |
| | | | | Therefore it was important to |
| | | | | include this as an inclusion criteria |

Table P: Serum antibodies of no clinical significance (FMU/AMU)

Table Q: Women who have had previous cervical treatment, now term (FMU/AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------------|--------------------------------|---|--|
| NICE (2014a), UK | Clinical Guideline 190 | Previous Cervical Treatment | Indicate individual assessment when planning place of birth for woman with 'previous Gynaecological history of Cone biopsy or large loop excision of the transformation zone'. | Table 9. The GDG considered this carefully and were of the view that once term was reached that women with this history may seek to be admitted to a MLU. |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|----------|---|---------------------------|
| NICE (2014a), UK | Clinical | PPH | 'Advise women with increased risk of the previous complication of | Table 7. |
| | Guideline 190 | | primary postpartum haemorrhage requiring additional treatment or | The GDG considered |
| | | | blood transfusion suggest planning to birth in an obstetric unit. | this carefully and agreed |
| | | | | that women with a |
| | | | | previous PPH, not |
| | | | | requiring blood |
| | | | | transfusion or surgical |
| | | | | intervention may plan to |
| | | | | birth in AMU only. |

Table R: Previous PPH, not requiring blood transfusion or surgical intervention (AMU)

Table S: Prostaglandin induction resulting in the onset of labour (AMU)

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|---------------------|---|-----------------------------|
| Thomas et al, | Cochrane | | Prostaglandins increase the chance of cervical change with no | P. 2 |
| (2014), UK | Review | | increase in operative delivery rates'. | The GDG considered the |
| | | Drootoglandin | | evidence and carefully |
| | | | | worded this criteria to |
| | | induction resulting | | include women who have |
| | | In the onset of | | gone into labour following |
| | | labour | | prostaglandin induction and |
| | | | | choose to be admitted to an |
| | | | | AMU. |

| Author/ Date/Country | Type of Evidence | Criteria | Results/Findings | Comment/Reference |
|-------------------------|---------------------|---------------|--|--------------------------|
| NICE (2014c), UK | Clinical | Group B | Pregnant women should not be offered routine antenatal screening | Section 1.8.9 |
| | Guideline 62 | Streptococcus | for group B streptococcus because evidence of its clinical and cost | |
| | | | effectiveness remains uncertain'. | |
| Ohlsson (2014), | Cochrane | | There is lack of evidence from well designed and conducted trials to | P.2 |
| Canada | Systematic | | recommend Intrapartum Antibiotic Prophylaxis (IAP) to reduce | PLEASE NOTE: |
| | Review | | neonatal Early Onset Group B Streptococcus Disease (EOGOSD)'. | Normal Practice in NI is |
| RCOG (2012), UK | Green-Top | • | Antenatal prophylaxis with oral benzylpenicillin for vaginal/rectal | to administer IAP as per |
| | Guideline | | colonisation does not reduce the likelihood of GBS colonisation | recommendations of |
| | | | at the time of delivery and so is not indicated in this situation. | GAIN (2013) |
| | | | IAP should be offered to GBS-colonised women. | |
| | | | IAP should be offered if GBS is detected on a vaginal swab in the | |
| | | | current pregnancy. Vaginal swabs should not be taken during | |
| | | | pregnancy unless there is a clinical indication to do so. If GBS is | |
| | | | present in a vaginal swab, it is likely that the risk of neonatal | |
| | | | disease is increased. A risk of disease of 2.3/1000 may be | |
| | | | assumed (overall UK incidence 0.5/1000; approximately 21% | |
| | | | women are carriers). | |

Table T: Group B Streptococcus positive in this pregnancy with no signs of infection (AMU)

References

- Ali, A Glennon K, Kirkham C, Yousif S, Eogan M (2014) Delivery outcomes and events in subsequent pregnancies after previous anal sphincter injury European Journal of Obstetrics & Gynecology and Reproductive Biology 174 (2014) 51–53 (Anal sphincter injury)
- 2. Arendas K, Qiu Q, Gruslin A. Obesity in pregnancy: preconceptional to postpartum consequences. Journal of Obstetrics & Gynaecology Canada 2008;30:477–88 (BMI).
- Aune D, Didrik Saugstad, O, Henriksen T, Tonstad S. Maternal body mass index and the risk of fetal death, stillbirth, and infant mortality: a systematic review and meta-analysis. JAMA. doi:10.1001/jama.2014.2269 (BMI)
- Confidential Enquiry into Maternal and Child Health (CEMACH). Perinatal Mortality 2007: United Kingdom. CEMACH: London, 2009 [http://www.cmace.org.uk/getattachment/1d2c0ebc-d2aa-4131-98ed-56bf8269e529/PerinatalMortality-2007.aspx] (Age)
- Devane, D; Brennan, M; Begley, C; Clarke, M; Walsh, D; Sandall, J; Ryan, P; Revill, P & Normand, C (2010) A systematic review, meta-analysis, meta-synthesis and economic analysis of midwife-led models of care. Royal College of Midwives: London (Social factors)
- DHSSPSNI (2013) Enhancing Healthcare Services for Children and Young People in Northern Ireland (From Birth to 18 Years) A review of paediatric healthcare services in hospital and the Community DHSSPSNI: Belfast (Age)
- Edozien LC, Gurol-Urganci I, Cromwell DA, Adams EJ, Richmond DH, Mahmood TA, van der Meulen JH. (2014) Impact of third- and fourth-degree perineal tears at first birth on subsequent pregnancy outcomes: a cohort study. BJOG 2014; DOI: 10.1111/1471-0528.12886 (Third & fourth degree perineal tears)
- EI-Chaar, D, Yang, Q, Gao, J. Bottomley, J, Leader A. Wen, Shi Wu, Walker, M (2009) Fertility and Sterility Risk of birth defects increased in pregnancies conceived by assisted human reproduction 92 (5):1557-61 (Assisted Conception)

- Fretts RC. Etiology and prevention of stillbirth. American Journal of Obstetrics & Gynecology 2005;193:1923–35 (Age & BMI)
- Froen JF, Arnestad M, Frey K, Vege A, Saugstad OD, StrayPedersen B. Risk factors for sudden intrauterine unexplained death: epidemiologic characteristics of singleton cases in Oslo, Norway, 1986-1995. Am J Obstet Gynecol 2001;184:694-702 (Age & BMI)
- 11. Guidelines and Audit Implementation Network (GAIN) (2013) Prevention of Early Onset Group B Streptococcal Disease – Northern Ireland Audit Report 2013 FINAL REPORT (Strep B)
- 12. GAIN (2014) Advice for guideline development in Northern Ireland manual GAIN: Belfast
- 13.GAIN (2016) Guideline for Admission to Midwife-Led Units in Northern Ireland and Northern Ireland Normal Labour & Birth Care Pathway. *Guidelines and Audit Implementation Network - GAIN* [Online] Available at: <u>https://rqia.org.uk/getattachment/3a7a37bb-d601-4daf-a902-6b60e5fa58c2/GAIN MLU</u> <u>Guideline.pdf.aspx.</u>
- 14. Healy DL, Breheny S, Halliday J, Jaques A, Rushford D, Garrett C, et al. Prevalence and risk factors for obstetric haemorrhage in 6730 singleton births after assisted reproductive technology in Victoria Australia. Human Reproduction 2010; 25: 265–74 (Assisted Conception)
- Hodnett ED, Downe S, Walsh D, Weston J. A Iternative versus conventional institutional settings for birth. Cochrane Database of Systematic Reviews 2010, Issue 9. Art. No.: CD000012. DOI: 10.1002/14651858.CD000012.pub3 (Social factors)
- 16. Hollowell, JA, Pillas, D, Rowe, R, Linsell, L, Knight, M and Brocklehurst, P. (2013) The impact of maternal obesity on intrapartum outcomes in otherwise low risk women: secondary analysis of the Birthplace national prospective cohort study. *British Journal of Obstetrics and Gynaecology*, September (BMI)
- 17. Huang DY, Usher RH, Kramer MS, Yang H, Morin L, Fretts RC. Determinants of unexplained antepartum fetal deaths. Obstet Gynecol 2000;95:215-21.(Age & BMI)

- 18. HSC PHA (2012) Integrated Perinatal Mental Health Care Pathway Belfast: HSC PHA (Maternal Mental Health)
- 19. Kristensen J, Vestergaard M, Wisborg K, Kesmodel U, Secher NJ. Pre-pregnancy weight and the risk of stillbirth and neonatal death. BJOG 2005;112:403–8 (BMI)
- 20. Lawrence A, Lewis L, Hofmeyr GJ, Styles C. Maternal positions and mobility during first stage labour. *Cochrane Database of Systematic Reviews* 2013, Issue 10. Art. No.: CD003934. DOI: 10.1002/14651858.CD003934.pub4 (Mobility)
- 21. Manktelow BM, Smith LK, Evans TA, Hyman-Taylor P, Kurinczuk JJ, Field DJ, Smith PW, Draper ES, on behalf of the MBRRACE-UK collaboration. Perinatal Mortality Surveillance Report UK Perinatal Deaths for births from January to December 2013. Leicester: The Infant Mortality and Morbidity Group, Department of Health Sciences, University of Leicester. 2015. (Age & Social Deprivation)
- 22. National Institute of Clinical Excellence (NICE) (2010) Public Health Guidance 27, July 2010, Weight management before, during and after pregnancy, London, http://www.nice.org.uk/guidance/PH27 (BMI)
- 23. National Institute of Clinical Excellence (2010), CG 110 Pregnancy and complex social factors: a model for service provision for pregnant women with complex social factors <u>https://www.nice.org.uk/guidance/cg110/chapter/1-Guidance</u> (Social Factors)
- 24. National Institute of Clinical excellence (NICE) (2012) CG 149 Antibiotics for early-onset neonatal infection: antibiotics for the prevention and treatment of early- onset neonatal infection London: NICE <u>https://www.nice.org.uk/guidance/cg149/chapter/research-recommendations</u> (Strep B)
- 25. National Institute of Clinical Excellence (NICE) (2014a) Clinical Guideline 190 Intrapartum care: care of the healthy woman and their babies during childbirth <u>http://www.nice.org.uk/guidance/cg190/resources/guidance-intrapartum-care-care-of-healthy-women-and-their-babies-during-childbirth-pdf</u> (Age, BMI, PPH, Maternal Haemoglobin, Meconium, 4 Previous births, SROM, Threatened Miscarriage, Threatened Preterm labour, Medical Condition, Previous Stillbirth, Cervical or Perineal Trauma, Serum antibodies)

- 26. National Institute of Clinical Excellence (NICE) (2014b) Clinical guideline 192, December 2014 Antenatal and postnatal mental health: clinical management and service guidance London <u>https://www.nice.org.uk/guidance/cg192</u> (Maternal Mental Health)
- 27. National Institute of Clinical Excellence (NICE) (2008, Modified 2014c) Clinical Guideline 62, March 2008, modified 2014, Antenatal Care, London: NICE <u>http://www.nice.org.uk/guidance/cg62/resources/guidance-antenatal-care-pdf</u> (Age, BMI, Social Deprivation & Strep B, Suspected low-lying placenta, Prostaglandin Induction)
- 28. Oakley L, Maconochie N, Doyle P, Dattani N, Moser K. Multivariate analysis of infant death in England and Wales in 2005-06, with focus on socio-economic status and deprivation. Health statistics quarterly / Office for National Statistics 2009; 42:22-39. (Social Deprivation)
- 29. Okun, N and Sierra, S. (2014) Pregnancy outcomes after assisted Human Reproduction Journal of obstetrics & Gynaecology Canada 36(1): 6483 (Assisted Conception)
- 30. Ohlsson A, Shah VS. Intrapartum antibiotics for known maternal Group B streptococcal colonization. Cochrane Database of Systematic Reviews 2014, Issue 6. Art. No.: CD007467. DOI: 10.1002/14651858.CD007467.pub4 (Group B Strep)
- 31. Reddy UM, Wapner RJ, Rebar RW, Tasca RJ. Infertility, assisted reproductive technology, and adverse pregnancy outcomes: executive summary of a National Institute of Child Health and Human Development workshop. Obstetrics and Gynecology 2007;109: 967–77. 70. (Assisted Conception)
- 32. Regulation and Quality Improvement Authority (RQIA) (2012) Baseline Assessment of the Care of Children Under 18 Admitted to Adult Wards in Northern Ireland Belfast: RQIA Accessed on 12 June 2015 <u>https://www.rqia.org.uk/getattachment/04aed398-20e6-430c-8b32-</u> <u>46f8bf52708a/Under-18s-on-Adult-Hosp-Wards-Overview-Report-Web-12-Dec-</u> <u>12 ISBN.pdf.aspx</u> (Age)
- 33. Renfrew, M; McFadden, A; Helena Bastos, M; Campbell, J; Amos Channon, A; Fen Cheung, N; Delage Silva, D.R.A; Downe, S; Powell Kennedy, H; Malata, A; McCormick, F; Wick, L & Declercq, E. (2014) Midwifery and quality care: findings from a new

evidence-informed framework for maternal and newborn care. *The Lancet,* 384, 9948, p1129-1145 (Accessed 17 August 2015) (Social Factors)

- 34. Royal College of Obstetricians and Gynecologists (RCOG) (2010) Late Intrauterine & Fetal Death & Stillbirth (Green-Top Guideline No 55) London: RCOG (Unexplained Stillbirth)
- 35. Royal College of Obstetricians and Gynaecologists (2011). *Management of Women with Mental Health Issues during Pregnancy and the Postnatal Period.* Good Practice No. 14. London: RCOG (Maternal Mental Health)
- 36. Royal College of Obstetricians and Gynecologists (RCOG) 2012 Green –Top Guideline No 26 The Prevention of Early Onset Group B Streptococcal Disease 2nd Edition July London: RCOG (Strep B)
- 37. Royal College of Obstetricians and Gynaecologists (2012) Bacteria Sepsis in Pregnancy Green Top Guideline No 64a London: RCOG (Strep B)
- 38. Royal College of Obstetricians and Gynecologists 2012 'In Vitro Fertilisation: Perinatal Risks and Early Childhood Outcomes' Scientific Impact Paper No. 8 May London: RCOG (IVF)
- 39. Sandall J. (2010) The contribution of continuity of midwifery care to high quality maternity care London: Royal College of Midwives <u>https://www.rcm.org.uk/sites/default/files/Continuity%20of%20Care%20A5%20Web.pdf</u> (Social factors)
- 40. Sandall, J; Soltani, H; Gates, S; Shennan, A; Devane, D; (2013) *Midwife-led continuity* models versus other models of care for childbearing women. CochraneDatabase Systematic Review; 8: CD004667 (Social factors)

- 41. Smith LK, Draper ES, Manktelow BN, Dorling JS, Field DJ. Socioeconomic inequalities in very preterm birth rates. Archives of Disease in Childhood Fetal & Neonatal Edition. 2007; 92(1):F11-F4 (Social Deprivation)
- 42. South Eastern Health & Social Care Trust (2013) Integrated Perinatal Mental Health Care Pathway Belfast: South Eastern Health & Social Care Trust (Maternal Mental Health)
- 43. Thomas J, Fairclough A, Kavanagh J, Kelly AJ. Vaginal prostaglandin (PGE2 and PGF2a) for induction of labour at term
- 44. Cochrane Database of Systematic Reviews 2014, Issue 6. Art. No.: CD003101. DOI: 10.1002/14651858.CD003101.pub3 (Prostaglandin induction)
- 45. Tracy, S.K; Hartz, D; Nicholl, M; McCann, Y; Latta, D (2005) *An integrated service network in maternity - the implementation of a midwifery-led unit* Australian Health Review 2005: 29(3): 332–339 (Social Factors)
- 46.Tracy, S.K; Hartz, D.L; Tracy, M.B; Allen, J.; Forti, A; Hall, B; White, J; Lainchbury, A; Stapleton, H; Beckmann, M; Bisits, A; Homer, C; Foureur, M; Welsh & A; Kildea, S (2013) Caseload midwifery care versus standard maternity care for women of any risk: M@NGO, a randomised controlled trial. The Lancet published online Sept 17 (Social Factors)

http://www.thelancet.com/journals/lancet/article/PIIS0140-6736(13)61406-3/abstract

47. Wiggins DA, Main E. (2005) Outcomes of pregnancies achieved by donor egg in vitro fertilization—A comparison with standard in vitro fertilization pregnancies American Journal of Obstetrics and Gynecology 192 (6) 2002–2006 (Donor Egg & IVF).