

Review Article





Screening for gestational diabetes mellitus

Abstract

Gestational diabetes is a carbonhydrate intolerance resulting in hyperglycaemia of variable severity with onset or first recognition during pregnancy. It is the most common metabolic disorder affecting pregnancy. Recognizing and treating diabetes or any degree of glucose intolerance in pregnancy results in lowering maternal and fetal complications. These patients present higher risk for excessive weight gain, preeclampsia, cesarean sections, a high risk of developing type 2 diabetes and cardiovascular disease in the future. Complications of infants are; higher risk for macrosomia, birth trauma, hypoglycemia, hypocalcemia, hyperbilirubinemia, respiratory distress syndrome, polycythemia, obesity and type 2 diabetes.

Gestational diabetes mellitus (GDM) screening should be performed at 24-28 gestational weeks and followed by definitive testing in women who are labeled as high-risk population. Two strategies are used to screen for gestational diabetes. In the two-step approach, the 50-g oral glucose challenge test (OGCT) is performed between 24 and 28 weeks of gestation in a nonfasting state. Alternatively, in the one-step approach, a 75-g glucose load is administered after measuring a fasting glucose level, and the plasma glucose levels are evaluated after 1 and 2 hours of the administration. There are the other methods for screening, including HbA1C and fasting blood glucose, but an exact approach for gestational diabetes screening has not yet been established.

Keywords: gestational diabetes mellitus, screening, one-step approach, two-step approach

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Abbreviations: GDM, gestational diabetes mellitus; OGCT, oral glucose challenge test; IADPSG, International association of the diabetes and pregnancy study groups; BMI, body mass index; NDDG, National diabetes data group; CC, carpenter and coustan criteria; WHO, World Health Organization; HAPO, hyperglycemia and adverse pregnancy outcomes; CDA, Canadian diabetes association; NICE, National Institute for health and clinical excellence; NIH, National Institutes of health; AACE, American association of clinical endocrinologists; ACOG, American college of obstetricians and gynecologists; OGTT, oral glucose tolerance test

Introduction

GDM is a carbonhydrate intolerance resulting in hyperglycaemia of variable severity with onset or first recognition during pregnancy.¹ GDM is an important public health problem. As older age at pregnancy become more common, GDM's prevalence is increasing as obesity and sedentary lifestyle.2-5 If GDM is unrecognized or left unregulated, there is an important risk of maternal, fetal and neonatal implications, including increased maternal weight gain, a higher rate of caesarean section, pre-eclampsia, shoulder dystocia and other birth traumas, macrosomia of newborn, and neonatal hypoglycemia, among others.6 All these perinatal complications can be prevented or reduced, if the hyperglicaemia is diagnosed early enough.^{7,8} During pregnancy, hyperplasia of the pancreatic β-cells occurs, leading to increased insulin secretion, and an early increase in insulin sensitivity is followed by progressive insulin resistance due to the placental secretion of diabetogenic hormones, including growth hormone, corticotropin-releasing hormone, placental lactogen, and progesterone. GDM screening is universal and testing for GDM is usually done between 24 and 28 weeks of gestation. 9,10 The approach to screening gestational diabetes in pregnant women will be reviewed here.

Prevalence

GDM is a common disease affecting approximately 6-7% of pregnancies, with variations in prevalence noted among different races, ethnicities, screening practises, testing methods, and diagnostic criteria. The prevalence has been increasing overtime, possibly due to increases in mean maternal age and weight. The prevalence of GDM is even higher among obese patients, ranging from 7-14%. 12,13 In 2010, International Association of the Diabetes and Pregnancy Study Groups (IADPSG) criteria proposed new screening and diagnostic criteria for gestational diabetes. The use of these criteria is predicted to increase the prevalence of gestational diabetes to 18%. 14

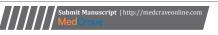
Risk factors

The risk factors for GDM include having a family history of diabetes mellitus type 2 (DM2), being a member of an ethnic group that is at increased risk for the development of DM2 (i.e. Hispanic, African, Native American, South Asian, East Asian, or Pacific Islands ancestry), previous GDM, a macrosomic infant (defined as 4kgs or more), a Body Mass Index (BMI) of $\geq 30 \text{kg/m}^2$, unexplained intrauterine foetal death, unexplained birth of a malformed infant, a maternal age of ≥ 35 years, glycosuria at the first prenatal visit, and a medical condition associated with the development diabetes, such as polycystic ovary syndrome, metabolic syndrome, or hypertension. Gestational diabetes is less likely to develop among women who are younger (<25 years of age), have a normal BMI ($<25 \text{kg/m}^2$), are white and non-Hispanic, have no first-degree relatives with diabetes, and have no history of previous glucose intolerance. If

Approaches to risk reduction

Physical activity

Nonpregnant women who engage in regular exercise have a lower risk of developing DM2 compared to nonpregnant women who





are sedentary. The effects of exercise on gestational diabetes have been explored with two types of studies: observational studies and randomised trials. The value of exercise was illustrated in 2011 with a meta-analysis that included seven pre-pregnancy studies and five early pregnancy studies. These studies have reported an association between increased physical activity and reduced risk of GDM.¹⁷ As for during gestation, a 2014 meta-analysis of six randomised trials revealed that initiating an exercise program during pregnancy did not significantly reduce the risk of developing GDM compared with routine care.¹⁸

Weight loss

In a cohort study, obese women who lost at least 4.5kgs between pregnancies decreased their risk of gestational diabetes relative to obese women whose weight reduced by less than 4.5kgs between pregnancies.¹⁹

Adverse outcomes during pregnancy

The risks of adverse outcomes increase as maternal fasting plasma glucose levels increase above 75mg/dl and as the one-hour and two-hour oral glucose tolerance test (OGTT) values increase. Associated foetal risks of GDM include macrosomia, shoulder dystocia, hypoglycemia, hyperbiluribinemia, respiratory distress syndrome, and childhood obesity. Maternal risks include preeclampsia, caesarean delivery, and increased risk of developing DM2 and cardiovascular disease later in life. Moreover, if maternal hyperglycaemia is present during the first trimester, it causes increasing the risk of miscarriage and congenital anomalies.20–28

History

In 1960s, O'Sullivan et al.²⁹ proposed diabetes improve outcomes, the first evidence that screening, diagnosis and treatment of hyperglycaemia in women not previously known to have.²⁹ These criteria were later modified by the National Diabetes Data Group (NDDG) in 1979 as well as by Carpenter and Coustan. The Carpenter and Coustan criteria (CC) were lower than the NDDG criteria and therefore resulted in a higher prevalence of GDM.^{29–31} In 1979-1980 international panels established the 2-h 75g OGTT as the diagnostic test for diabetes and glucose intolerance,³⁰ this recommendation was extended by the World Health Organization (WHO) to pregnant women.³² The 2-h 75g OGTT had been little examined during pregnancy, so the US(United States) National Diabetes Data Group (NDDG) continued to use the 3-h 100g OGTT.³⁰

In 2000, The NDDG recommendation had been followed by The American Diabetes Association (ADA). ADA recommended the use of the Carpenter and Coustan criteria for diagnosis of GDM. Over the last 3 decades these procedures and criteria were frequently adopted as a two-step procedure: a 50g 1-h challenge test and then a 100g 3-h OGTT for those positive at screening. In 2008, the results of the "Hyperglycemia and Adverse Pregnancy Outcomes (HAPO)" study were published. 25,505 pregnant women tested with a 2-h 75g OGTT for the classification of diagnostic criteria for GDM.²² This major observational study provided valuable information regarding the risks of adverse outcomes associated with various degrees of maternal glucose intolerance. Based on the results of this study, the International Association of Diabetes and Pregnancy Study Groups (IADPSG) proposed a new diagnostic criterion in 2010.14,22 The IADPSG Consensus Panel recommended the diagnostic criteria for GDM presented in Table 1.14 These cut points were also recommended by the ADA for a 2-h 75g OGTT in its 2011 position statement.³³

Most commonly used diagnostic criteria for GDM

The most commonly used guidelines for the diagnosis of GDM recommend the following diagnostic criteria (Table 2).^{1,14,34,35}

Table I Diagnostic thresholds for gestational diabetes mellitus

Dignostic criteria	S tep	Glucose load, g	Fasting glucose load mg/dl	lh	2h	3h	Abnormal values, n
NDDG	2-step	100	105	190	165	145	2
CC	2-step	100	95	180	155	140	2
ADA(2000- 2010)	2-step	75	95	180	155	-	2
CDA(2008)	2-step	75	95	191	160	-	2
IADPSG	I-step	75	92	180	153	-	1
WHO	I-step	75	110	-	140	-	1

ADA, American diabetes association; CC, carpenter–coustan; CDA, Canadian diabetes association; IADPSG, International association of the diabetes and pregnancy study groups; NDDG, National diabetes data group; WHO, World Health Organization

Table 2 Most commonly used guidelines for the diagnosis of GDM

Organisation	Fasting Plasma glucose, mmol/L	Glucose Challenge	I-h plasma glucose	2-h plasma glucose	3-h plasma glucose
WHO 1999*	≥7.0	75g OGTT	Not required	≥7.8	Not required
American Congress of Obstetricians and Gynecologists**	≥5.3	100g OGTT	≥10.0	≥8.6	≥7.8
Canadian Diabetes Association***	≥5.3	75g OGTT	≥10.6	≥8.9	Not required
IADPSG****	≥5.1	75g OGTT	≥10.0	≥8.5	Not required

*one value is sufficient for diagnosis

**two or more values are required for diagnosis

***two or more values required for diagnosis

Benefits of screening

Treating screen-detected GDM with dietary modifications, glucose monitoring, and insulin can significantly reduce the risk for preeclampsia, foetal macrosomia, and shoulder dystocia.³⁶

Screening for gestational diabetes mellitus

A screening test has to be sensitive, simple, cheap, defined with clear criteria, noninvasive, reproducible and available. Some authors are doubtfull about any health advantages in screening for GDM.³⁷ The Preventive Services Task Force (USPSTF), the United Kingdom National Health Service and the Canadian Task Force didn't report any sufficient high-level evidence about cost-effectiveness of routine GDM screening.³⁸ In the United States of America (USA), most obstetricians prefer universal screening with an initial 50-g glucose challenge test (GCT).³⁹⁻⁴¹ Very recently, the Canadian Diabetes Association (CDA) not only recommends the same sequential universal screening approach, but also offers an alternative with one-step 75-g oral glucose tolerance test (OGTT).⁴²

^{****}one value is sufficient for diagnosis

The National Institute for Health and Clinical Excellence (NICE) recommends selective screening in asymptomatic patients based on the risk factor assessment and does not advocate the use of the GCT, fasting glycemia, random blood glucose, or glucosuria.³⁹ The NICE is currently in the process of reviewing their guidelines. The Society of Obstetricians and Gynaecologists of Canada (SOGC) still recommends routine screening at 24-28 weeks' gestation with the 50-g GCT but is against screening for low-risk women with glucose testing.43

In 2013, the National Institutes of Health (NIH) published online the draft statement that the NIH panel did not find clear evidence to support adopting the one-step model, such as that proposed by IADPSG.⁴⁴ The American Association of Clinical Endocrinologists (AACE) and the ADA in 2011 recommend screening of all women without a pregestational diabetes at 24-28 weeks' gestation with a 75-g 2-h oGTT.40

Screening time

Screening is recommended after 24 weeks of gestation. Screening for GDM may occur earlier than 24 weeks of gestation in high-risk women (e.g. body mass index >30kg/m², prior history of gestational diabetes, or known impaired glucose metabolism), but there is little evidence of the benefits or harms of screening before 24 weeks of gestation. Performing oGTT before 24 weeks' gestation didn't recommended by the IADPSG Consensus Panel.45

In the absence of early screening or if early screening is negative, universal screening is performed at 24 to 28 weeks of gestation. A recent, randomised trial had shown the benefits of treating GDM and the possible reduction in healthcare costs that could be achieved with universal screening. Therefore, in 2011, the American College of Obstetricians and Gynecologists (ACOG) and ADA recommended universal screening because of the beneficial effects of screening, diagnosis and subsequent treatment.34,36 Before 28 weeks fetal growth acceleration has already began, so screening after 28 weeks isn't logical. For healty women without anamnestic risk factors GDM screening and diagnosis is performed between 24 and 28 weeks.^{7,38–40}

United States Preventive Services Task Force (USPTF) assessment for screening time

The USPSTF concludes with moderate certainty that screening for gestational diabetes after 24 weeks of gestation causes moderate reductions in maternal and foetal complications (specifically, the collective outcomes of preeclampsia, macrosomia, and shoulder dystocia). The USPSTF concludes that the evidence to support screening for gestational diabetes before 24 weeks of gestation is insufficient, and the balance of benefits and harms of screening cannot be determined.36

One-step and two-step approaches

Two strategies are used to screen for gestational diabetes. In the two-step approach, a 50-g oral glucose challenge test (OGCT) is performed between 24 and 28 weeks of gestation in a nonfasting state. If the screening threshold is met or exceeded (130mg/dL, 135mg/dl, or 140mg/dL), patients receive the oral glucose tolerance test (OGTT). During the OGTT, a fasting glucose level is obtained, followed by the administration of a 100-g glucose load, and glucose levels are evaluated after 1, 2 and 3 hours of the administration. A diagnosis of GDM is made when two or more glucose values fall at or above the specified glucose thresholds.36

These screening tests exhibited various sensitivities. For example, using a threshold of 140mg/dl in the 50-g GCT pooled an estimate of sensitivity ranging between 0.74 (95% CI 0.62-0.87) and 0.83 (95% CI 0.75-0.91). Lowering the threshold to 130mg/dL could increase the sensitivity of the test to 0.9.46 This false negative result might lead to false reassurance among the patients and the physicians. In contrast, the "one-step" approach could eliminate the problem of a false negative test and the potential drop-off after a positive screening test. 47 Alternatively, in the one-step approach, a 75-g glucose load is administered after the patient reaches a fasting glucose, and the plasma glucose levels are evaluated after 1 and 2 hours of the administration. Gestational diabetes mellitus is diagnosed if one glucose value falls at or above the specified glucose thresholds.

The one-step approach (75-g oral GTT) after release of the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) study findings of 2008, followed the 2010 recommendations of the IADPSG, which were endorsed by the ADA in 2011 are recommended. The most important reasoning for the shift from the 2-step to the 1-step method is the latter's ability to show GDM earlier, and thus doctors can treat the disease earlier, thereby decreasing the risks associated with the disease. In addition, proponents of the 1-step method affirm its ease of administration, its convenience for the patients, and its diagnostic accuracy.

The IADPSG recommended that a one-step 75-g OGTT be given to all pregnant women who do not have a diagnosis of overt diabetes. They also recommended that a single glucose value, rather than at least two abnormal values at or above diagnostic glucose thresholds on the OGTT, be accepted as sufficient evidence for a diagnosis of GDM. The diagnostic glucose thresholds recommended by the IADPSG were the maternal glucose values from the HAPO study that identified a 1.75-fold increase overall (adjusted odds ratio relative to the mean cohort glucose values) for gestational age, elevated C-peptides, high neonatal body fat, or a combination of these factors. In contrast, the 2-step approach to GDM screening has been shown to be more costeffective than the 1-step approach.48

Other screening tests for gestational diabetes mellitus

HgbA1c has limited efficacy in the diagnosis of GDM during the third trimester, demonstrating moderate sensitivity (85.7%) and poor specificity (61.1%). 40,49 The American Diabetes Association (ADA) and the International Association of Diabetes in Pregnancy Study Group (IADPSG) support its use during pregnancy. Four research articles evaluated the HbA1C thresholds of 5.0, 5.3, 5.5, 7.5 using different diagnostic criteria for gestational diabetes. 49-52

A threshold was lacking between the HgbA1C level and the probability of gestational diabetes across the four studies. However, O'Connor et al.53 investigated the normal values for HgbA1c in pregnancy in non-diabetic women and identified a normal reference interval of 4.3-5.4% in the first trimester.⁵³ Because of the high rates of false-positive and false-negative findings, serum HbA1c values cannot be recommended for screening GDM⁴⁰ USPSTF also supports this conclusion.38

Measuring fasting plasma glucose level has been suggested as an alternative to the OGCT. USPTF results show that a fasting plasma glucose test with a threshold of 85 mg/dL is similar to an OGCT in terms of sensitivity. 40 Herrera et al. 54 investigated the Importance of Fasting Blood Glucose (FBG) in Screening for Gestational Diabetes, and their data support the importance of incorporating a fasting

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blood glucose value into the initial screening or diagnostic testing for GDM.⁵⁴ The sensitivity of glucosuria for GDM screening is worse.⁵⁵

Various studies about screening methods

Nine studies provided data to estimate the sensitivity and specificity of OGCT using a cut-off point of 140mg/dL or greater. The joint estimates of sensitivity and specificity were 85% and 86%, respectively. Six studies reported results for a 50-g OGCT using a cutoff point of 130mg/dL or greater. Their joint estimates of sensitivity and specificity were 99% and 77%, respectively. A 50-g OGCT with a cut-off point of 130mg/dL had a higher sensitivity compared to a cut-off point of 140mg/dL; however, the specificity of the 130mg/ dL cut-off point was lower. 36,40 A study conducted in the United Arab Emirates using an HbA1C value of 5.5% or greater had a specificity of 21% and a sensitivity of 82%. Seven studies assessed the fasting plasma glucose test by using the Carpenter and Coustan's criteria, and confirmed the efficacy of GDM. Each study compared four fasting plasma glucose thresholds, and found that the sensitivity and specificity were 87% and 52% for 85mg/dL or greater, 77% and 76% for 90 mg/dL or greater, 76% and 92% for 92 mg/dL or greater, and 54% and 93% for 95mg/dL or greater, respectively.³⁶

Recommendations National Organisations

There is an agreement that universal GDM screening is costeffective. In 2013, the American Congress of Obstetricians and Gynecologists recommended the 2-step approach for screening GDM in all pregnant women. ACOG, SOGC and NICE recommend routine risk-factor-based screening. ADA recommended both the 1 and 2-step approaches with thresholds proposed by the IADPGS. ADA recommends screening all women with a 75-g 2-h oral glucose tolerance test (OGTT). In 2013, an independent panel supported by the NIH Consensus Development Program considered whether using the 75-g OGTT (1-step approach), as proposed by the International Association of Diabetes and Pregnancy Study Groups and supported by the American Diabetes Association, should be adopted instead of the 2-step approach. The panel released a statement that there is not enough evidence to support the 1-step approach over the 2-step. The American Academy of Family Physicians recommends screening for GDM in asymptomatic pregnant women after 24 weeks of gestation. It also concludes that the evidence is insufficient to assess the balance between benefits and harms that arise from screening for GDM in asymptomatic pregnant women before 24 weeks of gestation. The CAD, USPSTF, ADIPS, and DIP recommend that all asymptomatic women should be screened at 24-28 weeks' gestation. The Endocrine Society recommends universal screening for GDM using the 1-step approach. Australasian Diabetes in Pregnancy Society recommended the 1-step approach for screening GDM. 36,55

Conclusion

In all pregnant populations, screening for GDM, universal strategies are beneficial and better than the selective ones. Currently, the ACOG, the SOGC, and the NICE recommend routine risk-factorbased screening, and the CDA, the USPSTF, the ADA, the IADPSG, the ADIPS, and the ATLANTIC DIP recommend universal screening in asymptomatic pregnant women at 24-28 weeks' gestation. We have to concantrate on hyperglycemia because it is more important than trying to diagnose GDM itself. If it is possible, screening for hyperglycemia should be universally performed in every pregnant woman. The universal two-step screening is cost-effective. The best

strategy would be connecting the screening with diagnosing GDM in the same procedure with 75 g of glucose at 24-28 weeks' gestation. Despite the clear doubts the 50-g 1-h GCT has been stil used as the most popular screening method by a number of countries, including the USA.

In summary, there is still no exact approach to gestational diabetes screening. What is apparent is that screening and subsequent treatment is beneficial to short-term outcomes and possibly long-term outcomes. The most commonly evaluated screening test in the literature is the 2-step approach, and the most commonly used cut-off point is 140mg/ dL. Furthermore, the 2-step approach is more cost effective than the 1-step approach.

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None.

Conflicts of interest

The authors declare there is no conflict of interests.

References

- 1. World Health Organization. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: diagnosis and classification of diabetes mellitus. WHO/NCD/NCS/99. 2nd ed. Geneva: World Health Organization: 1999:1-66.
- 2. Guariguata L, Linnnenkamp U, Beagley J, et al. Global estimates of the prevalence of hyperglycaemia in pregnancy for 2013 for the IDF diabetes atlas. Diab Res Clin Pract. 2014;103(2):137-149.
- 3. National Institutes of Health consensus development conference statement: diagnosing gestational diabetes mellitus, March 4-6, 2013. Obstet Gynecol. 2013;122:358-369.
- 4. American Diabetes Association. Executive summary: standards of medical care in diabetes--2011. Diabetes Care. 2011;34(Suppl 1):S4-S10.
- 5. McCance DR. Pregnancy and diabetes. Best Pract Res Clin Endocrinol Metab. 2011;25(6):945-958.
- 6. Ferrara A. Increasing prevalence of gestational diabetes mellitus: a public health perspective. Diabetes Care 30. 2007; Suppl 2:S141-S146.
- 7. Hartling L, Dryden DM, Guthrie A, et al. Benefits and harms of treating gestational diabetes mellitus: a systematic review and meta-analysis for the US preventive services task force and the National Institutes of health office of medical applications of research. Ann Intern Med. 2013;159(2):123-129.
- 8. Power ML, Wilson EK, Hogan SO, et al. Patterns of preconception, prenatal and postnatal care for diabetic women by obstetrician-gynecologists. J Reprod Med. 2013;58(1-2):7-14.
- 9. Butte NF. Carbohydrate and lipid metabolism in pregnancy: normal compared with gestational diabetes mellitus. Am J Clin Nutr. 2000;71(5 Suppl):1256S-1261S.
- 10. Yamashita H, Shao J, Friedman JE. Physiologic and molecular alterations in carbohydrate metabolism during pregnancy and gestational diabetes mellitus. Clin Obstet Gynecol. 2000;43(1):87-98.
- 11. American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes Care. 2014;37(Suppl 1):S81-S90.
- 12. Kim SY, Saraiva C, Curtis M, et al. Fraction of gestational diabetes mellitus attributable to overweight and obesity by race/ethnicity, California, 2007-2009. Am J Public Health. 2013;103(10):e65-e72.
- 13. El-Chaar D, Finkelstein SA, Tu X, et al. The impact of increasing obesity class on obstetrical outcomes. J Obstet Gynaecol Can. 2013;35(3):224-

- Metzger BE, Gabbe SG, Persson B, et al. International association of diabetes and pregnancy study groups recommendations on the diagnosis and classification of hyperglycemia in pregnancy. *Diabetes Care*. 2010;33(3):676–682.
- Berger H, Crane J, Farine D, et al. Screening for gestational diabetes mellitus. J Obstet Gynaecol Can. 2002;24(11):894–912.
- Naylor CD, Sermer M, Chen E, et al. Selective screening for gestational diabetes mellitus. Toronto Trihospital Gestational Diabetes Project Investigators. N Engl J Med. 1997;337(22):1591–1596.
- Tobias DK, Zhang C, van Dam RM, et al. Physical activity before and during pregnancy and risk of gestational diabetes mellitus: a metaanalysis. *Diabetes Care*. 2011;34(1):223–229.
- Yin YN, Li XL, Tao TJ, et al. Physical activity during pregnancy and the risk of gestational diabetes mellitus: a systematic review and meta-analysis of randomised controlled trials. *Br J Sports Med*. 2014;48(4):290–295.
- Glazer NL, Hendrickson AF, Schellenbaum GD, et al. Weight change and the risk of gestational diabetes in obese women. *Epidemiology*. 2004;15(6):733–737.
- Dodd JM, Crowther CA, Antoniou G, et al. Screening for gestational diabetes: the effect of varying blood glucose definitions in the prediction of adverse maternal and infant health outcomes. *Aust N Z J Obstet Gynaecol*. 2007;47(4):307–312.
- 21. Temple R, Aldridge V, Greenwood R, et al. Association between outcome of pregnancy and glycaemic control in early pregnancy in type 1 diabetes: population based study. *BmMJ*. 2002;325(7375):1275–1276.
- 22. Metzger BE, Lowe LP, Dyer AR, et al. Hyperglycemia and adverse pregnancy outcomes. N Engl J Med. 2008;358(19):1991–2002.
- 23. Ferrara A, Weiss NS, Hedderson MM, et al. Pregnancy plasma glucose levels exceeding the American Diabetes Association thresholds, but below the National Diabetes Data Group thresholds for gestational diabetes mellitus, are related to the risk of neonatal macrosomia, hypoglycaemia and hyperbilirubinaemia. *Diabetologia*. 2007;50(2):298–306.
- Jensen DM, Korsholm L, Ovesen P, et al. Adverse pregnancy outcome in women with mild glucose intolerance: is there a clinically meaningful threshold value for glucose? *Acta Obstet Gynecol Scand*. 2008;87(1):59– 62
- Sermer M, Naylor CD, Gare DJ, et al. Impact of increasing carbohydrate intolerance on maternal-fetal outcomes in 3637 women without gestational diabetes. The Toronto Tri-Hospital Gestational Diabetes Project. Am J Obstet Gynecol. 1995;173(1):146–156.
- Sacks DA, Greenspoon JS, Abu-Fadil S, et al. Toward universal criteria for gestational diabetes: the 75-gram glucose tolerance test in pregnancy. Am J Obstet Gynecol. 1995;172(2 Pt 1):607–614.
- 27. Pettitt DJ, Knowler WC. Long-term effects of the intrauterine environment, birth weight, and breast-feeding in Pima Indians. *Diabetes Care*. 1998;21(Suppl 2):B138–B141.
- Hillier TA, Pedula KL, Schmidt MM, et al. Childhood obesity and metabolic imprinting: the ongoing effects of maternal hyperglycemia. *Diabetes Care*. 2007;30(9):2287–2292.
- O'Sullivan JB, Mahan CM. Criteria for the oral glucose tolerance test in pregnancy. *Diabetes*. 1964;13:278–285.
- National Diabetes Data Group. Classification and diagnosis of diabetes mellitus and other categories of glucose intolerance. *Diabetes*. 1979;28:1039–1057.
- 31. Carpenter MW, Coustan DR. Criteria for screening tests for gestational diabetes. *Am J Obstet Gynecol*. 1982;144(7):768–773.
- 32. WHO Expert Committee on Diabetes Mellitus. Second Report. World Health Organ Tech Rep Ser. 1980;646:1–80.

- 33. American Diabetes Association. Diagnosis and classification of diabetes mellitus. *Diabetes Care*. 2011;34(Suppl 1):S62–S69.
- 34. Committee Opinion No. 504: Screening and diagnosis of gestational diabetes mellitus. *Obstet Gynecol*. 2011;118(3):751–753.
- Canadian Diabetes Association 2008 Clinical Practice Guidelines for the Prevention and Management of Diabetes in Canada. *Canadian Journal of Diabetes*. 2008;32(Suppl 1):1–215
- Moyer VA. Screening for gestational diabetes mellitus: U.S. Preventive Services Task Force recommendation statement. *Ann Intern Med*. 2014;160(6):414–420.
- Simmons D, Moses RG. Gestational diabetes mellitus: to screen or not to screen? Is this really still a question? *Diabetes Care*. 2013;36(10):2877– 2878
- US Preventive Services Task Force. Screening for gestational diabetes mellitus: US Preventive Services Task Force recommendation statement. Ann Intern Med. 2008;148(10):759–765.
- Avalos GE, Owens LA, Dunne F. Applying current screening tools for gestational diabetes mellitus to a European population: is it time for change? *Diabetes Care*. 2013;36(10):3040–3044.
- Donovan L, Hartling L, Muise M, et al. Screening tests for gestational diabetes: a systematic review for the U.S. Preventive Services Task Force. Ann Intern Med. 2013;159(2):115–122.
- Van Leeuwen M, Louwerse MD, Opmeer BC, et al. Glucose challenge test for detecting gestational diabetes mellitus: a systematic review. *BJOG*. 2012;119(4):393–401.
- 42. Canadian Diabetes Association Clinical Practice Guidelines Expert Committee, Thompson D, Berger H, Feig D, et al. Diabetes and pregnancy. *Can J Diabetes*. 2013;37(Suppl 1):S168–S183.
- Berger H, Sermer M, Farine D. Should the SOGC guidelines on screening for gestational diabetes mellitus be changed once again? *J Obstet Gynaecol Can.* 2006;28(6):536–539.
- Vandorsten JP, Dodson WC, Espeland MA, et al. NIH Consensus Development Conference: diagnosing gestational diabetes mellitus. NIH Consens State Sci Statements. 2013;29(1):1–31.
- 45. Sacks DA, Hadden DR, Maresh M, et al. Frequency of gestational diabetes mellitus at collaborating centers based on IADPSG Consensus Panel recommended criteria: the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) Study. *Diabetes Care*. 2012;35(3):526–528.
- 46. American Diabetes Association. Gestational diabetes mellitus. *Diabetes Care*. 2004;27(Suppl 1):S88–S90.
- 47. Huynh J, Ratnaike S, Bartalotta C, et al. Challenging the glucose challenge test. *Aust N Z J Obstet Gynaecol*. 2011;51(1):22–25.
- 48. Coustan DR. Diagnosis of gestational diabetes. *Scand J Clin Lab Invest Suppl*. 2014;244:27–33.
- Rajput R, Yogesh Yadav, Rajput M, et al. Utility of HbA1c for diagnosis of gestational diabetes mellitus. *Diabetes Res Clin Pract*. 2012;98(1):104– 107.
- Uncu G, Ozan H, Cengiz C. The comparison of 50 grams glucose challenge test, HbA1c and fructosamine levels in diagnosis of gestational diabetes mellitus. Clin Exp Obstet Gynecol. 1995;22(3):230–234.
- Agarwal MM, Hughes PF, Punnose J, et al. Thomas. Gestational diabetes screening of a multiethnic, high-risk population using glycated proteins. *Diabetes Res Clin Pract*. 2001;51(1):67–73.
- Agarwal MM, Dhatt GS, Punnose J, et al. Gestational diabetes: a reappraisal of HBA1c as a screening test. Acta Obstet Gynecol Scand. 84(12):1159–1163.

- 53. O'Connor C, O'Shea PM, Owens LA, et al. Trimester-specific reference intervals for haemoglobin A1c (HbA1c) in pregnancy. Clin Chem Lab Med. 2012;50(5):905-909.
- 54. Herrera K, Brustman L, Foroutan J, et al. The importance of fasting blood
- glucose in screening for gestational diabetes. J Matern Fetal Neonatal Med. 2015;28(7):825-828.
- 55. Petrovic O. How should we screen for gestational diabetes? Curr Opin Obstet Gynecol. 2014;26(2):54-60.