



CLINICAL PRACTICE GUIDELINE

Nutrition During Pregnancy

Institute of Obstetricians and Gynaecologists,
Royal College of Physicians of Ireland
and

Directorate of Clinical Strategy and Programmes,
Health Service Executive

(Endorsements pending)

Version 2.0 Date of publication: Nov 2019

Guideline No.27 First Published: July 2013

Revision Date: Nov 2022

Table of Contents

Summary of Recommendations	1
1.0 Key Recommendations	2
2.0 Purpose and Scope	6
3.0 Background and Introduction	6
4.0 Methodology	8
5.0 Guidelines on Gestational Weight Gain	9
5.1 Body Mass Index (BMI)	9
5.2 Gestational Weight Gain (GWG)	10
5.3 Underweight	12
5.4 Multi-fetal Pregnancy	13
5.5 Bariatric Surgery	13
5.6 Lactation and Postpartum	13
6.0 Guidelines on Macronutrients	15
6.1 Energy	15
6.2 Carbohydrate and Fibre	16
6.3 Protein	18
6.4 Fat	20
7.0 Guidelines on Micronutrients	23
7.1 Folate/Folic Acid	23
7.2 Iron	25
7.3 Calcium	28
7.4 Vitamin D	30
7.5 Iodine	32
8.0 Hydration	34
9.0 Guidelines on Food Safety	35
9.1 Toxicological Substances	35
9.2 Food Borne Illness	38
10.0 Specific Diets	44
11.0 Allergies	44
12.0 Hospital Equipment and Facilities	44

CLINICAL PRACTICE GUIDELINE

NUTRITION DURING PREGNANCY

13.0 Provision of Information on Best Practice for Infant Feeding	. 45
14.0 Implementation Strategy	. 45
15.0 Key Performance Indicators	. 46
16.0 Qualifying Statement	. 48
17.0 References	. 49
Appendix 1 Useful websites	65
Appendix 2 Recommended daily macronutrient intakes	66
Appendix 3 Recommended daily micronutrient intakes	67

Summary of Recommendations for Nutrition in Pregnancy

- All women who may possibly become pregnant within the next three months, whether intentionally or not, are advised to take oral FA (FA) 400 micrograms daily to prevent Neural Tube Defects (NTDS); higher does are required preconception & in the first trimester for those with a higher risk of neural tube defects (NTDs) e.g. history of NTDs, obesity or pre-existing diabetes mellitus.
- 2. Pregnant women and women planning pregnancy should be encouraged first and foremost to **eat a healthy, balanced diet** incorporating foods based on the national Food Pyramid sources of iron, calcium, vitamin D and long chain omega-3 polyunsaturated fatty acids are particularly important.
- 3. **Additional vitamin D₃ 10μg is** recommended daily during pregnancy to support intake. A pregnancy multivitamin is a simple way to achieve this.
- 4. **Food safety** is enhanced by good food hygiene and avoiding foods or food supplements which may be teratogenic or harmful to development.
- 5. All women booking for antenatal care should have their **Body Mass Index** calculated accurately.
- 6. Ideally women who are underweight, overweight or obese should be seen by a dietitian for **pre-pregnancy dietary counselling in the community to optimise weight prior to conception** and therefore reduce associated risks during pregnancy.

Separate guidelines are available for obesity & pregnancy, diabetes in pregnancy and for nausea, vomiting and hyperemesis gravidarum.

1.0 Key Guideline Recommendations

1.1 Gestational Weight Gain

- All pregnant women must have an accurate weight, height and body mass index (BMI) measured and recorded at the first antenatal visit. Use BMI to screen women at increased risk for weight-related co-morbidities.
- Pregnancy is a time when women are keen to receive more information about healthy lifestyle interventions and are motivated to make positive choices. Antenatal appointments offer an opportunity to utilise the brief intervention model outlined in the Making Every Contact Count (MECC) Framework.
- Women who are identified as underweight (BMI <18.5 kg/m²) at booking require monitoring for weight gain and referral to a dietitian for assessment.
- Encourage women to achieve a healthy weight before conception to prevent associated complications. Advise postnatal weight loss in overweight and obese women and, where available, refer to a community dietitian.

1.2 Macronutrients

1.2.1 Energy

- Advise women that their dietary energy requirements in the beginning of pregnancy increase only marginally from pre-pregnancy levels. Encourage women to focus on eating well and not only on eating more.
- Energy requirements vary depending on a woman's age, BMI, number of fetuses and activity level. Caloric intake should be individualised based on these factors.

1.2.2 Carbohydrate and fibre

- An RDA of 175g per day is set, based on adequate blood glucose for utilization by the maternal and the fetal brain. This can be met with 3 servings of fruit, 2 servings of vegetables, 3 servings of milk or yoghurt and 3 servings of whole grains.
- Women are encouraged to eat a variety of whole grains, fruit, vegetables, legumes (starchy beans, lentils and pulses), nuts and seeds to meet fibre requirements.

1.2.3 Protein

- Adequate dietary protein intake throughout pregnancy is essential to ensure normal growth and development of the fetus.
- Two servings of protein foods per day in the first and second trimester and three servings in the third trimester will meet these requirements.
- For women who are undernourished or underweight, a balanced protein/energy nutrient supplement may be useful in pregnancy, but only under medical and dietetic supervision.

1.2.4 Fat

- National healthy eating guidelines are applicable to pregnant women in relation to fat intake.
- Pregnant women should be advised to minimise or avoid dietary intake of trans- fatty acids during pregnancy by avoiding fried foods, bakery items and other highly processed foods.

1.2.5 Hydration

- Adequate fluid intake during pregnancy is 2,300 ml/day (EFSA, 2010).
- Requirements increase in warmer ambient temperatures and with physical activity. Estimating requirements becomes vital within clinical settings for women who are unwell and not able to eat or drink liberally.
- Encourage women to drink to maintain pale, straw-coloured to transparent yellow urine with urination frequency of minimum 3-4 times per day

1.3 Micronutrients

1.3.1 Folate/Folic acid

- All women who may possibly become pregnant within the next three months, whether intentionally or not, are advised to take oral FA (FA) 400 micrograms daily to prevent Neural Tube Defects (NTDS)
- Women who intend to become pregnant are advised to start FA at least 6
 weeks before they start trying to conceive so that their folate levels are
 optimised before closure of the neural tube.
- Women who are at increased risk of a pregnancy complicated by a NTD should arrange to see their doctor because they may need a prescriptiononly higher dose of FA 5.0mg daily. Women who are prescribed 5.0 mg before pregnancy should continue on the same dose for the first trimester
- After the first trimester and during breastfeeding, all women are advised to take FA 400 micrograms to meet the World Health Organization's recommended daily intake for pregnancy and breastfeeding.

1.3.2 Iron

- During pregnancy, a supplement containing 16-20 mg of iron in addition to a balanced diet has the potential of reducing incidence of anaemia in the healthy population.
- Women suspected of iron deficiency require a full blood count (FBC) and, if possible, check serum ferritin. The treatment for iron deficiency anaemia is oral supplementation of 100-200 mg per day as elemental iron. Supplements

that contain 305 mg ferrous fumarate or 325 mg ferrous sulphate are equal to 100 mg elemental iron and readily available.

1.3.3 Calcium

- Pregnant women are advised to eat sufficient calcium rich foods to achieve an intake of 1000 mg per day.
- Three servings of milk, yogurt, cheese or fortified plant-based alternatives within a varied diet provide adequate calcium for most women.
- Women who have a low calcium intake are advised to take a daily supplementation to meet requirements.

1.3.4 Vitamin D

 All pregnant women in Ireland should take a supplement containing 10micrograms (400 IU) Vitamin D₃ per day in addition to inclusion of vitamin D rich foods such as fortified milk and oily fish.

1.3.5 Iodine

- Women of childbearing age, and especially those planning a pregnancy, should ensure that they meet the adult requirement of 150 micrograms iodine daily. Iodine requirements increase to 200 micrograms daily during pregnancy and breastfeeding.
- Two or more servings of milk or yogurt and eating white fish once a week is recommended.
- A prenatal multivitamin supplement containing 200 micrograms iodine daily may be required if dietary sources are inadequate.

1.4 Food Safety

• Practicing food safety is important during pregnancy to reduce exposure to toxicological substances and pathogens which may cause harm to the woman and fetus. (See Tables 10 and 11)

1.5 Specific Diets

Refer pregnant women who require therapeutic diets to a dietitian.

1.6 Allergies

- It is not recommended that women avoid potentially allergenic foods during pregnancy unless she herself is allergic to a specific food.
- Women who avoid food groups due to allergy or intolerance are at risk of inadequate intake of nutrients. A referral to a dietitian is warranted in these cases.

1.7 Hospital Equipment and Facilities

- A digital weighing scale and a wall-mounted stadiometer (height measure) should be available. The centre must ensure equipment is available in a discrete area and calibrated as per manufacturer instructions to accurately record measurements.
- Hospitals should have access to phlebotomy and laboratory testing.

1.8 Provision of Information on Best Practice for Infant Feeding

- It is the responsibility of all healthcare professionals to promote best practice for infant feeding.
- The National Infant Feeding Policy for Maternity and Neonatal Services and local implementation policy can be found at: https://www.breastfeeding.ie/Resources/Health-professional/Infant-Feeding-Policy-for-Maternity-Neonatal-Services-2019.pdf

2.0 Purpose and Scope

The purpose of this guideline is to improve healthcare professionals' knowledge and dissemination of optimal advice on nutrition during pregnancy while demonstrating the link between good nutrition during pregnancy and a favourable pregnancy outcome.

These guidelines are intended for healthcare professionals who are working in HSE-funded obstetric and gynaecological services in both community and acute services. They are designed to guide clinical judgment but not to replace it. In individual cases a healthcare professional may, after careful consideration, decide not to follow a guideline if it is deemed to be in the best interest of the woman.

Separate guidelines are available for the management of obesity (HSE 2011 Obesity and Pregnancy Clinical Practice Guideline), diabetes during pregnancy (HSE 2010 Guidelines for the Management of Pre-Gestational Diabetes Mellitus from Pre-Conception to the Postnatal Period) and hyperemesis gravidarum (HSE 2015 Hyperemesis and Nausea/Vomiting in Pregnancy). These guidelines can be accessed on the RCPI website: https://www.rcpi.ie/faculties/obstetricians-and-gynaecology/

3.0 Background and Introduction

Nutrition at time of conception and during pregnancy impacts immediate and long term maternal and infant. Pregnancy is seen as a teachable moment for health promotion, when women are motivated to change. Health care providers can take this unique opportunity to engage and collaborate with women to improve their nutritional intake. The medical team should endeavour to promote healthy lifestyle by "Making Every Contact Count" as part of the Healthy Ireland strategy.

Optimal maternal nutrition will give her baby the best start in life and may subsequently influence health and nutrition of the family unit. As outlined in the National Standards for Safer Better Maternity Services (HIQA, 2016), focus is on supporting women to make positive lifestyle changes while being mindful of social determinants and the ability or limitations to change. The Department of Health is also committed to underpinning health and wellbeing in policy and service delivery to ensure women are empowered to make necessary changes (DOH, 2016). On a global scale, the World Health Organization supports this strategy as they recognise that a woman's positive experience during pregnancy can create the foundations for healthy motherhood (WHO, 2016).

Over 25 years ago, the "Barker Hypothesis" first described the link between fetal nutrition and later development of chronic diseases in adulthood (Barker, 1993; Barker et al, 1995). Poor in utero nutrition can increase adult risk of cardiovascular disease (Kajantie et al, 2005), high blood pressure (Huxley et al, 2000), obesity (Tounian, 2011; Boney, 2005) and metabolic syndrome (Barker et al, 2005). The critical window of opportunity, occurring from conception until 24 months of age, is now commonly referred to as "the first 1000 days". During pregnancy a number of key nutrients are fundamental to the developing fetus. These nutrients and the recommended intakes are outlined in this guideline.

This guideline uses the current evidence to guide health care recommendations. Maternal weight at conception and gestational weight gain also play roles in health outcomes for pregnancy. Babies have increased risk of cardiac death in later life if malnourished in utero (Barker et al, 2012). Maternal diet can influence the type of fetal adipose tissue, which may explain the baby's risk of developing insulin resistance and subsequent diabetes in later life (Symonds et al, 2012). The fuel-mediated in utero hypothesis suggests that increased glucose (carbohydrate) and fat intake during pregnancy results in obesity for the baby in later life (Koletzko et al, 2012). Improved diet and lifestyle regimens potentially reduce this risk (Nelson et al, 2010; Moses et al, 2005.

Obese women are more than twice as likely to give birth to a large-for-gestational age baby compared to women of healthy weight (Sebire et al, 2012), which significantly increases the risk of complications associated with delivery (Boulet, 2003). Women from lower socioeconomic groups are at higher risk of unbalanced nutritional intake and advice must be tailored to their income and needs. Both of these groups of women can benefit from dietetic referral during pregnancy.

In addition to healthy eating advice, women need clear, consistent guidance on the foods to avoid during pregnancy to reduce infection and illness due to exposure of foodborne toxins and pathogens. This guideline provides the latest evidence on the foods to avoid and why they should be avoided. This advice is backed by scientific evidence and any potential risk factors need to be weighed against the benefits when making decisions. For further information and latest reports please refer to the Food Safety Authority Ireland (FSAI) and safefood.

4.0 Methodology

- Medline, EMBASE and Cochrane Database of Systematic Reviews were searched using terms relating to nutrition in pregnancy, fetal programming and nutrition, first 1000 days, diet and pregnancy and specific key nutrients in line with pregnancy.
- Searches were limited to humans and restricted to the titles of English language articles published between 1990-2019
- Relevant meta-analyses, systematic reviews, intervention and observational studies were reviewed.

Guidelines and Evidence Reviewed Included:

- Food Safety Authority of Ireland (FSAI). (2019) Healthy eating, food safety and food legislation: A guide supporting the Healthy Ireland Food Pyramid.
- DoH 2018. Folic Acid Supplementation Policy Report by the Department of Health Folic Acid Policy Committee.
- Safefood, 2017. The Folate Status of Pregnant Women in the Republic of Ireland.
- FSAI, 2016. Update Report on Folic Acid and the Prevention of Birth Defects in Ireland.
- FSAI 2012 Best Practice for Infant Feeding in Ireland: From pre-conception through the first year of infant's life.
- FSAI 2011 Scientific Recommendations for a National Infant Feeding Policy, 2nd Edition.
- HSE 2011. Obesity and Pregnancy Clinical Practice Guideline. Institute of Obstetricians and Gynaecologists, Royal College of Physicians of Ireland and Clinical Strategy and Programmes Directorate, Health Service Executive. April 2011.

The principal guideline developer was Ms. Fiona Dunlevy (CWIUH).

The guideline development team contributors were: Linda Culliney (CUMH), Alexandra Cunningham (Rotunda), Sinéad Curran (NMH), Laura Harrington (NMH), Laura Kelly (Rotunda), Dr Eileen O'Brien (UCD), Louise O'Mahony (CUMH), Dr. Ciara Reynolds (UCD), Dr. Niamh Ryan (CWIUH).

The guideline was peer-reviewed by: Dr Mary Flynn (FSAI), Mary Flynn (CUMH), Cliodhna Foley Nolan (Safefood), Mary Lenahan (FSAI), Joanne O'Halloran (HSE), Ursula O'Dwyer (National Health promotion adviser).

Edited by Prof Michael Turner & Laura Harrington.

5.0 Guidelines on Gestational Weight Gain

Summary Recommendation:

- All women should have their weight measured and Body Mass Index (BMI) calculated and documented at the first antenatal visit, ideally in the first trimester. BMI checks at the booking visit should be used as a screening mechanism for those at increased risk.
- Women who are identified as underweight at booking should be monitored for weight gain and may benefit from a referral to a dietitian. Women with a BMI <17kg/m² should be referred to a dietitian.
- Women should be encouraged to achieve a healthy weight before conception
 to prevent associated complications. Weight loss in the postnatal period
 before subsequent pregnancies is an effective strategy for improved
 outcomes. Postnatal weight loss in overweight and obese women should be
 encouraged and where available onward referral to community dietetics
 should be provided.

The amount of weight gained during pregnancy can impact pregnancy outcomes as well as the future health of a woman and her infant.

Prevalence of overweight and obesity has increased over the past number of decades, including among women of childbearing age. In Ireland, in 2016, the mean BMI of women was 27.2kg/m² (overweight) with 26.6% of these women classified as obese and 1.2% underweight (NCD-RisC, 2017). Less than half of Irish women have a healthy BMI, with one in two Irish women overweight or obese at their antenatal booking appointment (DOH, 2016).

Pre-pregnancy and postnatal weight loss is the most appropriate target for treatment of obesity. Women who restrict their intake during pregnancy, without a dietetic consultation, are at risk of nutrient deficiencies (IOG, 2011). Women who are underweight, overweight or obese should receive dietary counselling in the community to optimise weight prior to conception and therefore reduce associated risks during pregnancy. Obese women should be encouraged to avoid excessive gestational weight gain through healthy diet and exercise (Daly et al 2017).

5.1 Body Mass Index (BMI)

All women must have a weight and height measured and BMI calculated at the first antenatal visit, ideally in the first trimester (IOG, 2011). Measure the woman's height with her shoes off standing straight using a stadiometer (wall-mounted metre stick) to the nearest 0.1 cm. She should be weighed wearing light clothing (to the nearest 0.1 kg), and the BMI calculated and classified. Self-reporting of height and weight is unreliable (Fattah et al, 2010) therefore, it is essential to check maternal weight and height at an antenatal booking visit. BMI classifies weight from underweight to obesity class III. See Table 1 for BMI classifications.

Table 1: BMI classification

BMI (kg/m²)	Classification
<18.5	Underweight
18.5-24.9	Healthy
25.0-29.9	Overweight
30.0-34.9	Obesity Class I
35-39.9	Obesity Class II
>40	Obesity Class III

BMI is a surrogate marker of adiposity but does not measure adipose tissue directly or provide information on fat distribution (Fattah et al, 2010; Prentice and Jebb, 2001). It does however offer an affordable and safe screening tool for risk of complications on both ends of the spectrum. Women with BMIs outside the healthy range have increased risk of pregnancy complication. The highest risk appears to be in women within Obesity Class III (Santos et al, 2019; Lifecycle, 2019). Short stature and low weight have been associated with small-for-gestational age infants (SGA) (Goto, 2019), risk for caesarean section and intra-uterine growth restriction (WHO, 1995), while overweight and obesity increases the risk of complications in delivery (IOG 2011), development of gestational diabetes mellitus (Torloni et al, 2009), gestational hypertensive disorders, preterm birth, LGA infants (Santos et al, 2019) and chronic diseases in the infant (Barker, 2012). Preterm birth risk appears to be higher at both BMI extremes (Santos et al, 2019).

It is important that information about obesity and its risks are communicated in an informative, yet sensitive, manner (Schmied et al, 2010; Furber and McGowan, 2011). Women who are underweight may also be sensitive about their weight and compassion is equally important with this patient group.

5.2 Gestational Weight Gain (GWG)

Women gain weight at different rates during pregnancy. Table 2 shows the physiological contributors to weight gain in a singleton pregnancy (ACOG, 2000). Weight can be affected by nutritional intake, gestation, multiple fetuses, physiological stress, genetic factors and fluid retention and these should be taken into consideration on assessment (WHO, 1995). Women who are overweight or obese at booking tend to gain weight more slowly than women of healthy weight (Farah et al, 2011). Gestational weight loss or poor weight gain may increase the risk of SGA babies and pre-term births (Arendras et al, 2008; Kapadia et al, 2015; Xu et al, 2017; Goldstein et al, 2017), particularly in underweight mothers (Santos et al., 2019) but conversely may be more appropriate for women in the highest categories of obesity (Robillard et al, 2018; PMO Lifecycle, 2019). Obese women with high gestational weight gain appear to have the highest risk for pregnancy complications (Santos et al., 2019).

Referral to a dietitian is required to ensure nutritional adequacy in cases of weight concerns in pregnancy. Clinical judgment should be used to determine other clinically relevant causes for change in weight, for example oedema and fetal growth, before making recommendations to modify the rate of weight gain (Rasmussen and Yaktine, 2009).

Table 2: Distribution of weight gain in pregnancy

	Approximate weight gain (BMI 18.5-24.9 kg/m² at booking)
Baby	3.4 kg
Placenta	0.7 kg
Amniotic fluid	0.9 kg
Mother	
Breasts	0.9 kg
Uterus	0.9 kg
Body fluids	1.8 kg
Blood	1.8 kg
Stores of fat, protein and other nutrients	3.2 kg
TOTAL	13.6 kg

Adapted from ACOG, 2000

There is no international consensus on appropriate weight gain per BMI category in pregnancy (Alavi et al, 2013, NICE 2010). Normal and inappropriate GWG is not well defined, and the relationships with outcomes in pregnancy are possibly an association rather than a causative effect (O'Higgins et al, 2014). The IOM 2009 guidelines have been shown to be appropriate for women of normal weight however, women who are underweight and obese vary (Robillard et al, 2018). Research indicates that obese women have better outcomes if they gain less weight and class III obese women may have better outcomes if they lose weight (Robillard et al, 2018). For overweight or obese women that gain less than the recommended amount but have an appropriately growing fetus, no evidence exists that encouraging increased weight gain to conform with the current IOM guidelines will improve maternal or fetal outcomes (ACOG, 2013). Women who are overweight or obese at booking tend to gain weight more slowly than normal weight women (Farah et al, 2011) and it appears that maternal obesity rather than the rate of GWG is associated with increased risk of pregnancy complications (O'Dwyer et al, 2013).

A healthy diet and/or moderate exercise during pregnancy can reduce the risk of excessive GWG (Daly et al 2017, Muktabhant B et al, 2015; Rogozinska et al, 2017;), risk of caesarean delivery, macrosomia, and neonatal respiratory morbidity, particularly for high-risk women receiving combined diet and exercise interventions (Muktabhant B et al, 2015). Gestational weight gain is positively associated with weight retention in the years post-partum (O'Brien et al., 2019a), and therefore management of weight gain in pregnancy may be considered a public health priority in preventing women becoming overweight or obese between pregnancies.

On average, women who partake in physical activity during pregnancy have been shown to gain 1-2kg less than their inactive counterparts (Choi et al, 2013; Muktabhant et al, 2015). Not only are more physically active women less likely to gain excessive weight during pregnancy they also demonstrate a lower incidence of gestational diabetes and postpartum depression (Physical Activity Guidelines Advisory Committee, 2018). Guidelines suggest women with uncomplicated pregnancies partake in at least 150 minutes per week of moderate-intensity aerobic activity, spread throughout the week (RCPI, 2016; DoHHS, 2008; UK CMO, Recommendations, 2017; ACOG, 2015). Pregnant women who were sedentary before pregnancy should follow a more gradual progression of exercise, with 10 minute bouts of moderate intensity continuous exercise building up to a total of 150 minutes per week (UK CMO, 2017). Women with medical or obstetric complications should be carefully evaluated before making recommendations on physical activity participation during pregnancy (ACOG, 2015).

Gestational weight loss or slow weight gain may increase the risk of SGA babies and pre-term births (Arendras et al, 2008; Kapadia et al, 2015; Xu et al, 2017). In situations where a patient is losing weight in pregnancy dietetic intervention should be sought to ensure nutritional adequacy. Women should be weighed if there is a concern about inadequate weight gain or weight loss. This includes women on a restrictive diet during pregnancy.

Women who quit smoking after conception tend toward higher GWG and may benefit from additional support and targeted intervention (Hulman et al, 2016).

Antenatal appointments offer an opportunity to utilise the brief intervention model, outlined in the Making Every Contact Count (MECC) Framework (HSE, 2016), supporting the implementation of the Healthy Ireland 2013-2025 framework for chronic disease prevention (DOH, Ireland, 2013). Pregnancy is a time when women have regular healthcare appointments, are keen to receive more information about healthy lifestyle interventions, (Ronnberg et al, 2015) and are motivated to make positive choices toward a good pregnancy outcome (Thangaratinam et al, 2012). The frequent antenatal visits may be utilised to provide lifestyle information during pregnancy as qualitative evidence indicates women appreciate any advice that supports a positive pregnancy and healthy baby (WHO, 2016). This type of advice is more acceptable if delivered in an unhurried and supportive way (WHO, 2016).

Women with lower education level are at an increased risk of gaining both excessive and inadequate weight in pregnancy. International evidence suggests that dietary interventions (more so than physical activity interventions) are associated with greater compliance with gestational weight gain guidelines (O'Brien et al., 2019b).

5.3 Underweight

Nutrition education to increase energy and protein intake is recommended for undernourished pregnant women to reduce risk of SGA neonates (WHO, 2016). Women who are underweight pre-pregnancy that gain less than the IOM recommended range may have a higher than normal rate of infant mortality when compared with infants born to women with adequate GWG (Davis et al, 2014). The greater the severity of maternal underweight (<17kg/m²), the higher the risk of pre-term birth (Girsen, 2016; Santos et al, 2019). Women who have a BMI below 17kg/m² at booking may benefit from gaining more weight than recommended in the IOM guidelines, with a 21.6kg weight gain reducing risk for SGA (Robillard et al, 2018). Underweight women should be encouraged to meet requirements during pregnancy and referred to a dietitian to ensure these needs are met.

5.4 Multi-fetal Pregnancy

Limited evidence is available for multi-fetal pregnancies. IOM guidelines suggest that the weight gain of women pregnant with twins, who have good outcomes, varies with pre-pregnancy BMI. The guideline suggests that women with a healthy BMI gain 17-25 kg, overweight women 14-23 kg and obese women 11-19 kg at term (IOM, 2009). Evidence-based weight gain and nutritional guidelines for triplet pregnancies or higher are lacking. For triplet gestations, a healthy pre-pregnancy BMI and a total GWG of at least 15.9 - 20.5 kg have been associated with fewer pregnancy complications (Eddib et al, 2007). A minimum GWG of 16.3 kg by 24 weeks gestation and a cumulative total GWG of at least 23 kg by 33 weeks has been recommended (Dietitians of Canada, 2013; Brown, 2008). For women carrying quadruplets or higher, there are no set guidelines available and clinical judgment should be exercised.

5.5 Bariatric Surgery

The use of bariatric surgery for the treatment of obesity has led to an increase in the number of women who attend antenatal services following a bariatric procedure. After bariatric surgery, women are recommended to wait at least 12-24 months before conceiving as this period tends to be one of rapid weight loss, with higher risk of nutritional deficiencies (Aills et al, 2008; ACOG, 2009; Mechanick et al, 2013) during which an increased incidence of stillbirth has been reported (Gonzalez et al, 2015). Dietary advice and monitoring of food intake by dietitians with special knowledge of bariatric procedures and experience in advising pregnant women is recommended. If possible, appointments should be performed before pregnancy and at least every trimester, and if necessary even at closer intervals (ACOG, 2009; Mechanick et al, 2013). A greater focus on achieving nutritional requirements in all patients regardless of surgery type is required.

Women after bariatric surgery have low levels of micronutrient stores which require monitoring and repletion as needed (Devlieger et al, 2014). Women should be informed about the importance of nutritional supplementation.

No specific GWG guidelines exist for women following bariatric surgery, with recommendations in line with those of the general population (RCOG, 2015; Khan et al, 2013). Obese women post-bariatric surgery may gain less weight than their counterparts and demonstrate higher rates of IUGR, SGA and preterm births (Aills et al, 2008; Guelinckx et al, 2009). Thus, they represent a high risk group and warrant close surveillance before, during and after pregnancy. Intensive multidisciplinary follow-up is recommended (Busetto et al, 2017). Local clinical judgement should be used to determine the most effective care for individual patients based on GWG, tolerance and fetal growth (Jefferys et al 2013).

5.6 Lactation and Postpartum

Postpartum weight retention may contribute to a woman's lifelong development of obesity. Women are at risk for increased BMI between pregnancies (Crosby 2015), with this increase associated with a higher risk for perinatal complications (McBaina et al, 2016). Booking weight is higher in two-thirds of women presenting for antenatal care in subsequent pregnancies (Crosby, 2015). Excess GWG is also a risk factor for post-partum weight retention (Rong et al, 2015).

Breastfeeding may also decrease the likelihood of future obesity in the offspring (Woo & Martin, 2015). Obese women demonstrate lower intention to breastfeed, lower rates of breastfeeding initiation, as well as shorter duration of breastfeeding in the postpartum period (Turcksin et al, 2014). Thus, these women may benefit from extra support and encouragement in this regard during the antenatal and postnatal period.

Weight loss in the postnatal period before subsequent pregnancies is an effective strategy for improved outcomes (Kral et al, 2006). A reduction in BMI is associated with improved perinatal outcomes in subsequent pregnancies (McBaina et al, 2016).

Women from lower socioeconomic backgrounds are at higher risk of developing obesity postnatally and should be targeted for weight loss interventions (Turner and Layte, 2013). The average Irish woman is overweight beginning her pregnancy and postnatal weight retention will likely push many of these women into the obese category. Women referred to community dietetic services for weight management postpartum will be supported in adopting healthy lifestyle behaviours that may also have an impact on the future health of their offspring.

6.0 Clinical Guidelines on Macronutrients

See Appendix 2 for summary of requirements for energy, carbohydrate, protein and fat.

6.1 Energy

Summary Recommendation:

- Advise women that their dietary energy requirements in the beginning of pregnancy increase only marginally from pre-pregnancy levels. Encourage women to focus on eating well and not only on eating more.
- Energy requirements vary depending on a woman's age, BMI, number of fetuses and activity level. Caloric intake should be individualised based on these factors.

Energy requirements increase during pregnancy due to energy deposited in maternal and fetal tissue, increased basal metabolic rate (BMR) and to changes in the energy cost of physical activity. To define the energy cost of pregnancy, desirable GWG must be stipulated (Butte, 2005).

6.1.1 Requirements

In healthy well-nourished women the average increase in BMR over pre-pregnancy values is found to be 5%, 10% and 25% for the first, second and third trimesters, respectively (Butte, 2005). Total energy costs of pregnancy in well-nourished women are thus estimated from the increment in BMR plus the energy deposition associated with a mean GWG of 12kg. The BMR is calculated based on pre-pregnancy weight. Requirements can be calculated by adding activity factor and the appropriate incremental daily energy requirement per trimester. See Table 3.

Table 3: Additional daily energy requirements per trimester

Trimester	EFSA (2013)	WHO (2004)	IOM (2010)
1 st	70kcal	85kcal	NA
2 nd	260kcal	285kcal	340kcal
3 rd	500kcal	475kcal	452kcal

Energy requirements vary depending on a woman's age, BMI and activity level and calorie intake should be individualised based on these factors. Special consideration needs to be made for women who are either underweight or overweight prepregnancy. Women with lower BMI need to gain more weight to produce birth weight comparable to women with normal BMI. Conversely, women with increased BMI need to gain less weight, as reflected in the IOM recommendations for GWG.

There is no ideal prediction equation for energy requirements in pregnancy and any calculation should be given with frequent monitoring of maternal GWG and fetal growth (INDI, 2015). Thus monitoring weight gain versus calorie intake is beneficial.

6.1.2 Multi-fetal Pregnancy

Energy requirements for women carrying multiple foetuses are higher than for singleton. The increase in resting energy expenditure accounts for the increased mass of maternal tissues, including the breast, uterus, body fat, and muscle as well as the increase in blood volume. This can result in a 40% increase in energy requirements for twin gestations (Goodnight et al, 2009).

There are no standardised guidelines for women carrying twins or more, but they have been extrapolated from those for singletons. For women with a normal BMI carrying more than one fetus an estimated 40-45kcal/kg pre-pregnancy weight per day has been recommended, ensuring that adequate weight gain is used as an indicator of sufficient energy intake (Goodnight et al 2009). For underweight women carrying multiple foetuses, the estimate is 42-50kcal/kg pre-pregnancy weight per day, whereas for overweight women, 30-35kal/kg pre-pregnancy weight per day has been suggested.

6.1.3 Lactation

For women exclusively breastfeeding during the first six months after birth, the additional energy requirements during lactation has been estimated at 500kcal/day over pre-pregnancy requirements (EFSA, 2013). Women should again be advised to choose foods with a high nutrient density and quality.

6.2 Carbohydrate and Fibre

Summary Recommendation:

- An RDA of 175g per day is set, based on adequate blood glucose for utilization by the maternal and the fetal brain. This can be met with 3 servings of fruit, 2 servings of vegetables, 3 servings of milk or yoghurt and 3 servings of whole grains.
- Women are encouraged to eat a variety of whole grains, fruit, vegetables, legumes (starchy beans, lentils and pulses), nuts and seeds to meet fibre requirements.

Carbohydrates are important energy sources for every cell and system in the body. During pregnancy there is an increase in demand for carbohydrate in the form of glucose for maternal and fetal brain, central nervous system and red blood cell function, fetal growth and energy stores. The amount of dietary carbohydrate necessary for optimal human health is unknown.

6.2.1 Carbohydrate Requirements

An RDA of 175g per day is set, based on adequate blood glucose for utilization by the maternal and fetal brain, the only truly carbohydrate-dependant organ (IOM 2005).

6.2.2 Sources

Natural carbohydrates in the form of sugars, starches and fibres are found in grains, fruits, vegetables, nuts/seeds, milk and yoghurt. There is no evidence to suggest a certain portion of carbohydrates should come from starches or sugars (IOM, 2005). Given that natural carbohydrates are also valuable sources of vitamins, minerals, fibre, antioxidants and phytonutrients to support health, the National Healthy Eating Guidelines offer advice on a range of intakes to meet dietary requirements for the population. Based on suggested serving sizes from The Food Pyramid, the RDA of 175g of carbohydrate can be met with 3 servings of fruit, 2 servings of vegetables, 3 servings of milk or yoghurt and 3 servings of whole grains (HSE, 2016).

Carbohydrates foods such as crisps, sugar-sweetened breakfast cereals, added sugars, sweets, biscuits, desserts and sugary drinks are of low nutrient value and can contribute to excess weight gain and heart disease. These foods also replace other, more nutritious choices which can lead to insufficient intake of concomitant nutrients. Advice to limit foods high in sugar, fat and salt to no more than two servings per week applies to the whole population and does not change for pregnant women (HSE, 2016). Though serving sizes for these individual foods vary, a serving of added sugar is one teaspoon.

6.2.3 Fibre Requirements

Dietary fibre is non-digestible carbohydrate and lignin, therefore contribute little or no energy. Diets rich in fibre aid in regular bowel function, reduction in blood cholesterol and modulate blood glucose levels (Mousa, 2019). Adequate intake of fibre to support normal laxation for adults is 25g per day (EFSA, 2017). As constipation is common in pregnancy due to hormonal changes to bowel tone, an increase to 28g per day has been recommended (IOM 2005). Women are encouraged to take a variety of whole grains, fruit, vegetables, legumes (starchy beans, lentils and pulses), nuts and seeds daily to meet these requirements (FSAI 2011).

6.2.4 Multi-fetal Pregnancy

There is limited evidence to advise specific carbohydrate and fibre intakes for multifetal pregnancies (Bricker, 2015). Additional energy for fetal growth and maternal wellbeing can be met with an additional daily intake of 1-3 servings from each of these food groups above recommended for a singleton pregnancy: whole grains, fruit, vegetables and milk or yoghurt. Fetal growth, weight gain, maternal wellbeing should be monitored and individual dietary preferences should be taken into account.

6.2.5 Lactation

The RDA for carbohydrate during lactation is based on adequate intake to replace the content in average human milk and for maternal brain utilization (IOM 2005). Estimated requirements during lactation are 210g per day. Assuming the woman has a balanced diet, this increased energy requirement can be also be met concurrently with the recommended increased intake of calcium from milk and yoghurt to 5 servings daily plus an additional serving of fruit. Adequate intake of fibre during lactation is 25 g per day, as for the general adult population.

6.3 Protein

Summary Recommendations:

- Adequate dietary protein intake throughout pregnancy is essential to ensure normal growth and development of the fetus.
- Two servings of protein foods per day in the first and second trimester and three servings in the third trimester will meet these requirements.
- For women who are undernourished or underweight, a balanced protein/energy nutrient supplement may be useful in pregnancy, but only under medical and dietetic supervision.

Adequate dietary protein intake throughout pregnancy is essential to ensure normal growth and development of the fetus as it forms the structural basis for all new cells and tissues. Both excessive and insufficient dietary protein intake have been associated with intrauterine growth restriction and subsequently low birth weight and small for gestational age (Ota *et al.*, 2015).

6.3.1 Requirements

The fetus and placenta consume approximately 1kg of dietary protein throughout pregnancy, primarily in the second and third trimesters. To fulfil this need, women require 0.83g/kg with an additional 1g, 9g and 28g/day in the first, second and third trimesters respectively (EFSA, 2015). See Table 4.

Table 4: Protein requirements per trimester

Trimester	Additional protein	Recommended servings from Food Pyramid
1 st	1 g	2
2 nd	9 g	2
3 rd & lactation	28 g	3

6.3.2 Sources

In order to meet their requirements, pregnant women need to include at least 2 servings per day of a variety of lean protein rich foods in the first and second trimesters and 3 servings in the third trimester (DOH, 2016).

One serving is one of the following:

- 50-75g of cooked lean beef, lamb, pork, mince or poultry
- 100g of cooked fish, soya, tofu or tempeh
- 2 eggs
- 175g of kidney beans/lentils/black beans/dal
- 40g of unsalted nuts or seeds

It is important to consider the source of dietary protein. A high intake of processed meats such as sausages, chicken nuggets and burgers has been associated with a greater risk of small for gestational age infants (Knudsen et al 2008), while a diet containing less processed meats and more fish and eggs may reduce this risk (Ricci et al 2010). Therefore, pregnant women should be advised to limit salted processed meats and choose fresh meat, poultry, fish and plant sources of protein more frequently.

According to Irish food surveys, the majority of women in Ireland meet protein requirements (McGowan and McAuliffe, 2012). Women who have reduced nutritional intake due to nausea or vomiting of pregnancy are at risk of not meeting requirements. Women from lower socioeconomic groups may not meet requirements due to the associated costs of protein-rich foods (McCartney et al., 2013) and more likely to choose less expensive processed foods. This vulnerable group requires advice on lower cost, high quality proteins such as eggs, beans, tinned fish, chicken and lentils.

6.3.3 Multi-fetal Pregnancy

Protein requirements for women with multi-fetal pregnancy are estimated at an additional 50g protein/day in the second and third trimesters. Advise these women to include at least 4 servings per day of lean protein-rich foods (IOM, 2006).

6.3.4 Lactation

Lactating women require on average, an additional 19g of protein per day (EFSA, 2015). This can be achieved by including at least 3 servings per day of lean protein-rich foods.

6.3.5 Protein Supplements

The use of protein powders, beverages or any other high protein supplements is not recommended in pregnancy. Current best evidence states that high protein supplementation alone does not offer clinical benefits, even in undernourished women (Ota, 2012; Kramer & Kakuma, 2003). In women who have adequate protein intake, high protein supplements may increase the risk of small for gestational age (SGA) neonates (Blumfield, 2014; WHO, 2016). However, in women who are undernourished and underweight, a balanced protein/energy nutrient supplement may be useful in pregnancy, but only under medical and dietetic supervision (Imdad, 2011; Kramer & Kakuma, 2003).

6.4 Fat

Summary Recommendations:

- National healthy eating guidelines are applicable to pregnant women in relation to fat intake.
- Pregnant women should be advised to minimise the intake of saturated fatty acids during pregnancy, while ensuring that intake of other nutrients which coincide with saturated fat in the food supply (e.g. iron, vitamin B₁₂, calcium) remain adequate.
- Pregnant women should be advised to minimise or avoid dietary intake of trans- fatty acids during pregnancy.
- Encourage pregnant women to eat additional EPA and DHA by eating 1-2 portions of oily fish per week.

Optimal quality and quantity of dietary fat intake in pregnancy is unclear. National recommendations for fat intake in the general population apply. However, there is evidence which supports the minimisation of saturated and trans-fatty acids, as well as an increased intake of eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA) during pregnancy and lactation.

6.4.1 Saturated Fats

The US Healthy Heart Study (Crume, 2016) revealed that a higher proportional intake of saturated fatty acids, coincident with a lower intake of other macronutrients (including refined sugar), was associated with increased neonatal fat mass. Other studies link excessive saturated fat intake to higher neonatal adiposity, weight-for-age and waist-to-hip circumference ratio at 6 months old and development of obesity, insulin resistance and cardiovascular disease later in life (Murrin, 2013; Horan, 2016; Horan, 2014; Mennitti, 2015). Therefore women should be encouraged to limit fat especially.

The main sources of saturated fat include fatty beef, lamb, pork, poultry eaten with skin, beef fat (tallow), lard, cream, butter, cheese and other dairy products made from whole or reduced-fat (≤ 2 %) milk. Many baked goods and fried foods also contain high levels of saturated fats. It is recommended to advise pregnant women to choose lean, meats and low fat dairy and limit process proteins, baked goods and fried meats, while ensuring adequate intake of other nutrients which coincide with saturated fat in the food supply (e.g. iron, vitamin B_{12} , calcium).

6.4.2 Trans Fats

Maternal trans-fatty acid consumption during critical periods of fetal development may contribute to the development of metabolic diseases in childhood and later life (Innis, 2006; Mennitti, 2015).

Trans-fatty acids are transported across the placenta in proportion to maternal intake. They compete strongly for the same long-chain polyunsaturated fatty acid (LC-PUFA) binding sites on placental and other membranes and may inhibit the transport of LC-PUFAs across the placenta, compromising the mother's synthesis of LC-PUFA for the fetus (SACN, 2004). This may lead to alterations in the development of the hypothalamic centres that control appetite (Murrin, 2013), increasing the risk of metabolic diseases. Pregnant women are advised to minimise or avoid dietary intake of trans-fatty acids during pregnancy.

The main sources of trans-fatty acids include fried foods like doughnuts, and baked goods including cakes, pies, biscuits, pastries and confectionery (FSAI, 2016).

6.4.3 Polyunsaturated Fat (Omega-3 Fatty Acid) Requirements

Long-chain polyunsaturated fatty acids (LC-PUFAs) make up almost half of the lipid content of the brain. The fetus depends on maternal supply of LC-PUFAs due to its limited ability to synthesise them from precursor fatty acids (Herrera, 2002).

DHA and EPA (also known as omega-3 fatty acids) are, two important LC-PUFAs necessary for the normal development of the brain and retina. There is mixed evidence regarding the effectiveness of omega-3 supplementation during pregnancy to improve neurodevelopment in children. However, evidence from a double-blind, randomised controlled trial found that 600mg DHA per day in the second half of pregnancy resulted in longer gestation and larger infant size at birth (Carlson, 2013). In addition, higher DHA and EPA intakes have been associated

with lower neonatal skinfold thickness measures (adiposity), but further research is required in this area (Rytter, 2011).

Table 5: EPA plus DHA requirements (EFSA, 2010)

	Requirements	Recommended servings
Non-pregnancy	1750mg / week EPA + DHA	1-2 servings of oily fish per week
Pregnancy	Additional 700- 1400mg/week DHA	1-2 servings of oily fish per week*

^{*}See section 9.0 food safety recommendations regarding fish and seafood.

DHA and mercury content of fish vary independently: anchovies, Atlantic herring, Atlantic mackerel, mussels, salmon, sardines, snapper and trout are generally high in DHA and low in mercury content.

Note: foods fortified with omega-3 PUFA such as yoghurt, milk and eggs are becoming more widely available, however, many of these fortified foods contain the plant-based omega-3 PUFA (alpha-linolenic acid), which cannot be readily converted to their bioactive derivatives EPA and DHA.

Table 6: The main dietary sources of long-chain polyunsaturated fatty acids

	Omega-6	Sources
Essential fatty acids	Linoleic acid (LA)	Soybean, safflower, sunflower and corn oils; green leafy vegetables; nuts and seeds
LCPUFAs	Arachidonic acid (AA)	Egg yolk, beef, pork, chicken (particularly organ meats)
	Gamma-linoleic acid (GLA)	Evening primrose, blackcurrant oils
	Omega-3	Sources
Essential fatty acids	Alpha-linolenic acid (ALA)	Soybean, rapeseed (canola), flaxseed and walnut oils; nuts and seeds
LCPUFAs	Eicosapentaenoic acid (EPA)	Fish oils, oily fish (anchovies, Atlantic herring, Atlantic mackerel, mussels, farmed and wild salmon, sardines,
	Docosahexaenoic acid (DHA) / Docosapentaenoic acid (DPA)	snapper and trout)

6.4.4 Monounsaturated Fats

There is insufficient evidence at this time to make pregnancy specific recommendations for the intake of monounsaturated fatty acids. Therefore, pregnant women should be advised to follow existing dietary guidelines for the general population.

7.0 Clinical Guidelines on Micronutrients

See Appendix 3 for summary of recommendations for micronutrient intake in pregnancy and lactation.

7.1 Folate/ folic acid

Summary Recommendations:

- All women who may possibly become pregnant within the next three months, whether intentionally or not, are advised to take oral FA (FA) 400 micrograms daily to prevent Neural Tube Defects (NTDS)
- Women who intend to become pregnant are advised to start FA at least 6
 weeks before they start trying to conceive so that their folate levels are
 optimised before closure of the neural tube.
- Women who are at increased risk of a pregnancy complicated by a NTD should arrange to see their doctor because they may need a prescription-only higher dose of FA 5.0mg daily. Women who are prescribed 5.0 mg before pregnancy should continue on the same dose for the first trimester
- After the first trimester and during breastfeeding, all women are advised to take FA 400 micrograms to meet the World Health Organization's recommended daily intake for pregnancy and breastfeeding. This promotes fetal and neonatal development as well as reducing the risk of anaemia in the mother
- Women who are considered at increased risk include women who:
 - (a) experienced a previous pregnancy complicated by a NTD
 - (b) have pregestational Type 1 or 2 diabetes mellitus
 - (c) have a first degree relative diagnosed with a NTD
 - (d) are on certain medications (check medications on latest *Monthly Index of Medical Specialities Ireland MIMS*)
 - (e) with moderate or severe obesity (BMI > 34.9 kg/m^2).

All women should follow the National Guidelines for Healthy Eating but they should be aware that increasing their dietary intake of folate alone is unlikely in the absence of mandatory food fortification or FA supplementation to achieve optimal maternal folate levels.

(See Department of Health Folic Acid Policy, 2018)

Neural Tube Defects

Neural tube defects (NTDs) are the most common major malformation of the central nervous system in the developing fetus and include anencephaly, encephalocele, hydranencephaly, iniencephaly and spina bifida, as well as rarer forms. These defects arise at an early stage of pregnancy, between 21 and 28 days after conception, a time when many women do not yet realise they are pregnant as they have not yet experienced a missed menstrual period. The incidence of NTDs in Ireland remains high by European standards (EUROCAT, 2010; EUROCAT, 2014; McDonnell, 2014; McDonnell, 2018). The majority of young women in Ireland are not adequately protected against NTDs due to both low folate intakes from food and lack/insufficient use of pre-conceptual supplements (Safefood, 2017). Only 2.6% of women in Ireland presenting for antenatal care are achieving adequate dietary folate (Safefood, 2017).

A number of recent reviews reinforce the role of folate in the prevention of NTDs and in supporting the development of the fetus in utero, as well as in supporting maternal health during pregnancy and lactation (Molloy, 2008; Burke, 2009; Boilson, 2012; Lassi, 2013; WHO, 2015; McDonnell, 2018). In addition, the importance of sufficient folate both before and during pregnancy in preventing maternal anaemia is becoming increasingly apparent (O'Malley, 2018) with coexistent iron deficiency an additional consideration (Molloy, 2014).

7.1.2 Sources

Dietary sources include:

- naturally occurring folates in fruit, vegetables, beans, pulses and
- folic acid, the synthetic form used in food fortification of some breakfast cereals and milk.

Promotion of a healthy diet pre-pregnancy and during early pregnancy is important but dietary sources alone are insufficient to confer protection (Molloy, 2008; Obeid, 2014; Hopkins 2015; Kelly, 2015). Folic acid fortification is not mandatory in Ireland and foliates present in fruit and vegetables are reduced by heat, light and bruising and several factors impact on individual absorption rates.

A 'whole diet' approach is advocated to optimise bioavailability, with an emphasis on following the healthy eating guidelines in addition to daily FA supplementation to reduce the risk of NTDs.

7.1.3 Supplementation

Folic acid is the synthetic, stable version of folate with a predicable dose response effect on red blood cell folate. FA is used in food fortification and in oral supplements. Supplementation with FA has been demonstrated to protect against NTDs for 70% of the target population. All women who could conceive are encouraged to achieve adequate red cell folate by taking a 400 microgram supplement of folic acid daily in addition to dietary sources, for at least 12 weeks prior to trying to conceive. Supplemental FA can be continued throughout pregnancy and breastfeeding either as a single supplement of 400 micrograms/ day or as part of a pregnancy multivitamin if preferred.

Women who are at increased risk are advised to take a higher dose of folic acid to mitigate an increased likelihood of an NTD in their developing fetus. Women who are considered at increased risk include women who: experienced a previous pregnancy complicated by a NTD, have pregestational Type 1 or 2 diabetes mellitus, have a first degree relative diagnosed with a NTD, are on certain medications (check medications on latest *Monthly Index of Medical Specialities Ireland MIMS*), with moderate or severe obesity (BMI > 34.9 kg/m²).

The higher dose most widely available on the Irish market is 5mg and needs to be prescribed. This dose should be continued throughout the first trimester and 400 microgram/ day recommended as a single supplement or within a prenatal multivitamin for the duration of pregnancy and breastfeeding (FSAI, 2016; Safefood 2017).

7.2 Iron

Summary Recommendations:

- During pregnancy, a supplement containing 16-20 mg of iron in addition to a balanced diet has the potential of reducing incidence of anaemia in the healthy population.
- Women suspected of iron deficiency require a full blood count (FBC) and, if
 possible, check serum ferritin. The treatment for iron deficiency anaemia is
 oral supplementation of 100-200 mg per day as elemental iron. Supplements
 that contain 305 mg ferrous fumarate or 325 mg ferrous sulphate are equal
 to 100 mg elemental iron and readily available.

Iron is an essential component of hundreds of proteins and enzymes. As a component of haem it is responsible for oxygen transport, energy production and DNA synthesis (SACN, 2010).

7.2.1 Requirements

An iron intake of 16mg per day is essential throughout pregnancy to meet the increased requirement for fetal and placental growth, while maintaining adequate maternal stores to cover losses from childbirth. Several portions of iron rich foods daily are encouraged (table 7 & table 8).

During pregnancy, the fetus requires a large red blood cell mass to provide sufficient oxygen for development and growth. Iron requirements increase progressively after 25 weeks gestation to combat the lower oxygen environment in the womb (Dewey and Chaparro, 2007). Adaptations in iron transporters and maternal intestinal absorption facilitate delivery of iron to the fetus to meet these requirements (SACN 2010). Due to these maternal adaptations, along with cessation of menstrual losses, iron requirements are estimated to be the same as premenopausal women. Requirements for pregnancy are 16 mg per day (EFSA 2015).

7.2.2 Sources

Iron in the diet exists as haem iron and non-haem iron. Haem iron is readily absorbed and found nearly exclusively in animal tissues. Non-haem iron, sourced from animal and plant tissues, is absorbed less efficiently. Tables 7 and 8 list sources and iron content of common foods.

Table 7: Food sources and content of haem iron

Haem Iron Sources			
Food	Iron mg Per 100g	Recommended Serving size (g)	Iron content (mg)
Lean Beef	2.6	75	1.9
Beef Mince	2.3	75	1.7
Lamb cutlets	2.0	75	1.5
Pork Chops	1.3	75	1.0
Chicken Breast	1.0	75	0.8
Salmon	0.8	100	0.8
Tinned salmon	0.6	100	0.6
Tinned sardines	2.3	100	2.3
Cod	0.4	100	0.4

Table 8: Food sources and content of non haem iron

Non-haem Iron Sources		
Food	Portion (g)	Iron content (mg)
Fortified breakfast cereal	30	2.4-4.2
Baked Beans	150	2.1

Spinach	90	1.8
Egg (2 whole)	50	1
Wholemeal bread, slice	36	0.9
Broccoli	85	0.9
Dried Fruit	25	0.6

Absorption of non-haem iron can be enhanced with concomitant intake of vitamin C from fruit, fruit juice and certain vegetables. Tannins found in tea and coffee should be avoided at meal times as they can inhibit iron absorption from the diet (FSAI, 2011). Dietary sources of zinc, vitamin C, vitamin A, folate and copper are also important for haemoglobin synthesis. A food-based approach is best to meet requirements for all these nutrients.

7.2.3 Supplementation

Preventing anaemia

National nutrition surveys and research in maternity services show that the majority of Irish women do not meet their daily iron requirements during childbearing years or pregnancy (IUNA, 2011; SLÁN, 2007; McGowan and McAuliffe, 2012). An iron-rich diet in addition to appropriate supplementation has the potential of reducing incidence of anaemia in pregnancy and subsequent adverse outcomes, therefore the threshold for iron supplementation in pregnancy should be low (Barroso et al, 2011).

During pregnancy, a supplement containing 16-20 mg of iron in addition to a balanced diet is effective to maintain adequate stores in the healthy population (Health Canada, 2009) to reduce risk of anaemia (Pena-Rosas, 2015). Intermittent oral iron (120 mg with 2800 micrograms FA) once a week or every second day is recommended when daily iron supplementation is not tolerated due to side effects (WHO, 2016). As haemoglobin (Hb) levels >13g/dL have also been associated with poor obstetric outcomes, advice on iron supplementation should be individualised (SACN, 2010). Women with Hb >13~g/dL are advised to avoid supplementation.

Treating anaemia

A recent analysis of 312,281 pregnancies in 29 countries showed the odds of maternal death were twice as high in women with severe anaemia compared with those without (Daru et al 2018).

Symptoms of iron deficiency include shortness of breath during normal activity, fatigue and palpitations. Women suspected of iron deficiency should have a full blood count (FBC) and, if possible, serum ferritin analysed. Haemoglobin levels <11g/dL in the first trimester and <10.5g/dL in the second/third trimesters indicate

iron-deficiency anaemia (WHO, 2016). Serum ferritin <15 micrograms/L indicates iron depletion in all stages of pregnancy (Pavord et al, 2012). The treatment for iron deficiency anaemia is oral iron supplementation (NICE, 2008). Doses 100-200mg per day of elemental iron should be adequate to replenish stores (Pavord et al, 2012). Supplements that contain 305 mg ferrous fumarate or 325 mg ferrous sulphate are equal to 100 mg elemental iron and readily available. If a woman is diagnosed with anaemia, 100-120 mg elemental iron should be increased until her Hb level rises to > 11.0g/dl after which she can return to a standard dose (WHO, 2016).

Iron supplements can cause unpleasant gastrointestinal (GI) side effects resulting in poor compliance, impair absorption of other minerals, and increase risk of haemoconcentration (Zhou et al, 2009). Intermittent dosing (every second or third day) or lower dose iron preparations seem to be effective in preventing or treating anaemia without unpleasant GI symptoms (Pena-Rosas, 2015; SACN 2010). Note: Despite its high iron content, liver should be avoided in pregnancy due to the high level of pre-formed vitamin A (FSAI, 2011).

7.2.4 Multi-fetal Pregnancy

The iron requirement for twin pregnancy is estimated to be nearly twofold that of a singleton. Iron supplementation during the first and second trimesters has been associated with reductions in both preterm birth and LBW (Goodnight, 2009).

7.2.5 Lactation

Women should maintain an iron-rich diet postpartum to assist with wound healing and replenish stores. Women with anaemia postpartum are at increased risk and severity of infection (Ekiz et al, 2005), reduced work capacity and performance (Haas et al, 2001) and disturbances in cognition and emotions (Beard et al, 2005). There is limited evidence on the optimal treatment of postpartum anaemia (Markova et al, 2015). Iron supplementation in combination with monitoring, advice and treatment of GI symptoms is recommended.

7.3 Calcium

Summary Recommendations:

- Pregnant women are advised to eat sufficient calcium rich foods to achieve an intake of 1000 mg per day.
- Three servings of milk, yogurt, cheese or fortified plant-based alternatives within a varied diet provide adequate calcium for most women.
- Women who have a low calcium intake are advised to take a daily supplementation to meet requirements.

Calcium is an integral part of bone structure, giving the properties of rigidity, strength and elasticity. Ninety-nine percent of corporal calcium is present in bone and teeth. The remaining 1% acts an intracellular messenger in cells and tissues,

regulating metabolic processes, vascular contraction and dilation, muscle contraction, neural transmission and glandular secretion.

An inverse relationship has been described between calcium intake and hypertensive disorders of pregnancy from epidemiological and clinical studies since the 1980's (Belizan and Villar, 1980). It is suggested that low dietary calcium intake stimulates the release of parathyroid hormone or renin, resulting in increased intracellular calcium in vascular smooth muscle which causes vasoconstriction (Hofmeyr et al., 2014). The current consensus from the 2014 Cochrane review is that calcium supplementation (\geq 1000 mg/day) significantly reduces the risk of pre-eclampsia and preterm birth among women with low calcium diets (Hofmeyr et al., 2014).

Accretion of 30, 60 and 300 mg/day of calcium takes place in trimester one, two and three, respectively (Givens and Macy, 1933; Prentice, 2000; Institute of Medicine, 2011; Kovacs, 2016). Physiological adaptations occur during pregnancy to meet foetal calcium requirements, facilitated through doubling of maternal intestinal calcium absorption and mobilisation of calcium from maternal bones (Kovacs and Ralston, 2015). The maternal skeletal calcium can be resorbed if calcium is required for the fetus (Kovacs, 2016). The mechanisms involved in bone metabolism in pregnancy are not fully understood. An imbalance in bone metabolism with resorption exceeding formation may result in negative bone remodelling in the pregnant woman and could characterise pregnancy as a period of vulnerability for maternal skeletal health (Sanz-Salvador et al., 2015). Preliminary Irish evidence suggests that maternal low calcium intake in trimester three is associated with lower maternal bone mineral density five years after pregnancy (O'Brien et al., 2018).

7.3.1 Requirements

The European Food Safety Authority established a population reference intake (PRI) of 1,000 mg/day for adults 18 – 24 years and 950 mg/day for adults ≥25 years (EFSA, 2015). The PRI is similar to the RDA in that it meets the needs of virtually all (97.5%) members of the population. Considering the increases in intestinal calcium absorption during pregnancy, it is recommended that non-pregnant requirements be applied to pregnant women (IOM, 2011; EFSA, 2010; EFSA, 2015). In Ireland, calcium intakes are generally good among pregnant women with estimated intakes of 877 – 919 mg/day (McGowan and McAuliffe, 2012; Mullaney et al., 2016). A study of Irish women, which included contribution from calcium supplements, found higher intakes of 1,227 mg/day (O' Callaghan et al., 2016).

7.3.2 Sources

Calcium can be found in foods naturally and due to fortification. Three servings of calcium rich foods in addition to minor contributors (green leafy vegetables, almonds, beans, seeds) should be adequate for most women. Adolescents will require five servings to meet requirements for maternal and fetal growth. See sources in Table 9.

Calcium supplements may be needed for some women with low dietary intake. Absorption is greatest when supplements ≤500mg are consumed. If a woman is advised to take 1,000 mg/day of calcium from supplements, it would be preferable to split the dose and take 500 mg at two separate times during the day.

Table 9: Calcium-containing foods

Food source	Portion size	Calcium content
Cow's milk	200 ml	240 mg
Cheese	28 g	240 mg
Yoghurt	125 g	240 mg
Alternative milks, fortified	200 ml	240 mg
Rice pudding	200g (half large tin)	175 mg
Tinned sardines (eaten with soft bones)	60 g (half tin)	260 mg
Soya bean curd/tofu (Set with calcium chloride (E509) or calcium sulphate (E516), not nigari)	60 g	200 mg
Calcium fortified instant hot oat cereal	15g dry cereal (1 tbsp)	200 mg
Calcium-fortified orange juice	150 ml	180 mg
White bread	2 large slices	100 mg
Boiled broccoli	2 spears (85 g)	34 mg

7.3.3 Multi-fetal pregnancy

There is limited evidence regarding calcium requirements in pregnancies of twins or higher multiples. It has been suggested that bone resorption is greater in twin compared to singleton pregnancies (Nakayama et al., 2011), and perhaps need greater calcium intake. Recommending up to 5 servings of calcium rich foods daily is pragmatic and will increase women's intake of other essential micronutrients such as iodine.

7.3.4 Lactation

Calcium requirements during breastfeeding are similar to those in pregnancy. It has been recognised that during lactation, maternal bone resorption is transient, and calcium intakes greater than those recommended in pregnancy do not supress bone resorption or alter the calcium content of breast milk (Institute of Medicine, 2011).

7.4 Vitamin D

Summary Recommendations:

 All pregnant women in Ireland should take a supplement containing 10micrograms (400 IU) Vitamin D₃ per day in addition to inclusion of vitamin D rich foods such as fortified milk and oily fish.

Vitamin D is a fat-soluble vitamin essential for absorption of dietary calcium and may play a role in normal immune system function (Fronczak, 2003; Hypponen, 2001). Adverse effects of low serum 25(OH)D concentration on maternal reproductive health may include pre-eclampsia/PIH, increased risk of operative delivery, intrahepatic cholestasis of pregnancy and periodontal disease in pregnancy (SACN, 2016). There is also some evidence to suggest that prenatal supplemental vitamin D may reduce risk of adverse pregnancy outcomes including neonatal hypocalcaemia, rickets and osteomalacia and may reduce the risk of persistent/recurrent wheeze or asthma in children. (EFSA, 2018; Bärebring, 2018; Munns, 2016; Roth, 2017). Surveys of the Irish population show that the average intake of vitamin D is well below the recommended level (SLAN, 2007; IUNA, 2011). It is of particular concern that suboptimal intakes and low serum vitamin D levels have been reported in cohorts of pregnant women in Dublin, Cork and Belfast (McGowan, 2011; Mullaney, 2016; O'Riordan, 2008; Holmes, 2009). Observational studies of pregnant women in Ireland have also revealed significant levels of vitamin D deficiency and insufficiency (O'Brien, 2017; Kiely, 2016; Walsh, 2013).

7.4.1 Requirements

The Food Safety Authority set the RDA (recommended dietary allowance) for vitamin D at 5 micrograms (200 IU) per day for pregnant women (FSAI 1999). More recent guidelines from The European Food Safety Authority (EFSA), American College of Obstetrics and Gynaecologist and The Endocrine Society Global Consensus recommend an intake of 15 micrograms (600 IU) daily from all sources (EFSA 2018; Munns, 2016; ACOG, 2011) whereas The UK recommends that pregnant women take a vitamin D supplement of 10 micrograms daily (NICE 2008).

In the absence of updated Irish RDAs and given the known inadequacy of dietary intakes in Ireland, it is advisable for pregnant women in Ireland to take a daily supplement containing 10 micrograms (400 IU) vitamin D_3 . The majority of overthe-counter prenatal multivitamins contain 10 micrograms of vitamin D_3 , therefore if a woman chooses to take a prenatal multivitamin she will not require additional vitamin D_3 supplementation. Women who choose a strict vegetarian or vegan diet are advised to take a supplement in form of ergocalciferol (vitamin D_2).

If there is a history of rickets in a sibling or a known maternal vitamin D deficiency, a higher, treatment dose is warranted as the neonatal serum vitamin D will be 60% of the maternal level. The number of trials and outcomes reported are too limited, and in general too low in quality, to draw conclusions on the usefulness and safety

of high doses of vitamin D supplementation as a part of routine antenatal care. The EFSA established the upper limit of 100 micrograms/day (4000 IU) which applies to all adults, including pregnant and lactating women (EFSA, 2012).

7.4.2 Sources

Vitamin D_3 is found naturally in few foods. Dietary sources include the flesh of oilrich fish, some fish liver oils (fish liver oil should be avoided in pregnancy due to vitamin A content), and eggs from hens fed vitamin D. Fortified sources including margarine, milk and cereals are also widely available. Vegan-friendly vitamin D_2 is predominately sourced as supplements. Humans can synthesize vitamin D cutaneously; however, above latitudes of approximately 40°N such as in Ireland from October to March, vitamin D cannot be synthesized. The UV light required to promote vitamin D synthesis cannot penetrate the atmosphere during this time. Furthermore, sun exposure may increase the risk of melanoma, and so advising sun exposure is not a viable public health strategy to combat low vitamin D levels.

7.4.3 Multi-fetal Pregnancy

There is no guideline for vitamin D intake with twins or higher multiples. Considering the frequent occurrence of complications resulting in bed rest for women carrying multiples, and therefore limited cutaneous vitamin D synthesis, supplementation may be particularly important.

7.4.4 Lactation

Recommendations for vitamin D do not differ for lactating women. Again, as dietary sources are not consistently adequate, supplementation of 10 micrograms per day is prudent (EFSA, 2012). Vitamin D content of human milk is not sufficient to maintain stores in breastfed infants, therefore the HSE recommends that infants be given vitamin D drops orally (5 mcg vitamin D daily) until 12 month of age and when dietary sources are adequate (HSE, 2010; FSAI, 2011).

7.5 Iodine

Summary Recommendations:

- Women of childbearing age, and especially those planning a pregnancy, should ensure that they meet the adult requirement of 150 micrograms iodine daily. Iodine requirements increase to 200 micrograms daily during pregnancy and breastfeeding.
- Two or more servings of milk or yogurt and eating white fish once or twice a week or is recommended.
- A prenatal multivitamin supplement containing 200 micrograms iodine daily may be required if dietary sources are inadequate.

Iodine is an essential nutrient required for the biosynthesis of thyroid hormones, which are responsible for regulating growth, development and metabolism. Iodine requirements increase substantially during pregnancy and breastfeeding. Thyroid function is increased during pregnancy as thyroid hormones produced by the mother (and the fetus as the pregnancy progresses) are essential for growth of the fetus and to regulate the development of the fetal brain and nervous system. If women have too little iodine during pregnancy or infants have too little during early childhood, the damage may be irreversible. Research has shown that severe iodine deficiency can stunt children's normal physical growth and mental development, resulting in lower intelligence quotients (Harding 2017). A review of the impact of iodine status on the cognitive development of children aged 5 years and under indicated lower I.Q scores in iodine deficient children compared with iodine-replete children and concluded that iodine deficiency had a substantial impact on mental development (Bougma et al 2013). Less is known about the consequences of mild or moderate deficiency.

Research in the UK and Ireland has shown mild iodine deficiency in schoolgirls and pregnant women. (Vanderpump 2011) It is likely that many adult women may not be getting enough iodine, particularly in pregnancy. While there is as yet no available evidence of widespread thyroid hypofunction in the Irish obstetric population, one small Irish study observed levels suggestive of iodine deficiency in more than half (55%) of pregnant women, with a defined seasonal variation (Nawoor et al 2012).

7.5.1 Requirements

To ensure healthy thyroid function throughout pregnancy, adequate iodine intake is needed in the months prior to conception to ensure good thyroid stores. Following the National Healthy Eating Guidelines would be sufficient for most women to meet their iodine requirement preconception and during pregnancy. Including 2-3 servings of milk or yogurt daily and eating white fish once a week is recommended. Recent reviews have found insufficient data to reach any meaningful conclusions on the benefits and harms of routine iodine supplementation in women before, during or after pregnancy (Harding et al 2017). However, supplementation is advised for women who do not consume dairy foods or fish. As too much iodine has potential

to cause harm, only a multivitamin supplement designed for pregnancy containing 200 micrograms iodine is recommended.

7.5.2 Sources

Iodine is found in a range of foods, the richest sources being fish and dairy products. While seaweed is a concentrated source of iodine, it can provide excessive amounts and should be eaten in moderation.

The actual amount of iodine in food varies according to the iodine content of the soil, farming practice, fish species and season, making it difficult to accurately gauge iodine content per portion. In Ireland and the UK, milk and dairy products are the main sources of iodine for most people. Research in the UK has shown that organic milk has a 35-40% lower iodine content than conventional milk.

Pregnancy multivitamins currently available on the Irish market contain supplemental iodine between 100 and 200 micrograms, with the most widely available brands meeting 100% of requirements for pregnancy and lactation.

7.5.3 Multi-fetal Pregnancy

There is no specific recommended level of iodine for multiple pregnancy. Diets which include intakes of dairy foods and white fish sufficient to meet calcium requirements will also provide iodine. Where women follow restrictive diets for whatever reason, a pregnancy multivitamin daily is a practical recommendation.

7.5.4 Lactation

Iodine is a normal mineral in breastmilk that is essential for the infant's thyroid. The amounts of iodine in breastmilk vary with maternal iodine intake. Infant requirements are estimated to be 15 micrograms/kg daily in full-term infants and 30 micrograms/kg daily in premature infants (Ares, 2008). European research indicates that in iodine-sufficient areas, unsupplemented mothers excrete sufficient amounts of iodide in breastmilk, and routine supplementation is not necessary. (Gonzalez- Iglesesas 2012)

8.0 Clinical Guidelines on Hydration

Water is essential for all bodily functions, in particular thermoregulation. A water intake which balances losses and thereby assures adequate hydration of body tissues is essential for health and life (EFSA, 2010). Dehydration can result in constipation, headaches, lethargy and increased frequency of urinary tract infections. Fluid balance is generally self-regulated and managed without conscious involvement. Estimating requirements becomes vital within clinical settings for women who are unwell and not able to eat or drink liberally (Vivanti, 2012).

Poor hydration is a concern particularly among women experiencing hyperemesis gravidarum, a severe form of nausea and vomiting in pregnancy. Additional healthcare professional support, including referral to a dietitian, should be given to a woman with hyperemesis gravidarum to ensure her hydration status is maintained throughout pregnancy. HSE/RCPI guidelines for the management of hyperemesis gravidarum are available.

8.1 Requirements

Water requirements change depending on individual and environmental factors. Historically, it has been estimated in various ways from mL/kcal to mL/kg (Vivanti, 2012). Under ordinary circumstances, a reasonable estimate can be derived from 1 ml of water required for each 1 kcal of energy consumed (NRC, 1989). The EFSA suggest adequate intake for pregnant women to be that of non-pregnant women (2000 ml/day), plus an increase of 300 ml/day based on increased energy requirements. Therefore the recommended adequate fluid intake during pregnancy is 2,300 ml/day (EFSA, 2010).

Requirements increase in warmer ambient temperatures and with physical activity. Thirst may not be a reliable indicator of adequate hydration as mild dehydration triggers the sensation of thirst. A woman can be advised to monitor the colour of her urine as a practical indicator of hydration. Encourage women to drink to maintain pale, straw-coloured to transparent yellow urine with urination frequency of minimum 3-4 times per day (McKenzie, 2017).

8.2 Sources

Fluid requirements can be met with water from food and fluids, as well as oxidative processes in the body. Water is recommended as the main source of fluid for the general population as it does not contribute to dental caries or excessive weight gain. This advice does not change for pregnant women (FSAI, 2019).

8.3 Lactation

Estimates for adequate water intakes for lactating women are based on recommended intakes for women and additional water losses in breast milk production. This equates to 600-700 mL/day above the adequate intake for non-lactating women of the same age (EFSA, 2010).

9.0 Clinical Guidelines on Food Safety

Practicing food safety is important during pregnancy to reduce exposure to toxicological substances and pathogens which may cause harm to the woman and fetus. Table 10 provides a summary of toxicological agents, pathogens and sources women must be advised to avoid during pregnancy and lactation.

9.1 Toxicological substances

Summary recommendations:

Table 10: Toxicological substances and precautions in pregnancy and lactation

Substance	Reason for avoidance	Recommended upper limit (UL) and precautions	Sources
Caffiene	Stimulant	≤200mg Maximum 2 – 4 cups tea, 2 cups instant coffee, 1 filtered coffee a day	Coffee, tea, fizzy drinks, chocolate and medications
Alcohol	Teratogenic	Unknown – Avoid in pregnancy	Wine, beer, spirits, some medications and herbal tinctures
Mercury	Toxin	90-100 µg MeHg (2 cans Tuna or 1 tuna steak avoid other large fish)	Large fish high in food chain Shark, swordfish, marlin
Vitamin A (retinol equivalent)	Teratogenic	≤3000 µg	Liver, paté, some Supplements
Biotoxins	Neurotoxins	Not destroyed with heating or cooking Total avoidance recommended in pregnancy	Clams, mussels, scallops
Toxoplasma gondii	Parasite	Hand washing and good hygiene; Wear gloves when gardening or changing cat litter and wash hands afterward	Cat faeces/litter

9.1.1 Caffeine

There is sufficient evidence to indicate a risk to the health of the unborn with excess caffeine in pregnancy (WHO, 2016). Caffeine is a mildly addictive stimulant and is found in coffee, tea (including green and white tea), cocoa, chocolate, soft drinks, medications such as cold & flu remedies, pain relief, chewing gum and kola nuts. Caffeine can freely pass across the placenta but cannot be broken down by

the placenta or the fetus. Evidence suggests that excess caffeine intake (>400mg per day) can increase the risk of miscarriage and sudden infant death syndrome and lower intakes, may negatively affect fetal growth (FSAI, 2011).

Caffeine can be transferred to infants via breast milk. Infants do not synthesize the enzymes required to breakdown caffeine until approximately 3 months of age. Infants ingesting excessive caffeine may have poor sleep and become irritable (FSAI 2011).

- Pregnant women are advised to limit caffeine to the maximum recommended intake of 200mg per day. This equates to 2-4 cups of tea **or** 2 cups instant coffee **or** 1 cup filtered/brewed coffee.
- Lactating women are advised to limit caffeinated drinks to no more than 2-3 cups daily (200-300mg caffeine).

It should be noted that the caffeine content in teas and coffees vary depending on the brand, brewing time and method. Also, tea and coffee bought in a café may be stronger than that made at home.

9.1.2 Biotoxins

Biotoxins are produced in certain circumstances by various species of marine algae and accumulate in molluscs such as clams, mussels, oysters and scallops (FSAI, 2019). These toxins cannot be deactivated or destroyed by cooking or heating and can lead to gastrointestinal illness, neurotoxicity, multiple organ damage and in some situations, death for the woman. Studies are lacking to assess the risks to the fetus. Animal studies indicate these toxins can cross the placenta, lead to neurological damage that can present as learning and memory deficits later in life (Maucher et al, 2007; Munday, 2013).

 Advise women not to eat clams, mussels and oysters during pregnancy and lactation as exposure to biotoxins may cause severe illness in the woman and potential damage to the fetus.

9.1.3 Mercury

Mercury is a cumulative neurotoxin that is present both naturally in the environment and as a result of disposal of waste or releases from industrial activities. Excessive dietary mercury intake is mainly from large, predatory fish that are high in the food chain. Mercury readily crosses the placenta and fetal levels have been found to be higher than maternal levels (Ding *et al* 2013). Exposure to high levels of mercury can lead to significant neurological and behavioural disorders. Even lower levels of exposure can lead to adverse effects on neurological, cardiovascular and immune system functions (Karagas *et al* 2012).

 The advice for both pregnant and lactating women is to avoid intake of large predatory fish: shark, swordfish, ray and marlin. Tuna should be limited to one fresh steak per week or two 240g (8oz) tins per week.

9.1.4 Vitamin A

Pre-formed vitamin A (retinol equivalent) may be teratogenic at high doses (Hathcock *et al*, 1990). The maximum upper limit of pre-formed vitamin A, based on the risk of hepatotoxicity and teratogenicity, is 3000 micrograms or 3000 retinol equivalents (RE)/day (EFSA, 2006 & 2018). By contrast, precursors to vitamin A, beta-carotene and other carotenoids, pose no risk during pregnancy. The population reference intake for pregnant women is 700 micrograms per day (EFSA, 2015).

- Pregnant women are advised to avoid concentrated food sources of preformed vitamin A such as offal, liver and liver containing foods such as pâté.
- Women are advised to avoid dietary supplements (including fish liver oils)
 containing pre-formed vitamin A. Supplements containing beta-carotene as
 the sole source of vitamin A, which are used in many pregnancy-specific
 supplements, are suitable.

9.2 Food borne illness

Table 11: Summary sources and precautions of foodborne illness in pregnancy

Food	Reason for avoidance	Recommendation	Sources
Raw egg	Food borne disease (Samonella)	Thoroughly cooked eggs or dishes and desserts made with egg that are thoroughly cooked	Raw or partly cooked eggs, dishes containing part cooked eggs, homemade mayonnaise, custard or cheese cake made with raw fresh egg
Unpasteurised Dairy products	Food borne diseases	Avoid	All animal milk, cheese and yogurt which have not been pasteurised
Raw meat	Food borne disease (Toxoplasmosis)	Prevent cross contamination by storing raw meat/juices (bottom shelf) and ready-to-eat foods (upper shelves) separately in the refrigerator	Undercooked meat, burgers, BBQ
Raw Fish	Food borne disease	Avoid	Sushi, undercooked shellfish, cold smoked salmon
Mould ripened cheese	Food borne disease	Avoid or eat when cooked	Brie, camembert, gorgonzola, Danish blue and Roquefort.
Inappropriately prepared or stored food	Food borne disease	Hand hygiene and safe food preparation see FSAI/ safefood	All raw, cooked, ready-to- eat, reheated, stored foods

9.2.1 Safe Food Preparation

Food poisoning is caused by eating foods contaminated by either bacteria, viruses or parasites. Foods can become contaminated at any point during handling, storage or cooking. *Listeria monocytogenes* is a bacterium that is found in water and soil. Listeriosis is the illness that can result from ingestion of listeria from contaminated food. It can increase the risk of miscarriage, stillbirth, neonatal death and premature death. Listeria can be found in uncooked meats, uncooked vegetables, unpasteurised milk produce and processed foods. Listeria is killed by pasteurisation and cooking.

Toxoplasma gondii is a parasite, which can be found in raw and undercooked meat, unpasteurised goats milk, soil, cat faeces and untreated water. Toxoplasmosis, the resulting infection can be difficult to diagnose, as many patients do not exhibit any symptoms. If infection occurs during pregnancy, the parasite can be transmitted to the fetus, which causes congenital toxoplasmosis. The risk of transmission to the fetus is higher as the pregnancy progresses (Dunn, D et al 1999). Clinical manifestations of congenital toxoplasmosis if not treated include neurological deficits in early childhood or adulthood (Berrébi A et al, 2010).

To reduce the risk of food poisoning the following tips should be followed **during pregnancy and lactation**:

Handling and hygiene:

- Set fridge to 4°C or lower
- Set freezer to 18°C or lower
- Buy cold or frozen food at the end of shopping trip
- Keep food leftovers in the fridge and use within two days
- Wash hands, utensils and preparation surfaces well after handling raw meat.
- Always keep raw and cooked meat separate
- Reheat leftover meals to internal temperature of 70°C before consuming.

Meat, poultry and fish

- All meat poultry, fish, crab and lobster must be thoroughly cooked to internal temperature of 75°C before eating. This includes: cold cured meats such as pepperoni, salami, prosciutto, chorizo smoked fish (Avoid raw fish sushi)
- Avoid pâtés and molluscs (clams, mussels, oysters and scallops), both cooked and raw/cold.

Eggs

- Cook eggs until the white and yolk are solid.
- Pregnant women are advised not to consume foods in which raw eggs have been used: Homemade mayonnaise or aioli, Mousse, Cheesecake, Tiramisu

Fruit and vegetables

- Wash all fresh fruit and vegetables well prior to consumption to remove any traces of soil and visible dirt.
- These vegetables must be cooked thoroughly: Sprouts such as alfalfa, radish, mung bean and clover.

Dairy: milk, cheese and yogurt

- Consume only pasteurised or UHT (ultra-heat treated) milk and milk based produce.
- Avoid any unpasteurised milk (including goats and sheep's milk) and any products made using unpasteurised milks e.g. cheeses and yogurts.
- Avoid mould-ripened soft cheeses with a white rind unless thoroughly cooked: brie and camembert, soft blue veined cheeses e.g. gorgonzola, Danish blue and Roquefort. These cheeses can be eaten safely, cold or cooked, during lactation.
- All hard cheeses are safe to eat during pregnancy and lactation.
- Pasteurised soft cheeses, which are not mould-ripened as stated above, are also safe to consume e.g. cottage cheese, mozzarella, feta, halloumi, and ricotta.

For more information on general food safety guidance, see FSAI Healthy eating, food safety and food legislation

https://www.fsai.ie/science_and_health/healthy_eating.html for more details. And **safefood.eu** https://www.safefood.eu/Food-safety.aspx

10.0 Specific Diets

• Refer pregnant women who require therapeutic diets to a dietitian. For example, women with:

coeliac disease
pre-existing type 1 and 2 diabetes
phenylketonuria
cystic fibrosis
anaemia
post-bariatric surgery

 Women with chronic disease will ideally have had dietetic input as part of their preconception care and will continue to have input from their multidisciplinary team as part of their obstetric management.

11.0 Allergies

- There is insufficient evidence to suggest that avoiding allergenic foods during pregnancy and lactation can protect an infant from developing allergies.
- It is not recommended that women avoid potentially allergenic foods during pregnancy unless she herself is allergic to a specific food. If the pregnant woman herself has a food allergy, care should be taken that she does not consume that specific allergen.
- Women who avoid food groups, such as dairy, due to allergy or intolerance are at risk of inadequate intake of nutrients. A referral to a dietitian is warranted in these cases.

For more information on food allergy treatment and avoidance see the Irish Food Allergy Network (IFAN) website: http://ifan.ie/.

12.0 Hospital Equipment and Facilities

- All women must have an accurate weight and height measurement taken at their booking antenatal visit.
- The centre must ensure equipment is available in a discrete area and calibrated as per manufacturer instructions to accurately record measurements.
- Best practice dictates that a digital weighing scale and a wall-mounted stadiometer (height measure) are used.
- When necessary nutrient deficiency should be diagnosed or monitored through appropriate blood tests. Hospitals should have access to phlebotomy and laboratory testing.

13.0 Provision of Information on Best Practice for Infant Feeding

- It is the responsibility of all healthcare professionals to promote best practice for infant feeding.
- The National Infant Feeding Policy for Maternity and Neonatal Services and local implementation policy can be found here: https://www.breastfeeding.ie/Resources/Health-professional/Infant-Feeding-Policy-for-Maternity-Neonatal-Services-2019.pdf
- Information and support around breastfeeding should be given ante-natally, as it has been shown that many women make the decision on whether or not they will breastfeed before the birth of their infant (FSAI, 2011).
- Pregnant women should be educated during this time on the many benefits breastfeeding offers the mother and her infant including the health benefits in both short-term and throughout later life.

14.0 Implementation Strategy

- Distribution of guideline to all members of the Institute and to all maternity units.
- Implementation through National Women and Infants Health Programme.
- Distribution to other interested parties and professional bodies.
- Encourage Making Every Contact Count training
- Education or updates of staff in maternity hospitals by local dietitians.
- Encourage training on Mychild via HSEland.
- Education or updates of staff in maternity hospitals and universities by local dietitians.

15.0 Key Performance Indicators

Standard key performance indicators

Many KPI's are measurable though the MNCMS system or local hospital maternity booking system (K2). The following KPI's should be collected and audited on a regular basis as part of standard care and hospital metrics in all maternity units.

1. **Key recommendation**: "All women should have their weight checked and BMI calculated at the first antenatal visit, ideally in the first trimester. BMI checks at the booking visit should be used as a screening mechanism for those at increased risk."

Suggested KPI: Number/ proportion of women with weight and height and BMI at booking. GDM screening and anaesthetic clinic review should be provided as appropriate based on local policy for those with higher BMI's.

2. **Key recommendation**: "Refer women with a BMI <17kg/m² to a dietitian"

Suggested KPI: Number of women with BMI< 17kg/m² referred to the dietitian.

3. **Key recommendation**: "Antenatal appointments offer an opportunity to utilise the brief intervention model, outlined in the Making Every Contact Count Framework. Pregnancy is a time when women are keen to receive more information about healthy lifestyle interventions and are motivated to make positive choices. The frequent contact between pregnant women and the healthcare team at antenatal visits may be utilised to support patients to make lifestyle behaviour changes that may impact on weight gain, with a particular focus on physical inactivity and unhealthy eating."

Suggested KPI: Number of documented "Making Every Contact Count" records on MNCMS

4. **Key recommendation:** "All women who may possibly become pregnant within the next three months, whether intentionally or not, are advised to take oral FA (FA) 400 micrograms daily to prevent neural tube defects (NTDS). After the first trimester and during breastfeeding, all women are advised to take FA 400 micrograms to meet the World Health Organization's recommended daily intake for pregnancy and breastfeeding."

Suggested KPI: Number of women on folic acid supplementation at booking

Additional key performance indicators

Other KPI's should be collected and documented in the patient's record. They will require audit which is not directly available from hospital summary metrics. Audit

to review a proportion of patient records can be conducted on a regular basis to monitor local implementation of the guideline.

1. **Key recommendation**: "Women should be encouraged to achieve a healthy weight before conception to prevent associated complications. Weight loss in the postnatal period before subsequent pregnancies is an effective strategy for improved outcomes. Postnatal weight loss in overweight and obese women should be encouraged and where available onward referral to community dietetics should be provided."

Suggested KPI: Proportion of women with high BMI referred to community on discharge. This can be documented on discharge letter/ summary. Patient's GP can refer to the community dietitian the hospital should recommend this to GP.

2. **Key recommendation**: "During pregnancy, a supplement containing 16-20mg of iron in addition to a balanced diet has the potential of reducing incidence of anaemia in the healthy population."

Suggested KPI: Proportion of women on multivitamin or supplement containing 16-20mg iron

3. **Key recommendation**: "Women suspected of iron deficiency should have a full blood count (FBC) and, if possible, check serum ferritin. The treatment for iron deficiency anaemia is oral iron supplementation."

Suggested KPI: Number of women with iron deficiency anaemia

4. **Key recommendation**: "All pregnant women in Ireland should take a supplement containing 10micrograms (400 IU) vitamin D₃ per day."

Suggested KPI: Proportion of women on 10micrograms (400 IU) vitamin D₃

16.0 Qualifying Statement

These guidelines have been prepared to promote and facilitate standardisation and consistency of practice, using a multidisciplinary approach. Clinical material offered in this guideline does not replace or remove clinical judgment or the professional care and duty necessary for each pregnant woman. Clinical care carried out in accordance with this guideline should be provided within the context of locally available resources and expertise.

This guideline does not address all elements of standard practice and assumes that individual clinicians are responsible for:

- Discussing care with women in an environment that is appropriate and which enables respectful confidential discussion.
- Advising women of their choices and ensure informed consent is obtained.
- Meeting all legislative requirements and maintaining standards of professional conduct.
- Applying standard precautions and additional precautions, as necessary, when delivering care.
- Documenting all care in accordance with local and mandatory requirements.

17.0 References

Alavi N, Haley S, Chow K, McDonald SD (2013) Comparison of national gestational weight gain guidelines and energy intake recommendations. *Obesity Review*; 14(1):68-85.

Alwan NA, Greenwood DC, Simpson NA, McArdle HJ, Godfrey KM, Cade JE (2011) Dietary iron intake during early pregnancy and birth outcomes in a cohort of British women. *Human Reproduction*. 2011 Apr; 26(4):911-9.

Aills L, Blankenship J, Buffington C, Furtado M, Parrott J. (2008). Allied Health Sciences Section Ad Hoc Nutrition Committee: ASMBS Allied Health Nutritional Guidelines for the Surgical Weight Loss Patient. *Surg Obes Related Dis*; 4(suppl):S73-S108.

ACOG (American College of Obstetricians and Gynecologists) (2000) *Planning Your Pregnancy and Birth, 3rd ed.* Washington.

ACOG (American College of Obstetricians and Gynecologists),(2009) *Clinical Management Guideline, Bariatric Surgery and Pregnancy,* Washington, DC, USA, 2009.

ACOG (American College of Obstetricians and Gynecologists) (2011) Vitamin D: screening and supplementation during pregnancy. Committee Opinion No. 495. *Obstetrics Gynaecology;* 118:197–8.

American College of Obstetricians and Gynecologists (ACOG). (2013). Weight gain during pregnancy. Committee Opinion No. 548. *Obstet Gynecol*; 121:210–2.

American College of Obstetricians and Gynecologists (ACOG). (2015). Physical activity and exercise during pregnancy and the postpartum period. Committee Opinion No. 650 (Reaffirmed 2017). *Obstet Gynecol*.126:e135–e142

Arendras K, Qiu Q, Gruslin A (2008). Obesity in pregnancy: pre-conceptional to postpartum consequences. *Journal Obstetrics Gynaecology Canada*. 30(6):477-88.

Ares S, Quero J, de Escobar GM. Iodine balance, iatrogenic excess, and thyroid dysfunction in premature newborns. *Semin Perinatol.* 2008;32:407-12. <u>PMID:</u> 19007678

Barker DJP, Gluckman PD, Godfrey KM, Harding JE, Owens JA, Robinson JS (1993). Fetal nutrition and cardiovascular disease in adult life. *Lancet* 341:938-941

Barker DJP (1995) Fetal origins of coronary heart disease. *British Medical Journal* 311:171-174

Barker D, Osmond C, Forsén T, Kajantie E, and Eriksson J (2005) Trajectories of Growth among Children Who Have Coronary Events as Adults. *New England Journal of Medicine*. 353: 1802-1809

Barker DJ, Larsen G, Osmond C, Thornburg KL, Kajantie E, Eriksson JG (2012) The placental origins of sudden cardiac death. *International Journal of Epidemiology*. 41(5):1394-1399.

Barroso F, Allard S, Kahan BC, Connolly C, Smethurst H, Choo L, Khan K, Stanworth S (2011) Prevalence of maternal anaemia and its predictors: a multicentre study. *European Journal Obstetrics Gynecology Reproduction Biology*; 159(1):99-105.

Beard, J.L., Hendricks, M.K., Perez, E.M., Murray-Kolb, L.E., Berg, A., Vernon Feagans, L., Irlam, J., Isaacs, W., Sivem, A., Tomlinson, M. (2005) Maternal iron deficiency anemia affects postpartum emotions and cognition. *Journal of Nutrition* 135, 267-272.

Beyerlein A, Lack N, Von Kries R. (2010) Within population average ranges compared with Institute of Medicine recommendations for gestational weight gain. *Obstet Gynecol*; 116:1111-1118.

Bougma, K. Aboud, F.E. Harding, K.B. Marquis, G.S. (2013) Iodine and Mental Development of Children 5 Years Old and Under: A Systematic Review and Meta-Analysis. *Nutrients* 2013, *5*(4), 1384- 1416

Boilson, A. Staines, A. Kelleher, C.C. Daly, L. Shirley, I. Shrivastava, A. Bailey, S.W. Alverson, P.B. Ayling, J.E. McDermott, A.P. Scott, J.M. Sweeney, M.R. (2012) Unmetabolized FA prevalence is widespread in the older Irish population despite the lack of a mandatory fortification program. *The American Journal of Clinical Nutrition*, Vol 96. pp. 613-621

Boulet, Alexander, Salihu, Pass (2003) Macrosomic births in the United States: Determinants, outcomes and proposed grades of risk. *American Journal of obstetrics and Gynaecology*. 188; 1372-8.

Boney, Verma, Tucker, Vohr (2005). Metabolic syndrome in childhood: association with birth weight, maternal obesity and gestational diabetes. *Pediatrics*; 115; e290-6

Bricker L, Reed K, Wood L, Neilson JP (2015). Nutritional advice for improving outcomes in multiple pregnancies. Cochrane Database of Systematic Reviews, Issue 11. Art. No.: CD008867. DOI: 10.1002/14651858.CD008867.pub3.

Brown JE, Isaaca JS, Krinke UB, Murtaugh MA, et al. (2008). *Nutrition Throughout the Lifecycle, 3rd edition*. California: Wadsworth Cengage Learning; p517

Brownfoot FC, Davey MA, Kornman L. (2016). Routine weighing to reduce excessive antenatal weight gain: a randomised controlled trial. BJOG: an *international journal of obstetrics and gynaecology*; 123(2):254–61

Burke, B. Lyon Daniel, K. Latimer, A. Moran, K. Mulinare, J. Prue, C. Steen, J. Watkins, M. (2009) *Preventing Neural Tube Birth Defects: A Prevention Model and Resource Guide.* Atlanta: Centers of Disease Control and Prevention

Busetto L, Dicker D, Azran C, Batterham R, L, Farpour-Lambert N, Fried MHjelmesæth J, Kinzl J, Leitner D, R, Makaronidis J, M, Schindler K, Toplak H, Yumuk V. (2017) Practical Recommendations of the Obesity Management Task Force of the European Association for the Study of Obesity for the Post-Bariatric Surgery Medical Management. *Obes Facts*; 10:597-632

Butte NF, King JC. (2005). Energy requirements during pregnancy and lactation. *Public Health Nutrition*. 8 (7A), 1010-1027.

Choi J, Fukuoka Y, Lee JH. The effects of physical activity and physical activity plus diet interventions on body weight in overweight or obese women who are pregnant or in postpartum: a systematic review and meta-analysis of randomized controlled trials. *Prev Med* 2013;56:351–64

Crosby DA, Collins M, O'Higgins A, Mullaney L, Farah N, Turner MJ. Interpregnancy changes in maternal weight and body mass index. *American Journal Perinatology*. 2015 Feb;30(2):199-204.

Cuskelly G, McNulty H, Scott JM. Effect of increasing dietary folate on red-cell folate: implications for prevention of neural tube defects. *Lancet.* 1996 Mar 9;347(9002):657-9.

Daley A, Jolly K, Lewis A, Clifford S, Kenyon S, Roalfe A, et al. (2014). The feasibility and acceptability of regular weighing of pregnant women by community midwives to prevent excessive weight gain: RCT. *Pregnancy Hypertens*;4:233–4.

Daly N, Farren M, McKeating A, O'Kelly R, Stapleton M, Turner MJ. (2017) A Medically Supervised Pregnancy Exercise Intervention in Obese Women: A Randomized Controlled Trial. *Obstetrics Gynecology*;130(5):1001-1010.

Daru J, Zamora J, Fernández-Félix B.M, Vogel J, Oladapo OT, Morisaki N, Tunçalp Ö, Torloni MR, Mittal S, Jayaratne K, Lumbiganon P, Togoobaatar G, Thangaratinam S, Khan KS. (2018). Risk of maternal mortality in women with severe anaemia during pregnancy and postpartum: a multilevel analysis, *Lancet Global Health* 6(5):e548-e554

Davis RR, Hofferth S, Shenassa E. (2013) Gestational Weight Gain and Risk of Infant Death in the United States. *American Journal of Public Health*, 2013; 104 Suppl 1:S90-5.

De-Regill LM, Palacios C, Ansary A, Kulier R, Pena-Rosas JP, Editorial Group: Cochrane Pregnancy and Childbirth Group (2012) *Vitamin D supplementation for women during pregnancy* The Cochrane database of systematic reviews, Issue 6. Wiley & Sons.

Dennedy MC, Dunne F (2010) The Maternal and Fetal impacts of Obesity and Gestational Diabetes on Pregnancy Outcome. *Best practice and research Clinical Endocrinology and Metabolism* Aug;24(4):573-89.

Department of Health (2018) Folic Acid Policy Committee. *Folic Acid Supplementation Report* September 2018

Department of Health (DOH). (2016) Health and Wellbeing Programme. *Healthy Ireland Survey*, 2015. Dublin: Irish Social Science Data Archive [distributor], March 2016

Department of Health (DOH). (2016) Creating a Better Future Together. National Maternity Strategy 2016-2026.

Department of Health (DOH), Ireland. (2013) Healthy Ireland - A Framework for Improved Health and Wellbeing 2013 - 2025 [Online]. Available at: http://www.thehealthwell.info/node/773887 [Accessed: 11th May 2018]

Department Of Health (DoH) 2012 The National Healthy Eating Guidelines - Your Guide to Healthy Eating using the Food Pyramid.

(DoHHS) Department of Health and Human Services. (2008) Physical activity guidelines for Americans. Washington, DC: DHHS; 2008. Available at: http://health.gov/paquidelines

Devlieger R, Guelinckx I, Jans G, Voets W, Vanholsbeke C, Vansant G. (2014) Micronutrient levels and supplement intake in pregnancy after bariatric surgery: a prospective cohort study; *PLOS One*; 9:e114192

Dewey and Chaparro (2007) Session 4: Mineral metabolism and body composition iron status of beast-fed infants. *Proceeding of Nutrition Society.* 66 (3): 412-422.

Dietitians of Canada. Multi-fetal Pregnancy Practice Guidance Summary. In: Practice-based Evidence in Nutrition (PEN). 2013 Available at: http://www.pennutrition.com

Eddib A, Penvose-Yi J, Shelton JA, Yeh J. (2007). Triplet gestation outcomes in relation to maternal pre-pregnancy body mass index and weight gain. *J Matern Fetal Neonatal Med.*; 20(7):515-9

Ekiz, E., Agaoglu, L., Karakas, Z., Gurel, N., Yalcin, I. (2005) The effect of iron deficiency anemia on the function of the immune system. *The Hematology Journal* 5, 579–583.

EUROCAT Central Registry (2010) EUROCAT Special Report: Prevalence of Neural Tube Defects in younger mothers in Europe 2000–2008. Antrim: EUROCAT Central Registry

EUROCAT Central Registry (2014) EUROCAT Special Report: Geographic Inequalities in Public Health Indicators Related to Congenital Anomalies. Antrim: EUROCAT Central Registry

European Food Safety Authority (EFSA). (2018) Overview on Tolerable Upper Intake Levels as derived by the Scientific Committee on Food (SCF) and the EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA). Version 4. EFSA

European Food Safety Authority (EFSA). (2015). Scientific Opinion on Dietary Reference Values for protein. EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA). *EFSA Journal*;10(2):2557

European Food Safety Authority (EFSA). (2015). Scientific Opinion on Dietary Reference Values for vitamin A. EFSA Journal;13(3):4028

European Food Safety Authority. (2014) *Draft Scientific Opinion - Scientific Opinion on Dietary Reference Values for folate. EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA).* Parma: European Food Safety Authority

European Food Safety Authority. (2014) Draft Scientific Opinion - Scientific Opinion on Dietary Reference Values for Iodine. EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA). EFSA Journal 2014;12(5):3660

Parma: European Food Safety Authority DOI: 10.2903/j.efsa.2014.3660

European Food Safety Authority (EFSA). (2013). EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA); Scientific Opinion on Dietary Reference Values for energy. *EFSA Journal* 11(1):3005. [112 pp.] Available online: www.efsa.europa.eu/efsajournal

European Food Safety Authority (EFSA). (2010) EFSA Panel on Dietetic Products, Nutrition, and Allergies (NDA); Scientific Opinion on Dietary reference values for water. *EFSA Journal*; 8(3):1459. [48 pp.]. Available online: www.efsa.europa.eu

European Food Safety Authority (EFSA) Panel on Dietetic Products, Nutrition, and Allergies (2010) Scientific Opinion on Dietary Reference Values for fats, including saturated fatty acids, polyunsaturated fatty acids, monounsaturated fatty acids, trans fatty acids, and cholesterol. *EFSA Journal*; 8(3):1461

European Food Safety Authority (EFSA) Panel on Dietetic Products, Nutrition and Allergies (NDA) (2009); Scientific Opinion on substantiation of health claims related to iodine and thyroid function and production of thyroid hormones (ID 274), energy-yielding metabolism (ID 274), maintenance of vision (ID 356), maintenance of hair (ID 370), maintenance of nails (ID 370), and maintenance of skin (ID 370) pursuant to Article 13(1) of Regulation (EC) No 1924/2006 on request from the European Commission. *EFSA Journal*; 7(9):1214.

European Food Safety Authority (EFSA). (2006) Scientific Committee on Food

Scientific Panel on Dietetic Products, Nutrition and Allergies Tolerable Upper Intake Levels for Vitamins and Minerals. Available at:

http://www.efsa.europa.eu/sites/default/files/efsa_rep/blobserver_assets/ndatolerableuil.pdf

Farah N, Stuart B, Donnelly V, Kennelly MM, Turner MJ (2011) The influence of maternal body composition on birth weight. European Journal of Obstetrics and *Gynecology Reproduction Biology*. Jul;157(1):14-7

Fattah C, Farah N, Barry S, O'Connor N, Stuart B, Turner MJ (2010) Maternal weight and body composition in the first trimester of pregnancy. *Acta Obstetricia et Gynecologica Scandinavia*; 89:952-5.

Fealy SM, Taylor RM, Foureur M, Attia J, Ebert L, Bisquera A, Hure AJ (2017) Weighing as a stand-alone intervention does not reduce excessive gestational weight gain compared to routine antenatal care: a systematic review and meta-analysis of randomised controlled trials. *BMC Pregnancy and Childbirth* 17, 36.

Fronczak CM, Barón AE, Chase HP, Ross C, Brady HL, Hoffman M, Eisenbarth GS, Rewers M, Norris JM. (2003) In utero dietary exposures and risk of islet autoimmunity in children. *Diabetes Care*. 26(12):3237-42.

Furber CM, McGowan L (2011) A qualitative study of the experiences of women who are obese and pregnant in the UK. *Midwifery*. 27(4):437-44.

Food Safety Authority of Ireland (FSAI). (2019) *Healthy eating, food safety and food legislation: A guide supporting the Healthy Ireland Food Pyramid*. Available online: https://www.fsai.ie/science and health/healthy eating.html

Food Safety Authority of Ireland (FSAI). (2016) Scientific Committee of the FSAI *Update Report on Folic Acid and the Prevention of Birth Defects in Ireland* March 2016 ISBN 978-1-910348-04-8

Food Safety Authority of Ireland (FSAI). (2016) Survey of the trans fatty acid content in processed food products in Ireland. Dublin

Food Safety Authority Ireland (FSAI) 2011 Best Practice for infant feeding in Ireland. Dublin.

Food Safety Authority of Ireland (FSAI) 2011 *Scientific Recommendations for a National Infant Feeding Policy, 2nd Edition*. Dublin.

Girsen AI, Mayo JA, Carmichael SL, Phibbs CS, Shachar BZ, Stevenson DK, Lyell DJ, Shaw GM, Gould JB On behalf of the March of Dimes Prematurity Research Center at Stanford University School of Medicine. (2016) Women's prepregnancy underweight as a risk factor for preterm birth:a retrospective study. *BJOG*.2016 123:2001–2007.

Goldstein et al (2017) Association of Gestational Weight Gain with Maternal and Infant Outcomes: A Systematic Review and Meta-analysis. *JAMA*. 2017 Jun 6;317(21):2207-2225.

Gonzalez-Iglesias H, de la Flor St Remy RR, Lopez-Sastre J et al. Efficiency of iodine supplementation, as potassium iodide, during lactation: A study in neonates and their mothers. *Food Chem.* 2012;133:859-65

Gonzalez I, Rubio MA, Cordido F, Breton I, Morales MJ, Vilarrasa N, Monereo S, Lecube A, Caixas A, Vinagre I, Goday A, Garcia-Luna PP: Maternal and perinatal outcomes after bariatric surgery: a Spanish multicenter study. *Obes Surg* 2015;25:436-442

Goodnight W, Newman R, Society of Maternal-Fetal Medicine. Optimal Nutrition for Improving Twin Pregnancy Outcome. Obstet Gynecol. 2009;114 (5):1121-1134

Goto E (2019) Dose-response associations of maternal height and weight with small for gestational age: a meta-analysis. *EJCN* 10.1038/s41430-019-0432-y

Guelinckx I, Devlieger R, Vansant G. (2009). Reproductive outcome after bariatric surgery: a critical review, *Human Reproduction Update*; 15, 2, 189–201, https://doi.org/10.1093/humupd/dmn057

Haas, J.D., Brownlie, T. (2001) Iron deficiency and reduced work capacity: A critical review of the research to determine a causal relationship. *Journal of Nutrition* 131, 676S-690S.

Harding KB, Peña-Rosas JP, Webster AC, Yap CMY, Payne BA, Ota E, De-Regil LM. Iodine supplementation for women during the preconception, pregnancy and postpartum period. Cochrane Database of Systematic Reviews 2017, Issue 3

Health Canada, Expert Advisory Group on *National Nutrition Pregnancy Guidelines* (2009) Prenatal Nutrition Guidelines for Health Professionals, Iron.

Hofmeyr GF, Duley I, Atallah A. Dietary calcium supplementation for prevention of pre-eclampsia and related problems: a systematic review and commentary (2007) *BJOG: An International Journal of Obstetrics & Gynaecology.* 114 (8) pages 933–943

Holmes VA, Barnes MS, Alexander HD, McFaul P, Wallace JM (2009) Vitamin D deficiency and insufficiency in pregnant women: a longitudinal study. *British Journal Nutrition*. 102(6):876-81.

Hopkins, S. M., Gibney, M. J., Nugent, A. P., McNulty, H, Molloy, A. M., Scott, J. M., Flynn, A., Strain, JJ, Ward, M, Walton, J. and McNulty, B. A. 2015. 'Impact of voluntary fortification and supplement use on dietary intakes and biomarker status

of folate and vitamin B-12 in Irish adults'. *American Journal of Clinical Nutrition*. 2015 Jun;101(6):1163-72.

Health Service Executive (HSE) National Steering Group on Vitamin D (2010). *Policy on Vitamin D Supplementation for Infants in Ireland*. Available at: https://www.hse.ie/eng/health/child/vitamind/vitamin%20d%20policy.pdf; Accessed May 2019.

Health Service Executive (HSE). (2016). *Making every contact count: a health and behaviour change framework and implementation for health professionals in the Irish health service.* Health and Wellbeing Division of the HSE. ISBN No 978-1-78602-038-3

Health Service Executive (HSE). (2016). *Healthy Eating Guidelines*. Available at: https://www.hse.ie/eng/about/who/healthwellbeing/our-priority-programmes/heal/healthy-eating-guidelines/.

HSE 2010 Guidelines for the management of pre-gestational diabetes mellitus from preconception to the postnatal period. July 2010 available at: https://www.hse.ie/eng/services/publications/nursingmidwifery%20services/onsdg uidelinesgestationaldiabetes.pdf.

Hulman A, Lutsiv O, Park CK, Krebs L, Beyene J, McDonald SD. Are women who quit smoking at high risk of excess weight gain throughout pregnancy? BMC Pregnancy Childbirth. (2016);16:263

Huxley RR, Shiell AW, Law CM (2000) The role of size at birth and postnatal catchup growth in determining systolic blood pressure: a systematic review of the literature. *Journal of Hypertension*. Jul;18(7):815-31.

Hyppönen E, Läärä E, Reunanen A, Järvelin MR, Virtanen SM (2001) Intake of vitamin D and risk of type 1 diabetes: a birth-cohort study. *Lancet*. 3;358(9292):1500-3.

IOM (Institute of Medicine). (2009). Weight Gain during Pregnancy: Re-examining the Guidelines. Washington, DC: The National Academies Press.

IOM (Institute of Medicine). (2005) *Dietary Reference Intakes for Energy, Carbohydrate, Fiber, Fat, Fatty Acids, Cholesterol, Protein, and Amino Acids.* Washington, DC: The National Academies Press.

Irish Nutrition and dietetic Institute (INDI). (2015). *Nutrition Support Reference Guide*. Dublin. INDI. 309-313.

Institute of Obstetricians and Gynaecologists (IOG), Royal College of Physicians of Ireland and Clinical Strategy and Programmes Directorate, Health Service Executive (2011) *Obesity and Pregnancy Clinical Practice Guideline*. HSE Dublin.

IUNA Irish Universities Nutrition Alliance, edited by Walton J (2011) National Adult nutrition survey. Food and Nutrient intakes, Physical Measurements, Physical Activity Patterns and Food Choice Motives. IUNA March 2011.

Irish Universities Nutrition Alliance (IUNA). (2011) *National Adult Nutrition Survey: Summary Report.* Available at: http://iuna.net/index.php/national-adult-nutrition-survey-nans/. Accessed 04/04/2018

Irish Universities Nutrition Alliance. *North/South Ireland Food Consumption Survey*. Dublin: Food Safety Promotion Board; 2001. Available at: http://www.iuna.net/docs/NSIFCSummary.pdf

Jefferys AE, Siassakos D, Draycott T, Akande VA, Fox R. (2013). Deflation of gastric band balloon in pregnancy for improving outcomes (Review). *Cochrane Database Syst Rev.* Issue 4. DOI:10.1002/14651858.CD010048.pub2.

Kajantie E, Osmond C, Barker DJ, Forsén T, Phillips DI, Eriksson JG (2005) Size at birth as a predictor of mortality in adulthood: a follow-up of 350000 person- years. *International Journal of Epidemiology*. Jun;34(3):655-63.

Kapadia, M. Z., Park, C. K., Beyene, J., Giglia, L., Maxwell, C. and McDonald, S. D. (2015). Can we safely recommend gestational weight gain below the 2009 guidelines in obese women? A systematic review and meta-analysis. *Obes Rev*, 16: 189–206.

Kelly, F. Gibney, E.R. Boilson, A. Staines, A. Sweeney, M.R. (2015) FA levels in some food staples in Ireland are on the decline: implications for passive FA intakes? *Journal of Public Health*. 2016 Jun; 38(2):265-9.

Khan R, Dawlatly B, Chappatte O. Pregnancy outcome following bariatric surgery. (2013). *The Obstetrician & Gynaecologist*;15:37–43

Knight-Agarwal CR, Kaur M, Williams LT, Davey R, Davis D. (2013) The views and attitudes of health professionals providing antenatal care to women with a high BMI: a qualitative research study. *Women and Birth*. 27(2):138-44

Kral JG, Biron S, Simard S, Hould FS, Lebel S, Marceau S, Marceau P (2006) *Large maternal weight loss from obesity surgery prevents transmission of obesity to children who were followed for 2 to 18 years. Pediatrics* 118(6):e1644-9.

Knudsen VK, Orozova-Bekkevold IM, Mikkelsen TB, Wolff S, Olsen SF (2008) Major dietary patterns in pregnancy and fetal growth. *European Journal Clinical Nutrition*. Apr;62(4):463-70.

Koletzko B, Brands B, Poston L, Godfrey K, Demmelmair H; Early Nutrition Project (2012) Early Nutrition Programming of long term health. *Proceedings of Nutrition Society*. Aug; 71(3):371-8.

Lassi, Z.S. Salam, R.A. Haider, B.A. Bhutta, Z.A. (2013) *FA supplementation during pregnancy for maternal health and pregnancy outcomes (Review).* London: The Cochrane Collaboration

LifeCycle Project-Maternal Obesity and Childhood Outcomes Study Group. Association of Gestational Weight Gain With Adverse Maternal and Infant Outcomes. JAMA. 2019;321(17):1702–1715.

Mark P. J. Vanderpump; The epidemiology of thyroid disease, *British Medical Bulletin*, Volume 99, Issue 1, 1 September 2011, Pages 39–51.

Markova V, Norgaard A, Jørgensen KJ, Langhoff-Roos J. (2016) Treatment for women with postpartum iron deficiency anaemia. *Cochrane Database Syst Rev*. 13;(8):CD010861. doi: 10.1002/14651858.CD010861.pub2.

Maucher, J. M., & Ramsdell, J. S. (2007). Maternal-fetal transfer of domoic acid in rats at two gestational time points. *Environmental health perspectives*, *115*(12), 1743–1746. doi:10.1289/ehp.10446

McBaina, Gustaaf, Dekkera, Clifton, Mola and Grzeskowiak (2016). Impact of interpregnancy BMI change on perinatal outcomes: a retrospective cohort study. *European Journal of Obstetrics & Gynecology and Reproductive Biology*. 205: 98-104

McDonnell, R. Delany, V. O'Mahony, MT. Mullaney, C. Lee, B. Turner, MJ. (2014) Neural tube defects in the Republic of Ireland in 2009–11. *Journal of Public Health.* 2015 Mar; 37(1).

McDonnell, R. Delany, V. O'Mahony, M.T. Lynch, C. McKeating, A. (2018) An Audit of Neural Tube Defects in the Republic Of Ireland for 2012-2015. *Irish Medical Journal* 111(3)

McGowan C, McAuliffe FM (2012). Maternal dietary patterns and associated nutritient intakes during each trimester of pregnancy. *Public Health Nutrition*. Available on CJO 2012 doi:10.1017/S1368980012000997

McGowan C, Byrne J, Walsh J, McAuliffe FM. (2011). Insufficient vitamin D intakes among pregnant women. *European Journal of Clinical Nutrition*: 65, 1076-1078

McKenzie, A. L., Muñoz, C. X., Ellis, L. A., Perrier, E. T., Guelinckx, I., Klein, A., ... Armstrong, L. E. (2017). Urine color as an indicator of urine concentration in pregnant and lactating women. *European journal of nutrition*, *56*(1), 355–362. doi:10.1007/s00394-015-1085-9

Mechanick J, Youdim A, Jones DB et al. (2013), "Clinical practice guidelines for the perioperative nutritional, metabolic, and nonsurgical support of the bariatric surgery patient–2013 update: cosponsored by American Association of Clinical Endocrinologists, the Obesity Society, and American Society for Metabolic & Bariatric Surgery," Endocrine Practice, vol. 19, no. 2, pp. 337–372, 2013

Molloy, A.M. Kirke, P.N. Brody, L.C. Scott, J.M. Mills, J.L. Effects of folate and vitamin B12 deficiencies during pregnancy on fetal, infant, and child development. *Food and Nutrition Bulletin*; 2008, 29(2) pp.101-111

Molloy, A.M. Einri, C.N. Jain, D. Laird, E. Fan, R. Wang, Y. Scott, J.M. Shane, B. Brody, L.C. Kirke, P.N. Mills, J.L. (2014) Is low iron status a risk factor for neural tube defects? *Birth Defects Res A Clin Mol Teratol*; Feb;100(2):100-6.

Moses RG, Luebcke M, Davis WS, Coleman KJ, Tapsell LC, Petocz P, Brand-Miller JC (2006) Effect of a low-glycemic-index diet during pregnancy on obstetric outcomes. *American Journal Clinical Nutrition*. Oct;84(4):807-12.

Mousa A, Naqash A, Lim S (2019). Macronutrient and Micronutrient Intake During Pregnancy: An Overview of Recent Evidence. *Nutrients*. 11; 443; doi:10.3390/nu11020443

Muktabhant B, Lawrie TA, Lumbiganon P, Laopaiboon M. (2015). Diet or exercise, or both, for preventing excessive weight gain in pregnancy. *Cochrane Database of Systematic Reviews*; 15;(6)

Mullaney, O'Higgins, Cawley, Daly, McCartney, Turner. (2016) Maternal weight trajectories between early pregnancy and four and nine months postpartum. *Public Health*; 135; 144-146

Munday, R., & Reeve, J. (2013). Risk assessment of shellfish toxins. *Toxins*, *5*(11), 2109–2137. doi:10.3390/toxins5112109

National Institute for Health and Clinical Excellence (NICE) (2010) Weight management before during and after pregnancy.

National Institute for Health and Clinical Excellence (NICE) (2008) Improving the nutrition of pregnant and breastfeeding mothers and children in low income families. NICE public health guidance.

National Institute for Health and Clinical Excellence (NICE). (2008). *Antenatal care:* routine care for the healthy pregnant woman. NICE public health guidance. NICE guideline CG62

National Institute for Health and Clinical Excellence (NICE) (2008) (B) *Maternal and Child Nutrition*. PH11 available at http://publications.nice.org.uk/maternal-and-child-nutrition-ph11/introduction

National Research Council (NRC). (1989). *Recommended Dietary Allowances*. 10th ed. National Academy of Sciences Press, Washington, D.C.

Nawoor Z, Burns R, Smith DF, Sheehan S, O'Herlihy C, Smyth PP (2006). Iodine intake in pregnancy in Ireland--a cause for concern? *Irish Journal Medical Science*. 175(2):21-4.

NCD Risk Factor Collaboration (NCD-RisC). (2017) Worldwide trends in body-mass index, underweight, overweight, and obesity from 1975 to 2016: a pooled analysis of 2416 population-based measurement studies in 128.9 million children, adolescents, and adults. *Lancet*, 390:2627–2642

Nelson SM, Mathews P and Poston L (2010) Maternal metabolism and obesity: modifiable determinants of pregnancy outcome. *Human Reproduction Update* 16, 255-275.

Obeid, R. Koletzko, B. Pietrzik, K. 2014. 'Critical evaluation of lowering the recommended dietary intake of folate'. *Clinical Nutrition*; Apr 33(2):252-9.

Obican S, Jahnke G, Soldin O, Scialli A. (2012) Teratology Public Affairs Committee Position Paper: Iodine Deficiency in Pregnancy. *Birth Defects Research: Clinical and Molecular Teratology* 94:677-682.

O'Brien EC, Geraghty AA, O'Sullivan EJ, Kilbane MT, Horan MT, McKenna MJ, McAuliffe FM (2018). Bone resorption and calcium in pregnancy –A window to futurematernal bone health. *Proceedings of the Nutrition Society*, 77 (OCE3), E52

O'Brien EC, Geraghty AA, O'Sullivan EJ, Riordan JA, Horan MK, Larkin E, Donnelly J, Mehegan J, Twomey PJ, McAuliffe FM (2019a). Five-year follow up of a low glycaemic index dietary randomised controlled trial in pregnancy—no long-term maternal effects of a dietary intervention. BJOG; 126: 514–524.

O'Brien EC, Segurado R, Geraghty AA et al. (2019b). Impact of maternal education on response to lifestyle interventions to reduce gestational weight gain - Individual participant data meta-analysis. BMJ Open. In press.

O'Higgins A, Doolan A, Mullaney L, Daly N, McCartney D, Turner MJ. The relationship between gestational weight gain and fetal growth: time to take stock? *J Perinat Med.* 2014;42(4):409-415.

O'Kane, M. Pourshahidi, K. Farren, K.M. Mulhern, M.S. Strain, J.J. Yeates, A.J. (2015) Knowledge and awareness of iodine nutrition among women of childbearing age. *Proceedings of The Nutrition Society 74(OCE4)*

O'Malley, EG. Cawley, S. Kennedy, RK. Reynolds, CME. Molloy, A. and Turner, MJ (2018) Maternal anaemia and folate intake in early pregnancy. *Journal of Public Health* Jan 31.

O'Neill JL, Keaveney EM, O'Connor N, Cox M, Regan A, Shannon E, Turner MJ (2011) Are Women in Early Pregnancy Following the National Pyramid Recommendations?. *Irish Medical Journal* Oct; 104 (9):270-2

O'Riordan MN, Kiely M, Higgins JR, Cashman KD (2008) Prevalence of suboptimal vitamin D status during pregnancy. *Irish Medical Journal*. 101(8):240, 242-3.

Ota, Tobe, Mori, Farrar (2012) Antenatal dietary advice and supplementation to increase energy and protein intake. *Cochrane Database Systematic Review*. Sep 12;9

Pavord, S, Myers, B, Robinson S, et al. (2012) *UK Guidelines on the management of iron deficiency in pregnancy*. British Society for Haematology Committee for Standards in Haematology. Available at: https://doi.org/10.1111/j.1365-2141.2011.09012.x. Accessed 04/04/2018,

Peña -Rosas JP, De-Regil LM, Dowswell T, Viteri FE (2015) Intermittent oral iron supplementation during pregnancy (Review). *Cochrane database of systematic reviews;* Issue 10.

Peña-Rosas JP, De-Regil LM, Garcia-Casal MN, Dowswell T. (2015). Daily oral iron supplementation during pregnancy (Review). *Cochrane database of systematic reviews;* Issue 7.

Physical Activity Guidelines Advisory Committee (2018). *Physical Activity Guidelines Advisory Committee Scientific Report.* Washington, DC: U.S. Department of Health and Human Services, 2018

Prentice AM, Jebb SA (2001) Beyond body mass index. Obesity Review; 2:141-7.

Rasmussen, Habicht (2010) Maternal Supplementation Differentially Affects the Mother and Newborn. *The Journal of Nutrition*. 140, 402-406.

Rasmussen KM and Yaktine A, (2009) Editors; Committee to reexamine IOM Pregnancy Weight Guidelines; Institute of Medicine; National Research Council. Weight Gain During Pregnancy Reexamining the Guidelines. Washington, The National Academies Press.

Ricci E, Chiaffarino F, Cipriani S, Malvezzi M, Parazzini F (2010) Diet in pregnancy and risk of small for gestational age birth: results from a retrospective case-control study in Italy. *Maternal and Child Nutrition*. 6(4):297-305.

Robillard, Dekker, Boukerrou, Le Moullec, Hulsey (2018) Relationship between prepregnancy maternal BMI and optimal weight gain in singleton pregnancies. *Heliyon*; 4, 5.

Rogozinska E, Marlin N, Jackson L, Rayanagoudar G, Ruifrok AE, Dodds J, et al. (2017) Effects of antenatal diet and physical activity on maternal and fetal outcomes: individual patient data meta-analysis and health economic evaluation. *Health Technol Assess*;21(41)

Rong, K, Yu K, Han X, Szeto, I, Qin, X, Wang, J, Ma, D. (2015). Pre-pregnancy BMI, gestational weight gain and postpartum weight retention: A meta-analysis of observational studies. *Public Health Nutrition*, 18(12), 2172-2182. doi:10.1017/S1368980014002523

Ronnberg AK, Ostlund I, Fadl H, Gottvall T, Nilsson K. (2015) Intervention during pregnancy to reduce excessive gestational weight gain—a randomised controlled trial. *BJOG.*;122(4):537-44. doi: 10.1111/1471-0528.13131

Royal College of Obstetricians and Gynaecologists (RCOG). (2015). *The Role of Bariatric Surgery in Improving Reproductive Health*, (Scientific Impact Paper No. 17). RCOG. Available at www.rcog.org.uk/en/guidelines-research-services/guidelines/sip17

Royal College of Physicians Ireland (RCPI) Policy Group on Physical Activity (2016) *A prescription for a wonder drug.* RCPI. Available at https://www.rcpi.ie/policy-and-advocacy/rcpi-policy-group-on-physical-activity/

Safefood (2017). The Folate Status of Pregnant Women in the Republic of Ireland; the current position December 2017. Available at ISBN: 978-1-905767-77-9, 978-1-905767-76-2

Santos et al (2019) Impact of maternal body mass index and gestational weight gain on pregnancy complications: an individual participant data meta-analysis of European, North American, and Australian cohorts BJOG. 2019; 126:984-995

Santos S, Eekhout I, Voerman E, et al (2018). Gestational weight gain charts for different body mass index groups for women in Europe, North America, and Oceania. *BMC Med.* 16(1):201. Published 2018 Nov 5. doi:10.1186/s12916-018-1189-1

Schmied VA, Duff M, Dahlen HG, Mills AE, Kolt GS (2011) 'Not waving but drowning': a study of the experiences and concerns of midwives and other health professionals caring for obese childbearing women. *Midwifery*. Aug;27(4):424-30.

Scientific Advisory Committee on Nutrition (SACN) (2010). *Iron and Health* Available at https://www.gov.uk/government/publications/sacn-iron-and-health-report. Accessed 04/04/2018.

Scientific Advisory Committee on Nutrition (SACN) (2016) Vitamin D and Health Available at: https://www.gov.uk/government/groups/scientific-advisory-committee-on-nutrition. Accessed May 2019.

Scott C, Andersen CT, Valdez N, Mardones F, Nohr EA, Poston L, et al. No global consensus: a cross-sectional survey of maternal weight policies. *BMC Pregnancy Childbirth* 2014;14:167.

Sebire NJ, Jolly M, Harris JP, Wadsworth J, Joffe M, Beard RW, Regan L, Robinson S (2012) Maternal obesity and pregnancy outcome: a study of 287,213 pregnancies in London. *International Journal Obesity Related Metabolism Disorders*. Aug;25(8):1175-82.

Siega-Riz AM, Viswanathan M, Moos MK, Deierlein A, Mumford S, Knaack J, Thieda P, Lux LJ, Lohr KN (2009) A systematic review of outcomes of maternal weight gain

according to the Institute of Medicine recommendations: birthweight, fetal growth, and postpartum weight retention. *American Journal of Obstetrics Gynecology*. Oct; 201(4):339.e1-14.

SLAN (Survey of Lifestyle Attitudes and Nutrition) (2008). *The National Health and Lifestyles Survey*. In Health. Promotion. Unit (Ed.). Dublin: Health Promotion Unit.

Stagnaro-Green A, Abalovich M, Alexander E, Azizi F, Mestman J, Negro R, Nixon A, Pearce EN, Soldin OP, Sullivan S, Wiersinga W; American Thyroid Association Taskforce on Thyroid Disease During Pregnancy and Postpartum. (2011). Guidelines of the American Thyroid Association for the diagnosis and management of thyroid disease during pregnancy and postpartum. *Thyroid* Oct;21(10):1081-125.

Symonds, Pope, Sharkey, Budge (2012) Adipose tissue and fetal programming. *Diabetologia*. 55:1597-1606.

Thangaratinam S, Rogozinska E, Jolly K, Glinkowski S, Duda W. (2012) Interventions to reduce or prevent obesity in pregnant women: a systematic review. *Health Technol Assess*;16(31)

Torloni, M. R., Betrán, A. P., Horta, B. L., Nakamura, M. U., Atallah, A. N., Moron, A. F. and Valente, O. (2009). Pre-pregnancy BMI and the risk of gestational diabetes: a systematic review of the literature. *Obes Rev.* 10(2):194-203.

Tounian P (2011) Programming towards childhood obesity. *Annals of Nutritional Metabolism.* 58, Suppl. 2, 30-41.

Trumpff C, De Schepper J, Tafforeau J, Van Oyen H, Vanderfaeillie J, Vandevijvere S. (2013). Mild iodine deficiency in pregnancy in Europe and its consequences for cognitive and psychomotor development of children: A review. *Journal of Trace Elements in Medicine and Biology* Feb 7. S0946-672X(13)00004-7.

Turcksin, R., Bel, S., Galjaard, S., and Devlieger, R. Maternal obesity and breastfeeding intention, initiation, intensity and duration: A systematic review. Matern Child Nutr. 2014; 10: 166–183

Turner MJ and Layte R (2013) Obesity levels in a national cohort of women 9 months after delivery. American Journal of Obstetrics and Gynecology 209(2): 124.

UK Chief Medical Officers (CMO) Recommendations (2017). Physical Activity in Pregnancy. Available at: https://www.health-ni.gov.uk/sites/default/files/publications/health/Physical%20activity%20for%20pregnant%20women%20infographic_FINAL.pdf

Villamor and Cnattingius (2016) Interpregnancy Weight Change and Risk of Preterm Delivery. *Obesity*. 24 (3): 727-734

Vivanti AP (2012). Origins for the estimations of water requirements in adults. *European Journal Of Clinical Nutrition* 66, 1282-1289

Walsh JM, Murphy DJ (2007) Weight and pregnancy. *British Medical Journal*; 335: 169.

Walsh JM, McGowan CA, Mahony R, Foley ME, McAuliffe FM. (2012) Low glycaemic index diet in pregnancy to prevent macrosomia (ROLO study): randomised control trial. *British Medical Journal*. Aug 30;345

Woo JG, Martin LJ. Does breastfeeding protect against childhood obesity? Moving beyond observational evidence. *Curr Obes Rep* (2015); 4: 207-216

World Health Organisation (WHO) expert committee technical report series 1995, *Physical status: the use and interpretation of and interpretation of anthropometry*. WHO technical report series 854. Geneva: World Health Organization

World Health Organisation (WHO). (2004). *Human energy requirements Report of a Joint FAO/WHO/UNU Expert Consultation*. Rome: World Health Organization World Health Organization (2015) *Guideline: Optimal serum and red blood cell folate concentrations in women of reproductive age for prevention of neural tube defects*. Geneva: World Health Organization

World Health Organisation (WHO). (2016). WHO recommendations on antenatal care for a positive pregnancy experience. Luxembourg: World Health Organisation.

Xu et al (2017) Inadequate weight gain in obese women and the risk of small for gestational age (SGA): a systematic review and meta-analysis. *J Matern Fetal Neonatal Med.*; 30(3):357-367

Zhou SJ, Gibson RA, Crowther CA, Makrides M.b. 2009. Should we lower the dose of iron when treating anaemia in pregnancy? A randomized dose-response trial. *European Journal of Clinical Nutrition*. Feb;63(2):183-90.

Appendix 1:

Useful Websites

HSE Nurture Programme

https://www.hse.ie/eng/health/child/nurture/intro.html

Food Safety Authority of Ireland website:

http://www.fsai.ie/

HSE Health Promotion website:

http://www.healthpromotion.ie/

Safe food website:

http://www.safefood.eu/

The Irish Nutrition and Dietetic Institute (INDI) website:

https://www.indi.ie/

Making Every Contact Count

https://www.hse.ie/eng/about/who/healthwellbeing/making-every-contact-count/about/html

HSE My Child

https://www2.hse.ie/my-child/

HSE Land

https://www.hseland.ie

Irish Food Allergy Network

http://ifan.ie/

Appendix 2:

Daily macronutrient requirements for pregnant and lactating women

	1 st Trimester	2 nd Trimester	3 rd Trimester	Lactation
Energy	Additional 0-85	Additional 260-	Additional 450 -	Additional
	kcal/ day	340 kcal/ day	500 kcal/ day	500 kcal/ day
Carbohydrate	175 g/ day	175 g/ day	175 g/ day	210 g/ day
	minimum	minimum	minimum	minimum
Protein	Additional 1g/ day	Additional 9g/day	Additional ≤28g/day or one extra serving from food pyramid	Additional 19g/day or one extra serving from food pyramid
Fat	Additional 700-	Additional 700-	Additional 700-	Additional 700-
	1400mg/week	1400mg/week	1400mg/week	1400mg/week
	DHA	DHA	DHA	DHA

Appendix 3:

Daily micronutrient requirements for pregnant and lactating women

	1 st Trimester	2 nd Trimester	3 rd Trimester	Lactation
Iron	16 milligrams	16 milligrams	16 milligrams	16 milligrams
Folic acid	400	400	400	400
	micrograms	micrograms	micrograms	micrograms
Calcium	1000	1000	1000	1000
	milligrams	milligrams	milligrams	milligrams
Vitamin D	10 micrograms	10 micrograms	10 micrograms	10 micrograms
	(400 IU)	(400 IU)	(400 IU)	(400 IU)
Iodine	200	200	200	200
	micrograms	micrograms	micrograms	micrograms