

# Postpartum Screening Following GDM: How Well Are We Doing?

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**Abstract** Once diagnosed with gestational diabetes mellitus (GDM), a woman has a sevenfold increased risk of developing type 2 diabetes relative to women who do not have diabetes during pregnancy. In addition, up to one third of women with GDM have overt diabetes, impaired fasting glucose, or impaired glucose tolerance identified during postpartum glucose screening completed within 6 to 12 weeks. Therefore, the American Diabetes Association, the World Health Organization, and the American College of Obstetricians and Gynecologists currently recommend postpartum glucose screening following GDM. However, despite this recommendation, in many settings the majority of women with GDM fail to return for postpartum glucose testing. Studies conducted to date have not comprehensively examined the health care system, the physician, or the patient determinants of successful screening. These studies are required to help develop standard clinical procedures that enable and encourage all women to return for postpartum glucose screening following GDM.

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## Introduction

Gestational diabetes mellitus (GDM), defined as glucose intolerance with onset or first recognition during pregnancy, is estimated to affect 2% to 10% of the pregnancies in the United States, with estimates being higher for racial and ethnic minority groups than for non-Hispanic white individuals [1]. Once diagnosed with GDM, a woman has a high chance of developing type 2 diabetes following delivery, with studies estimating cumulative incidence of 15% to 50% [2–9] in the decades following delivery, and a recent meta-analysis reporting a sevenfold increased risk of developing type 2 diabetes in women who had a pregnancy with GDM relative to women who did not have diabetes during pregnancy [4••]. In addition, recent studies indicate that women with GDM not only have increased risk of developing type 2 diabetes, but have increased cardiometabolic and cardiovascular disease risk [10–12].

Although the majority of women with GDM have normal glucose regulation postpartum, up to one third of women will have overt diabetes, impaired fasting glucose, or impaired glucose tolerance identified during postpartum screening completed within 6 to 12 weeks [9, 13–15]. However, although women are often motivated when pregnant to improve their health and often successfully control their diabetes during pregnancy, in many settings the majority of women with GDM fail to return for postpartum glucose testing despite clinical guidelines recommending such testing [6, 13, 16–24].

The current article outlines clinical guidelines recommending postpartum glucose screening following GDM,

reviews current estimates of postpartum glucose screening following GDM including the type of screening used, and discusses factors associated with receiving postpartum glucose screening.

### Postpartum and Long-Term Glucose Screening Guidelines

The definitions of diabetes and impaired glucose regulation have changed during the past 15 years, and may change again with the recent emphasis the American Diabetes Association (ADA) has placed on the value of hemoglobin A<sub>1c</sub> for screening. However, the present review considers the current diagnostic criteria for diabetes, including the 1985 and 1999 World Health Organization (WHO) criteria [25] that require a 2-hour 75-g oral glucose tolerance test (OGTT) and the 1997 ADA criteria [26] that are based on fasting plasma glucose, but also recognize a casual or 2-hour 75-g OGTT glucose level  $\geq 200$  mg/dL as diagnostic of diabetes. Additionally, the ADA in 1997 and the WHO in 1999 focused on impaired glucose tolerance (defined as a 2-hour 75-g OGTT level of 140–199 mg/dL) as a marker of abnormal glucose regulation and increased risk of subsequent type 2 diabetes [25, 26], while more recently the ADA has considered individuals with impaired glucose tolerance or impaired fasting glucose (defined as a fasting plasma glucose 100–125 mg/dL) as being “prediabetic” [27, 28].

Because a fasting plasma glucose test is quicker and easier to perform than a 2-hour 75-g OGTT, but does not identify individuals with impaired glucose tolerance and is therefore less sensitive, there is controversy as to which test should be used to screen individuals at high risk for developing diabetes. Erring on the comprehensive side, the Fifth International Workshop on GDM recommended that all women with GDM undergo a 2-hour 75-g OGTT 6 to 12 weeks postpartum [29]. The American College of Obstetricians and Gynecologists (ACOG) Committee on Obstetric Practice and the ADA recommend that all women with GDM be screened at 6 to 12 weeks postpartum with a fasting plasma glucose or a 2-hour 75-g OGTT [30, 31]. Interestingly, neither ACOG nor ADA clearly states whether the 2-hour 75-g OGTT is preferred over the fasting plasma glucose test. ACOG acknowledges that the 2-hour 75-g OGTT is more sensitive and ADA merely recognizes the 2-hour 75-g OGTT as a valid diagnostic test. The WHO guidelines recommend a 2-hour 75-g OGTT 6 weeks postpartum [25].

Whereas the ACOG does not make a statement about long-term follow-up of women with GDM, the ADA recommends that high-risk individuals, including women with previous GDM, be screened for diabetes every 3 years [31].

### Estimates of Postpartum Glucose Screening

A literature search was conducted to identify recently published articles (ie, published since January 1, 2004) specific to postpartum diabetes screening following GDM. A decision was made not to focus on articles in which the primary objective was to determine rates of postpartum type 2 diabetes or abnormal glucose regulation because implementation of these studies likely impacted postpartum screening rates.

During the past 5 years, a number of studies using medical record review were published regarding the prevalence of postpartum glucose screening following GDM [18–24]. These studies are summarized in Table 1. All are retrospective, with the exception of a single study completed using a GDM registry established by Kaiser Permanente Medical Care Program in Northern California [20]. The prevalence of postpartum glucose screening with a fasting plasma glucose or a 2-hour 75-g OGTT in these studies ranges from 23% to 58% [18–24]. The two studies reporting the highest prevalence of postpartum glucose screening were the Kaiser Permanente Medical Care Program in Northern California and the Kaiser Permanente Northwest Health Maintenance Organization in Oregon and Washington State [19, 20]. They were the only studies to report postpartum glucose screening of over 50%. In the single study conducted in Canada, where there is a publically funded universal health care system, the prevalence of postpartum glucose testing was 48% [22].

In addition to the studies completed based on medical record review, three surveys were conducted that collected information on postpartum glucose screening rates following GDM (Table 2) [32–34]. In a survey of ACOG Fellows and Junior Fellows, 74% of physicians who provide prenatal care reported providing postpartum glucose screening following GDM and 58% reported performing a fasting plasma glucose or a 2-hour 75-g OGTT postpartum following GDM [33]. In a second survey of North Carolina in-state practitioners who provided prenatal care, 21%, 43%, 20%, and 16% reported that they always, usually, sometimes, or “rarely or never” screen for abnormal glucose levels following GDM [32]. A total of 54% reported that they used a 2-hour 75-g OGTT when they screened. Interestingly, the survey of North Carolina in-state practitioners was the only study that reported rates of routine screening after the postpartum period; 35% reported that they screen every year, 14% reported that they screen every 3 years, and 47% reported no routine screening [32]. Finally, in a postpartum survey of women with GDM conducted in Australia, 73% of women reported they had received some type of glucose screening at any point postpartum; and 61% reported that they had received some type of screening within the 6- to 8-week window

**Table 1** Studies published within the past 5 years reporting postpartum diabetes screening rates following GDM based on medical record review

Study	Time frame	Population	Follow-up	n	Screened postpartum (%; method)	Factors associated with increased screening
Smirnakis et al. [24]	Retrospective; 2000–2001	Massachusetts General Hospital and Baystate Medical Center, MA, required postpartum follow-up	6 wk–4.5 y	197	67%; any type of screen 37%; FPG or OGTT	Higher GDM diagnostic glucose levels
Kim et al. [21]	Retrospective; 1997–2002	University of Michigan Hospital, MI	>6 wk	533	38%; any type of screen 23%; FPG or OGTT	Being married; saw an endocrinologist after delivery; more provider contacts after delivery
Russell et al. [23]	Retrospective; 2001–2004	Diabetes Pregnancy Clinic, Women and Infants' Hospital of Rhode Island, RI	5–9 wk	344	45%; FPG or OGTT	Being Hispanic compared to white; attending 6-wk postpartum visit
Almario et al. [18]	Retrospective; 2004–2006	Thomas Jefferson University Hospital, PA, required postpartum follow-up	5–12 wk	90	20%; screen ordered 33%; referral/screen ordered	High-risk pregnancy office; higher GDM diagnostic glucose levels; insulin use during pregnancy
Dietz et al. [19]	Retrospective; 2004–2006 <sup>a</sup>	Kaiser Permanente Northwest Health Maintenance Organization, OR and WA	6–12 wk	461	79%; screen ordered	Practice site where care received was associated with clinician order; Asian or Hispanic compared to white; attending 6-wk postpartum visit
Kwong et al. [22]	Retrospective; 1999–2006	Diabetes Pregnancy Clinic, Alberta, Canada	6 wk–6 mo	909	48%; FPG or OGTT	Lower parity; insulin use during pregnancy
Ferrara et al. [20]	GDM registry; 1995–2006	Kaiser Permanente Medical Care Program in Northern California, CA	6 wk–1 y	14,448	1995, 20%: FPG or OGTT 2006, 56%: FPG or OGTT	Older age, Asian or Hispanic compared to white; higher education, earlier GDM diagnosis, use of diabetes medication during pregnancy; more provider contacts after delivery; being nonobese; lower parity

FPG fasting plasma glucose; GDM gestational diabetes mellitus; OGTT oral glucose tolerance test

<sup>a</sup>The time period for the study was actually 1999–2006, but we chose to focus on the period from 2004 to 2006

**Table 2** Surveys, RCTs, and prospective studies published within the past 5 years reporting postpartum diabetes screening rates following GDM

Study	Design/time frame	Population	Follow-up	n	Screened postpartum (%; method)	Factors associated with increased screening
Gabbe et al. [33]	Survey; 2003	ACOG Fellows and Junior Fellows; doctors response rate, 41%	NA	441	74%; any type of screen 58%; FPG or OGTT	Doctors <40 years of age were more likely to report routinely performing postpartum glucose screening
Baker et al. [32] <sup>a</sup>	Survey; 2005–2006	North Carolina in-state practitioners who provided prenatal care; doctors response rate, 40%	NA	327	21%; always screen 43%; usually screen 20%; sometimes screen 16%; rarely or never screen 54%; use OGTT to screen when they screen	Factors reported to impact screening: lost to follow-up (50%), patient inconvenience (32%), inconsistent guidelines (27%), patient refusal (18%), patient cost (17%), and reimbursement (16%)
Morrison et al. [34]	Survey; 2003–2005	National Diabetes Service Scheme database, Australia; GDM patients response rate, 36%	6–8 wk	1372	73%; any type of screen 61%; any type of screen within 6–8 wk 27%; OGTT within 6–8 wk	Lower age, written postnatal information, individualized risk reduction advice, receiving care from an endocrinologist among those with less education and receiving care from a diabetes educator among those who saw an obstetrician
Clark et al. [36]	RCT: 2 × 2 factorial; postal reminders to patients and physicians 2002–2005	High-risk pregnancy clinic, Ontario, Canada; GDM patients follow-up	6 wk–1 y	234	OGTT—test used 60.5%; patient and doctor 55.3%; patient only 51.6%; doctor only 14.3%; no reminder 57%; FPG or OGTT	Postal reminders to patients and/or doctors increased postpartum OGTT screening
Hunt and Conway [14]	Prospective; 2001–2003	Diabetes pregnancy clinic; UT Health Science Center at San Antonio, TX; GDM patients	6–12 wk	707	Less severe GDM, lower GDM diagnostic glucose levels, did not require insulin during pregnancy, and no prior history of GDM	Older age and insulin use during pregnancy
Ogonowski and Miazgowski [35]	Prospective; 2005–2007	Diabetes pregnancy clinic, Szczecin, Poland; GDM patients	5–9 wk	855	37%; OGTT	

ACOG American College of Obstetricians and Gynecologists; FPG fasting plasma glucose; GDM gestational diabetes mellitus; NA not available; OGTT oral glucose tolerance test; RCT randomized controlled trial

<sup>a</sup>This study also reported rates of routine screening after the postpartum period: 35% reported that they screen every year, 14% reported no routine screening

recommended in Australia; however, only 27% reported that they had received a 2-hour 75-g OGTT within the 6- to 8-week postpartum window [34].

### Factors Associated with Increased Screening

Successful postpartum glucose screening following GDM is dependent on the health care system, the physician, and the patient. The health care system is responsible for establishing clinical practice recommendations and facilitating their implementation, the physician is responsible for following clinical practice recommendations and ordering the recommended tests, and the patient is responsible for completing the test. Studies conducted to date have not comprehensively examined health care system, physician, or patient determinants of successful screening. However, targeting even a single area may significantly increase postpartum glucose screening following GDM.

In the studies based on medical record review (Table 1), factors consistently associated with increased screening across at least two studies could be grouped into three categories: GDM severity, health care/provider, and patient characteristics. GDM characteristics associated with increased screening included higher GDM diagnostic glucose levels and insulin use to treat diabetes during pregnancy [18, 20, 22, 24]. Health care/provider characteristics associated with increased screening included completion of a 6-week postpartum visit, type of practice site where care was received, having more provider contacts after delivery, and in a single study seeing an endocrinologist after delivery [18–21]. Patient characteristics associated with increased screening included being Asian or Hispanic compared with non-Hispanic white, and lower parity [19, 20, 22].

Factors identified to impact postpartum glucose screening through the two physician surveys were physician age (ie, physicians <40 years of age were more likely to report routine screening), loss to follow-up, patient inconvenience, inconsistent guidelines, patient refusal, patient cost, and reimbursement (Table 2) [32, 33]. Factors identified to impact postpartum glucose screening through the postpartum survey of GDM patients included patient age (ie, younger age was associated with increased screening), written postnatal information, individualized risk reduction advice, receiving care from an endocrinologist among less educated women, and receiving care from a diabetes educator among those who saw an obstetrician (Table 2) [34].

Factors identified to impact postpartum glucose screening in two prospective studies that examined characteristics of women who did and did not return for postpartum glucose screening included patient age, insulin use during pregnancy, GDM diagnostic glucose levels, and prior history of GDM (Table 2) [14, 35]. Interestingly, in contrast

to the postpartum survey of women in Australia [34], in the prospective study conducted in Poland [35], older patient age was associated with increased glucose screening. Also interesting, in the study conducted in San Antonio, Texas [14], in which a case manager was used to increase postpartum glucose screening and all patients were encouraged to complete screening, factors associated with increased severity of GDM (ie, higher diagnostic glucose levels, prior history of GDM, and insulin use during pregnancy) were associated with failure to return for postpartum glucose screening.

Finally, in a single randomized clinical trial designed to increase postpartum glucose screening following GDM, postal reminders to patients and their physicians were successfully used as the intervention to improve postpartum glucose screening (Table 2) [36]. A two-by-two factorial design was used: 61% of women completed a 2-hour 75-g OGTT when both the patient and physician received the postal reminder, 55% when only the patient received the postal reminder, 52% when only the physician received the reminder, and only 14% when neither the patient nor the physician received the reminder.

### Conclusions

The ADA, the WHO, and the ACOG all currently recommend postpartum glucose screening following GDM [25, 28–30••]. However, there is disagreement over whether an OGTT is required or if obtaining fasting plasma glucose levels is sufficient, as well as what cutpoint should be used to diagnose impaired fasting glucose. These inconsistencies across clinical practice recommendations likely contribute to the observed low postpartum glucose screening rates. However, even in studies aimed at improving postpartum glucose screening rates and in integrated health management organizations with a focus on improving postpartum glucose screening rates, completion rates hovered around 60% [14, 19, 20].

Recently, several clinical trials have indicated that through diet and exercise, or with the aid of a pharmacologic agent, it was possible to lower the incidence or delay the onset of diabetes among individuals at high risk for the disease [37–41]. Thus, the optimum way to reduce the risk associated with diabetes may be preventing diabetes itself, either by altering lifestyle, or by using pharmacologic agents. Women with impaired fasting glucose or impaired glucose tolerance in the early postpartum period following GDM are at very high risk for developing type 2 diabetes and the burden of diabetes is especially high in these women because of their young age. Reducing the incidence of type 2 diabetes following GDM also reduces the inherent risks to future offspring of exposure to a diabetic intrauterine environment.

Studies conducted to date have not comprehensively examined the health care system, the physician, and the patient determinants of successful screening. For instance, although studies have identified loss to follow-up and failure to return for 6-week postpartum visit as risk factors for failure to complete glucose screening postpartum, studies have not examined what factors facilitate or impede a patient's ability to return for postpartum medical care. Studies have also not examined to what extent changing from obstetrical to primary care postpartum may impact postpartum glucose screening. Finally, studies have not been completed that focus on long-term screening for diabetes following GDM. These studies are required to help develop standard clinical procedures that enable and encourage all women to return for postpartum glucose screening.

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