Accuracy of glycosuria, random blood glucose and risk factors as selective screening tools for gestational diabetes mellitus in comparison with universal diagnosing

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ABSTRACT

Objective Despite the short-term and long-term health implications of gestational diabetes mellitus (GDM), opinions are divided on selective vis-à-vis universal screening. We validated the accuracy of screening tests for GDM.

Research design and methods Pregnant women (n=491) were recruited to this prospective, blind comparison with a gold standard study. We did selective screening between 13 and 20 weeks using reagent-strip glycosuria, random capillary blood glucose (RBG) and the presence of ≥ 1 risk factor(s). Between 20 and 34 weeks, we did universal screening following the 'one-step' approach using glycated hemoglobin (HbA1c), fasting venous plasma glucose (FPG), and the 1-hour and the 'gold standard' 2-hour oral glucose tolerance test (OGTT). Tests accuracy was estimated following the WHO and the National Institute for Health and Care Excellence (NICE) diagnostic criteria. Overall test performance was determined from the area under the receiver operating characteristic curve (AUC).

Results GDM prevalence per 2-hour OGTT was 9.0% for the WHO criteria and 14.3% for the NICE criteria. Selective screening using glycosuria, RBG and risk factors missed 97.4%, 87.2% and 45.7% of cases, respectively. FPG threshold ≥ 5.1 mmol/L had the highest clinically relevant sensitivity (68%) and specificity (81%), but FPG threshold ≥ 5.6 mmol/L had higher positive predictive value. Although sensitivity of 1-hour OGTT was 39.5%, it had the highest accuracy and diagnostic OR. Regarding test performance, 1-hour OGTT and FPG were very good (AUC >0.8), RBG was poor (AUC ≈ 0.60), whereas HbA1c was invaluable (AUC <0.5).

Conclusions Selective screening using glycosuria and random blood glucose is unnecessary due to its low sensitivity. Fasting glucose ≥ 5.1 mmol/L could be applicable for screening at the population level. Where 2-hour OGTT is not available, FPG ≥ 5.6 mmol/L, complemented by the presence of risk factors, could be useful in making therapeutic decision.

INTRODUCTION

Gestational diabetes mellitus (GDM), a form of pregnancy-specific glucose intolerance, has

Significance of this study

What is already known about this subject?

- ▶ Despite recent updates on the diagnostic criteria for gestational diabetes mellitus (GDM), opinions are divided on the need for selective screening using risk factors, glycosuria and random blood glucose.

What are the new findings?

- ▶ The presence of risk factors was more sensitive compared with glycosuria, random blood glucose and glycated hemoglobin, which were highly insensitive and diagnostically poor, and hence missed majority of the cases.
- ▶ Fasting plasma glucose threshold ≥ 5.1 mmol/L was more sensitive than values ≥ 5.6 mmol/L.

How might these results change the focus of research or clinical practice?

- ▶ Findings reaffirm the rising prevalence of GDM and highlight the need to integrate fasting blood glucose monitoring into all GDM screening/diagnostic procedures.
- ▶ When deciding to treat, the presence of risk factors could be complementary to fasting plasma glucose, particularly where oral glucose tolerance test is not available.

been identified as the most prevalent metabolic disorder in pregnancy. The condition is associated with adverse maternal, fetal and neonatal outcomes,^{1 2} and increases the risk of type II diabetes in mother-child dyads in later years.³ In 2013, the global prevalence of hyperglycemia in pregnancy in women aged 20–49 years was 17%. Over 90% of cases occurred in low-income and middle-income countries.⁴ In Africa, the prevalence of hyperglycemia in pregnancy has considerably increased from negligible levels to almost 30% in some settings within the past four decades.^{5–7} The rising prevalence of

hyperglycemia has been attributed to epidemiologic transition associated with increasing type II diabetes mellitus, unhealthy dietary habits, obesity, physical inactivity, and advanced maternal age at childbirth.^{3,8} However, consideration of gross variations in screening and diagnostic procedures and the reference thresholds used is necessary when interpreting these trends.

Screening here denotes the scheduled measurement of blood glucose in all pregnant women whether asymptomatic or symptomatic, and whether 'at risk' or not, followed by diagnostic testing in screen-positive clients. However, the specific screening procedures and outcomes that should provide indication to conduct a diagnostic test are still controversial.^{9–11} Despite recent establishment of diagnostic criteria by various health regulating bodies, opinions are divided on selective versus universal screening. Selective screening, sometimes known as routine screening, is where only pregnant women identified from screening procedures to be at high risk for GDM are requested to perform diagnostic test to evaluate their actual glycemic status. Conversely, universal screening is where all pregnant women irrespective of their glycemic status perform a recommended diagnostic test to assess their likelihood for developing GDM.

In line with evidence from the hyperglycemia and adverse pregnancy outcome study,¹ and the recommendations from the International Association of Diabetes and Pregnancy Study Group (IADPSG),¹² the WHO updated its clinical guidelines for detecting hyperglycemia in pregnancy in 2013.¹⁰ Similarly, the American Diabetes Association (ADA) updated its guidelines in 2015 in accordance with the recommendations of the IADPSG.¹¹ The WHO recommends one-step diagnosing of GDM using fasting plasma glucose (FPG) ≥ 5.1 mmol/L or 75 g oral glucose intake, followed by 1-hour postprandial glucose ≥ 10.0 mmol/L. However, 2-hour oral glucose tolerance test (OGTT) ≥ 8.5 mmol/L performed at any time during pregnancy but preferably between 24 and 28 weeks is highly recommended.

On the other hand, the National Institute for Health and Care Excellence (NICE) considers 2-hour 75 g OGTT performed between 24 and 28 weeks as the 'gold standard'. Compared with the current WHO/IADPSG/ADA guidelines, NICE's diagnostic threshold for 2-hour OGTT is 0.7 mmol/L lower (≥ 7.8 mmol/L), whereas the FPG threshold is 0.5 mmol/L higher (≥ 5.6 mmol/L).⁹ Meanwhile, the 1999 WHO diagnostic criteria for FPG was ≥ 7.0 mmol/L, whereas the 2-hour OGTT was ≥ 7.8 mmol/L, same as the current NICE guideline. Even though this paper focuses on GDM, it is worth noting that the WHO/IADPSG recommend random plasma glucose or 2-hour OGTT ≥ 11.1 mmol/L at any time during pregnancy as suggestive of clinical (pre-existing) diabetes.^{10,13} In this paper, the WHO, IADPSG and ADA diagnostic recommendations are collectively denoted as the WHO criteria.

Screening protocol for GDM in Ghana comprises reagent-strip urinalysis to detect glycosuria, tested at

every antenatal care (ANC) visit. Values of 1+ and 2+ on two occasions or 3+ and 4+ on one occasion warrant an OGTT.¹⁴ Although fasting blood glucose is recommended at the first ANC booking, and at 28–32 weeks, no diagnostic cut-offs are provided in the standard treatment guidelines. Adherence to the screening and diagnostic protocol is discretionary by the clinician and dependent on the level of healthcare provision. Making diagnostic decision based on maternal risk factor assessment is often the norm. However, there is no clear guideline on the type or number of risk factors that should necessitate diagnostic testing. Previous macrosomia delivery (birth weight >4.0 kg), a cardinal adverse neonatal outcome of GDM, has identified 3%¹⁵ of pregnant women to be at risk of developing the condition. Meanwhile, the reported GDM prevalence according to 2-hour OGTT using the WHO reference threshold of ≥ 8.5 mmol/L is 9% in the largest referral facility in Ghana.¹⁶

Urinalysis through reagent-strip glycosuria is not without challenges. During pregnancy, the renal threshold for glomerular glucose reabsorption is reduced, leading to increased glycosuria at some point in about 50% of all pregnancies.¹⁷ However, hyperglycemia without detectable glycosuria is not unlikely.¹⁸ Although dipstick glycosuria test is cheap and convenient, we suspect that its usage as the main routine screening test is resulting in missed (false negative) opportunities for diagnosis and management. Therefore, this diagnostic accuracy study was conducted to compare the prevalence of GDM in primary and secondary healthcare settings in Ghana using the current WHO and NICE diagnostic criteria. We validated the performance of dipstick glycosuria, random capillary whole blood glucose (RBG) and the presence of risk factors as screening tests for accurately detecting GDM. Outcomes were validated against fasting venous plasma glucose (FPG), glycated hemoglobin (HbA1c), and postprandial 1-hour and 2-hour OGTT considered as the diagnostic tests. The effectiveness of universal and selective screening approaches was assessed. Finally, cut-off points that optimized sensitivity and specificity were suggested from the coordinates of the receiver operating characteristic (ROC) curve.

METHODS

Design

This diagnostic accuracy study, reported according to the Standards for Reporting Diagnostic Accuracy guidelines,¹⁹ was designed as a prospective, blind comparison with a gold standard study. This design compares the diagnostic accuracy of different tests in the same individual. This study is part of a prospective cohort study aimed at assessing the effects of hyperglycemia in pregnancy on maternal, fetal and neonatal outcomes. The index screening tests assessed were dipstick glycosuria test, RBG and maternal risk factors. One-hour OGTT and HbA1c were the index diagnostic tests evaluated.

Performance of each test within the same individual was validated against 2-hour OGTT and FPG.

Study sites

The study was conducted at the antenatal clinics of one primary-level and four secondary-level public health-care facilities in the Volta Region of Ghana. Forty-seven percent (516 461) of the 1 098 854 women in the region are in their reproductive ages (15–49 years).²⁰ ANC and delivery services are provided predominantly in secondary-level hospitals in Ghana. Hence although the study was performed predominantly in secondary care settings, it was designed to replicate results that would have been achieved at all healthcare levels. The largest of the secondary facilities was the Volta Regional Hospital, a 240-bed capacity regional referral hospital in the process of upgrade to a university teaching hospital. The other three facilities were district hospitals. These included the Hohoe Municipal Hospital with a bed capacity of 178, Margret Marquart Catholic Hospital with a bed capacity of 152 and the Ho Municipal Hospital with a bed capacity of 130. The Jasikan District Hospital, a 45-bed capacity facility, represented maternal healthcare provision at the primary level.

Sample size and eligibility

Cochran's formula,²¹ which accounts for finite population corrections, was used to determine the minimum sample size required. We considered 9.3% reported prevalence of GDM in Ghana¹⁶ and a total target population of 516 461²⁰ women in their reproductive age in the region. A 95% confidence level corresponding to 1.96 Z-score and a 5% permitted error margin generated a sample size of 130. The primary and secondary facilities were each treated as separate clusters. To account for variability of health service provision at each level, the effect of clustering was adjusted for, ensuring maximized statistical accuracy of GDM estimate that is nationally representative.

Aided by the G*Power software (V.3.1.9.2),²² a design effect of 3.2 was determined based on a two-tailed t-test statistic with an alpha error probability of 0.05, 1- β error probability of 0.95 and an intraclass correlation of 0.16 derived from 0.03 coefficient of determination. The sample size was multiplied by the design effect, resulting in an effective sample size of 416. An a priori statistical power analysis yielded 91%, an indication that the sample size was adequately powered. Participant recruitment from the primary and secondary facilities was done proportional to size in the ratio of 1:6.

Irrespective of the presence or absence of GDM risk factors, any pregnant woman aged 15 years and above identified in the first trimester accessing ANC in any of the five study facilities between June and November 2016 and who provided written consent was eligible to participate. However, pregnant women with prior diabetes, identified either from the medical records or self-affirmed, were excluded. Every consecutive pregnant

woman who met the inclusion criteria was recruited until the required sample size was attained.

Test procedures and diagnostic cut-offs

Maternal risks factors were assessed from the sociodemographic data collected, as well as medical and obstetric history. Women with three or more known risk factors (maternal age above 35 years, body mass index above 29.9 kg/m², parity above three children, history of diabetes and previous macrosomia delivery) were regarded to be at high risk for GDM.

The one-step strategy recommended by the IADPSG for diagnosing GDM was followed.¹² Glycosuria, RBG and risk factors were considered as screening tests, whereas FPG, HbA1c, and 1-hour OGTT were diagnostic. These tests were all evaluated with reference to the 2-hour OGTT.

Glycosuria urinalysis was tested at every ANC visit as a qualitative urine dipstick test using the 'Urit 2V^{PG}' urine reagent strips to detect the presence of glucose in about 10 mL of urine. A color change corresponding to trace test result or above (1+ to 4+) at any one point during pregnancy was considered to be screen-positive for GDM. Between 13 and 20 weeks, participants were screened for GDM through RBG. About 0.8 μ L capillary blood was drawn from the middle fingertip prick and measured on the point-of-care testing device called 'On Call Plus Blood Glucose Meter'. Random blood glucose >7.0 mmol/L was considered to be screen-positive for GDM.⁹

All participants were scheduled for diagnostic testing between 20 and 28 gestational weeks. Participants who failed to turn up were rescheduled between 30 and 34 weeks. The evening prior to the test, participants were called and reminded to fast overnight for 10–12 hours. On arrival at the laboratory on the morning of the test, they were made to rest for about 15 min before the tests were conducted. Three milliliters of preprandial venous blood were drawn from the antecubital fossa. One milliliter of the blood was used to measure the FPG on a fully automated 'Selectra ProM' clinical chemistry analyzer operating on kinetic enzymatic peroxidase-antiperoxidase principle. In accordance with the WHO¹⁰ and NICE⁹ diagnostic criteria, FPGs ≥ 5.1 mmol/L and ≥ 5.6 mmol/L were considered to be GDM diagnostic-positive. HbA1c was checked irrespective of clients' fasting status using 2 mL of venous blood. It was analyzed on an automated HA-8160 ADAMS analyzer operating on the ion exchange High Performance Liquid Chromatography (HPLC) principle. The results were aligned to the Diabetes Control and Complications Trial method. HbA1c results $\geq 6.5\%$ (7.8 mmol/L) were suggestive of GDM.⁹

After drawing the preprandial blood, participants were supervised to drink within 3 min 75 g anhydrous glucose dissolved in 300 mL of water at room temperature. Postprandial venous blood was collected at 1-hour and 2-hour intervals and plasma glucose levels checked following similar biochemical procedure as for the FPG. As per the WHO diagnostic criteria, 1-hour OGTT ≥ 10.0 mmol/L and 2-hour OGTT ≥ 8.5 mmol/L¹⁰ were indicative of

GDM. Also, as per the NICE diagnostic criteria, 2-hour OGTT ≥ 7.8 mmol/L was indicative of GDM.¹¹ Test quality was ensured by running controls after every 20th test and/or on the morning of each test day as the case might be. Where thresholds for two out of these four tests were obtained (RBG ≥ 11.1 mmol/L, 2-hour OGTT ≥ 11.1 mmol/L, HbA1c $\geq 6.5\%$ or FPG ≥ 5.6 mmol/L), it was considered to be a case of pre-existing diabetes first detected in pregnancy.

Statistical analysis

The data were entered into EpiData software and exported into STATA software (V.14.0) for analysis. All missing inputs for FPG and 2-hour OGTT were automatically excluded from the analysis. Dichotomous outcomes indicating positive or negative disease state were assigned to each screening and diagnostic test. Differences among primary and secondary facility users were tested using independent sample t-test for continuous scale glucose values and presented as mean with SD. Categorical data were tested using χ^2 test. The results were presented with the corresponding 95% CI of the level. P value < 0.05 was statistically significant.

The numbers of true positive, true negative, false positive, and false negative were estimated with reference to the diagnostic criteria for 2-hour OGTT based on the

WHO (≥ 8.5 mmol/L) and NICE (≥ 7.8 mmol/L) guidelines. Similar measures were calculated for FPG ≥ 5.1 mmol/L and ≥ 5.6 mmol/L in accordance with the WHO and NICE diagnostic criteria. Paired diagnostic accuracy was estimated using standard formulas for disease measurement to evaluate the discriminative, predictive and diagnostic properties of each test. Discriminatory and predictive abilities were determined by test sensitivity, specificity, positive and negative predictive values, diagnostic accuracy, and diagnostic OR (DOR). Higher values indicated better diagnostic properties.

To determine overall test performance, the ROC curve was obtained from the raw glucose values. The closer the curve was to the left-hand upper border of the ROC space, the more accurate the test. Test performance was evaluated based on the area under the curve (AUC). Rating ranged from excellent (AUC=0.9–1.0), very good (0.8 to < 0.9), good (0.70 to < 0.80), sufficient (0.60 to < 0.70), poor (0.50 to < 0.60) to invaluable (AUC < 0.5).^{23 24} The p value for each variable tested the null hypothesis that the true area under the ROC curve (AUC) is equal to 0.5 (no effect). This corresponds to the diagonal where the true positive rate equals the false positive rate. A p value < 0.05 indicates that we can reject the null hypothesis, meaning the test is clinically relevant.²³ The lower

Figure 1 Flow of participants through the study and the proportions who tested positive and negative to each test. HbA1c, glycated hemoglobin; Neg, negative; OGTT, oral glucosetolerance test; Pos, positive.

the p value, the better the test performance. Reference cut-offs of each test that provided analogous and clinically useful sensitivity and specificity for the study population were estimated from the coordinates of the ROC curve.

RESULTS

Overall, 491 pregnant women aged 15–54 years were screened for GDM between 12 and 20 gestational weeks, while 435 performed the diagnostic tests between 20 and 34 weeks. The 56 fallout was a result of spontaneous abortions, inability to fast overnight and difficulty tolerating the OGTT. The flow of participants through the study, proportions who performed each screening and diagnostic test, and the main reasons for withdrawal from each stage of the screening/diagnosing are shown in figure 1.

Presented in figure 2 is a comparison of the GDM positive proportions based on screening and diagnostic outcomes. Per the WHO diagnostic criteria, the prevalence of GDM based on the 'gold standard' 2-hour OGTT was 9.0% (n=39, 95% CI 6.5 to 12.0). The prevalence based on FPG was 23.8% (n=106, 95% CI 19.8 to 27.9), whereas based on 1-hour OGTT was 4.5% (n=20, 95% CI 2.7 to 6.7). On the other hand, the NICE diagnostic criteria for

2-hour OGTT yielded a GDM prevalence of 14.3% (n=63, 95% CI 11.0 to 17.5), whereas the diagnostic cut-off for FPG yielded a prevalence of 10.8% (n=48, 95% CI 7.9 to 13.7). Meanwhile, 29.1% (n=130, 95% CI 24.7 to 32.6) had at least one risk factor for GDM. If the WHO 1999 criteria for FPG ≥ 7.0 mmol/L was in use, the prevalence of GDM would have been 2.7% (n=12, 95% CI 1.1 to 4.3). Cases of pre-existing diabetes first detected in pregnancy were 2.0% (n=10, 95% CI 1.0 to 3.5).

Presented in the box and whisker plots in figure 3 are the interquartile values for RBG, HbA1c, FPG, and 1-hour and 2-hour OGTT among participants accessing ANC in primary (n=78) and secondary (n=357) facilities. The solid line represents the WHO diagnostic criteria, whereas the dotted lines represent the NICE diagnostic criteria. In comparison with participants who received ANC from the secondary facilities, participants who received ANC from the primary healthcare facilities had significantly higher mean FPG (4.44 ± 0.98 vs 5.12 ± 0.85 mmol/L), and 1-hour (6.70 ± 1.88 vs 7.20 ± 1.83 mmol/L) and 2-hour (6.24 ± 1.71 vs 6.74 ± 1.72 mmol/L) postprandial glucose. However, no statistical differences were found for RBG and HbA1c values.

Overall, the mean 1-hour postprandial glucose (7.2 ± 1.8 mmol/L) was 0.5 mmol/L higher than the

Figure 2 Percentage of participants who tested positive to the screening and diagnostic tests. HbA1c, glycated hemoglobin; NICE, National Institute for Health and Care Excellence; OGTT, oral glucose tolerance test.

Figure 3 Mean and interquartile glucose values for (A) random blood glucose, (B) fasting plasma glucose, (C) HbA1c, (D) 1-hour OGTT, (E) 2-hour OGTT and (F) triglycerides among participants receiving antenatal care in the primary and secondary facilities. HbA1c, glycated hemoglobin; OGTT, oral glucose tolerance test; solid line, WHO criteria; dotted line, the NICE criteria.

2-hour postprandial glucose (6.7 ± 1.7 mmol/L). Also, the mean random blood glucose (5.4 ± 1.1 mmol/L) was 0.3 mmol/L higher than the FPG (5.1 ± 0.9 mmol/L). Breakdown of the diagnostic-positive, screen-positive, and risk factor-positive proportions among primary and secondary facility users is presented in [table 1](#).

Even though GDM tended to be more prevalent among primary healthcare users, no significant differences existed except for the prevalence based on FPG and HbA1c. Regarding risk factors, first trimester obesity was found to be significantly higher among participants attending the secondary-level facilities.

Table 1 Prevalence of GDM and risk factors among users of primary and secondary healthcare facilities

Test and reference cut-off values	Mean±SD	% (n)	95% CI*	Facility (%)		P values†
				Primary	Secondary	
Diagnostic tests						
2-hour OGTT ≥8.5 mmol/L‡	6.7±1.7	9.0 (39)	6.5 to 12.0	10.3	8.7	0.663
2-hour OGTT ≥7.8 mmol/L§	6.7±1.7	14.3 (63)	11.0 to 17.5	20.5	12.9	0.106
1-hour OGTT ≥10.0 mmol/L‡	7.2±1.8	4.5 (20)	2.7 to 6.7	6.4	4.1	0.368
FPG ≥5.1 mmol/L‡	5.1±0.9	23.8 (106)	19.8 to 27.9	48.7	18.7	<0.0001
FPG ≥5.6 mmol/L§	5.1±0.9	10.8 (48)	7.9 to 13.7	23.4	8.2	<0.0001
HbA1c ≥6.5%§	5.5±0.9	6.3 (30)	4.2 to 8.5	11.7	5.2	0.040
Screening tests						
RBG >7.0 mmol/L§	5.4±1.1	4.7 (23)	2.9 to 6.7	6.2	4.4	0.562
Glycosuria ≥trace	–	3.1 (15)	1.6 to 4.7	1.2	3.4	0.484
Risk factors						
Maternal age >35 years	28.4±6.3	13.8 (62)	10.5 to 17.2	13.3	13.9	1.000
Gravidity above 5	2.7±1.5	5.4 (24)	3.4 to 7.7	6.7	5.1	0.577
Parity above 3 children	1.5±1.3	7.1 (30)	4.8 to 9.7	5.8	7.4	0.801
Body weight >90 kg¶	63.0±13.1	3.4 (15)	1.8 to 5.3	1.4	3.8	0.483
Height <150 cm¶	161.8±8.2	4.2 (19)	2.4 to 6.2	5.9	3.9	0.509
BMI ≥29.9 kg/m²¶	24.7±4.9	12.0 (52)	9.0 to 15.0	4.3	13.5	0.027
MUAC >30 cm¶	28.3±3.9	23.3 (103)	19.5 to 27.6	16.9	24.5	0.220
Family history of diabetes	–	6.4 (29)	4.2 to 8.6	8.0	6.1	0.604
Systolic BP >140 mm Hg	106.6±12.4	2.0 (9)	0.7 to 3.3	1.4	2.1	1.000
Diastolic BP >90 mm Hg	66.2±9.4	1.8 (8)	0.7 to 3.1	1.4	1.9	1.000
Proteinuria ≥1+	–	4.2 (19)	2.4 to 6.0	1.3	4.8	0.338
Triglycerides >2.25 mmol/L	3.0±1.7	59.5 (290)	55.0 to 63.9	35.0	64.4	<0.0001

*95% CI for the positive proportions (% (n)).

†Differences among primary and secondary facility estimated by χ^2 test.

‡Same tests but diagnostic criteria based on WHO guidelines.

§Same tests but diagnostic criteria based on the NICE guidelines.

¶Maternal weight, height and MUAC measured in the first trimester.

BMI, body mass index; BP, blood pressure; FPG, fasting plasma glucose; HbA1c, glycated hemoglobin; MUAC, mid-upper arm circumference; NICE, National Institute for Health and Care Excellence; OGTT, oral glucose tolerance test; RBG, random blood glucose.

Table 2 shows the diagnostic accuracy of the test instruments with reference to the 'gold standard' 2-hour OGTT and FPG estimated based on the WHO and NICE cut-offs. Generally, the screening tests had low sensitivity and positive predictive value (PPV), whereas their specificity and negative predictive value (NPV) were relatively high. Using the WHO diagnostic criteria for 2-hour OGTT as reference, FPG ≥5.1 mmol/L had the highest sensitivity (68%) and NPV (96%). However, 1-hour OGTT had the highest specificity, PPV (75%) and DOR. Glycosuria, HbA1c and RBG were the least sensitive (<15%) and hence yielded a clinically irrelevant specificity >95%. Although the presence of at least one risk factor was 54% sensitive, its accuracy was the least among the test instruments (≈70%). Similar pattern was observed when the NICE diagnostic criteria for 2-hour OGTT (≥7.8 mmol/L) was used as the reference. But here the test instruments had relatively lower sensitivity and DOR.

Using the WHO diagnostic criteria for FPG (≥5.1 mmol/L) as the reference yielded generally lower diagnostic measures. Here, 2-hour OGTT ≥7.8 mmol/L had the highest sensitivity and NPV. This was followed by risk factors with a sensitivity of 32%. However, the 1-hour OGTT ≥10.0 mmol/L had the highest PPV and DOR. When the NICE diagnostic criteria for FPG (≥5.6 mmol/L) was used as the reference, sensitivity of the test instruments was slightly higher.

If selective screening based on trace/above glycosuria was used, only 3.2% would have performed a diagnostic test for GDM. Out of this total, only 2.6% would be positive if the test of choice was 2-hour OGTT. Should glycosuria cut-off be changed to 1+ and above, screen-positive would have further declined to less than 1% and none of the participants would have needed a diagnostic test. If selective diagnostic testing based on RBG >7.0 mmol/L

Table 2 Diagnostic measure of test instruments using 2-hour OGTT (≥ 8.5 mmol/L or ≥ 7.8 mmol/L) and fasting plasma glucose (≥ 5.1 mmol/L or ≥ 5.6 mmol/L) as diagnostic criteria

Reference test	Diagnostic measure	TP	FN	FP	TN	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)	DOR
2-hour OGTT ≥ 8.5 mmol/L*	Risk factors	19	16	97	267	54.3	73.4	16.4	94.4	71.7	3.3
	Glycosuria	1	38	13	383	2.6	96.7	7.1	91.0	88.3	0.8
	RBG	5	33	16	381	13.2	96.0	23.8	92.0	88.7	3.6
	HbA1c	1	37	26	364	2.6	93.3	3.7	90.8	85.3	0.4
	FPG*	26	12	77	318	68.4	80.5	25.2	96.4	79.5	9.0
	FPG†	16	21	29	367	43.2	92.7	35.6	94.6	88.5	9.6
	1-hour OGTT	15	23	5	391	39.5	98.7	75.0	94.4	93.6	51.0
2-hour OGTT ≥ 7.8 mmol/L†	Risk factors	23	35	93	248	39.7	72.7	19.8	87.6	67.9	1.8
	Glycosuria	3	59	11	362	4.8	97.1	21.4	86.0	83.9	1.7
	RBG	6	56	15	358	9.7	96.0	28.6	86.5	83.7	2.6
	HbA1c	2	59	25	342	3.3	93.2	7.4	85.3	80.4	0.5
	FPG*	38	22	65	308	63.3	82.6	36.9	90.9	79.9	8.2
	FPG†	21	39	24	349	35.0	93.6	46.7	90.0	85.5	7.8
	1-hour OGTT	17	44	3	370	27.8	99.2	85.0	89.4	89.2	47.7
FPG ≥ 5.1 mmol/L*	Risk factors	31	67	83	217	31.6	72.3	27.2	76.4	62.3	1.2
	Glycosuria	5	101	9	330	4.7	97.3	35.7	76.6	75.3	1.8
	RBG	11	95	10	329	10.4	97.1	52.4	77.6	72.2	3.8
	HbA1c	10	93	18	317	9.7	94.6	35.7	77.3	70.3	1.9
	1-hour OGTT	17	89	3	334	16.0	99.1	85.0	79.0	75.2	21.3
	2-hour OGTT*	27	76	12	318	26.2	96.4	69.2	80.7	79.7	9.4
	2-hour OGTT†	38	65	22	308	36.9	93.3	63.3	82.6	79.9	8.2
FPG ≥ 5.6 mmol/L†	Risk factors	18	27	98	265	40.0	73.0	15.5	90.8	69.4	1.8
	Glycosuria	4	44	10	387	8.3	97.5	28.6	89.8	87.9	3.5
	RBG	8	40	13	384	16.7	96.7	38.1	90.6	88.1	5.9
	HbA1c	6	41	22	369	12.8	94.4	21.4	90.0	85.6	2.5
	1-hour OGTT	13	35	7	388	27.1	98.2	65.0	91.7	90.5	20.6
	2-hour OGTT*	17	28	21	367	37.8	94.6	44.7	92.9	88.7	10.6
	2-hour OGTT†	21	24	39	349	46.7	89.9	35.0	93.6	85.5	7.8

Screen-positive is ≥ 1 risk factors, trace/above reagent-strip glycosuria and random capillary blood glucose > 7.0 mmol/L. Diagnostic-positive is HbA1c $\geq 6.5\%$ †, fasting venous plasma glucose ≥ 5.1 mmol/L* and ≥ 5.6 mmol/L†, and 1-hour OGTT ≥ 10.0 mmol/L* and 2-hour OGTT ≥ 8.5 mmol/L* and ≥ 7.8 mmol/L† per the *WHO and †NICE criteria. Higher sensitivity, specificity, positive and negative predictive values, accuracy and DOR are indicative of better test performance. DOR values range from 0 to infinity.

DOR, diagnostic OR; FP, false positive; FPG, fasting plasma glucose; FN, false negative; HbA1c, glycated hemoglobin; NICE, National Institute for Health and Care Excellence; NPV, negative predictive value; OGTT, oral glucose tolerance test; PPV, positive predictive value; RBG, random blood glucose; TN, true negative; TP, true positive.

were used, only 4.8% would have performed a diagnostic test for GDM, out of which 12.8% would have tested positive. If selective diagnostic testing based on the presence of at least one risk factor was used to inform diagnostic testing, 29.1% would have performed a diagnostic test, out of which 53.4% would be positive.

Figure 4 presents the ROC curves showing the AUC as indication of test performance. Interpretation of the ROC curves is presented in table 3. In figure 4A and B, the reference test was 2-hour OGTT based on the WHO (≥ 8.5 mmol/L) and NICE (≥ 7.8 mmol/L) guidelines. Here, we observed that the 1-hour OGTT and FPG were very good tests as the AUC was between 0.88 and 0.86. Random blood glucose was a poor test for detecting

GDM (AUC ≈ 0.6), whereas HbA1c was invaluable because the AUC was < 0.5 and the CI of the AUC was statistically insignificant ($p = 0.686$). In figure 4C and D, FPG ≥ 5.1 mmol/L and ≥ 5.6 mmol/L were used as the reference test. Here, 1-hour and 2-hour OGTT were found to be 'good' test because the AUC was between 0.76 and 0.78. Although AUC values for RBG and HbA1c were slightly higher when compared with figure 4A and B, the test rating was similar as when the 2-hour OGTT was used as the reference.

Finally, from the coordinates of the ROC curve where 2-hour OGTT ≥ 8.5 mmol/L was the reference, we estimated the diagnostic thresholds for the study population that provided optimized and clinically relevant sensitivity

Figure 4 Receiver operating characteristic analyses showing area under the curves with reference to 2-hour OGTT (A,B) and fasting plasma glucose (C,D). HbA1c, glycated hemoglobin; OGTT, oral glucosetolerance test.

and specificity. A random blood glucose threshold of 5.2 mmol/L corresponded to a sensitivity of 60% and specificity of 54%. Similar sensitivity and specificity (62% vs 41%) were obtained for the HbA1c threshold of 5.2%. Increasing the sensitivity of both RBG and HbA1c to 70% corresponded to thresholds of 5.0 mmol/L and 5.0%, respectively, but the specificity of both was not clinically relevant (38% vs 25%). Regarding fasting glucose, 90% sensitivity and 72% specificity corresponded to 4.8 mmol/L. At the same threshold, the NICE criteria (2-hour OGTT ≥ 7.8 mmol/L) yielded 80% sensitivity and

74% specificity. In the case of 1-hour OGTT, 7.8 mmol/L threshold corresponded to 82% sensitivity and 81% specificity.

DISCUSSION

Through this diagnostic accuracy study, we found that GDM prevalence from 2-hour OGTT (9.0%) was similar to the 9.3% recorded in a tertiary facility in Ghana in 2015.¹⁶ Glycemic values of primary healthcare users tended to be higher. Although the presence of at least one

Table 3 Test performance estimated from area under the ROC curves using 2-hour OGTT and fasting plasma glucose values as reference test according to the WHO and NICE diagnostic criteria

Reference criteria	Test assessed	AUC‡	95% CI	P values§
2-hour OGTT ≥8.5 mmol/L*	1-hour OGTT	0.88	0.81 to 0.94	<0.0001
	Fasting glucose	0.86	0.80 to 0.91	<0.0001
	Random glucose	0.60	0.51 to 0.69	0.035
	HbA1c	0.48	0.40 to 0.57	0.686
2-hour OGTT ≥7.8 mmol/L†	1-hour OGTT	0.85	0.80 to 0.91	<0.0001
	Fasting glucose	0.83	0.78 to 0.88	<0.0001
	Random glucose	0.62	0.55 to 0.69	0.002
	HbA1c	0.48	0.40 to 0.55	0.567
Fasting glucose ≥5.1 mmol/L*	1-hour OGTT	0.77	0.71 to 0.82	<0.0001
	2-hour OGTT	0.78	0.72 to 0.83	<0.0001
	Random glucose	0.63	0.57 to 0.69	<0.0001
	HbA1c	0.51	0.44 to 0.57	0.880
Fasting glucose ≥5.6 mmol/L†	1-hour OGTT	0.76	0.68 to 0.84	<0.0001
	2-hour OGTT	0.78	0.70 to 0.86	<0.0001
	Random glucose	0.65	0.56 to 0.74	0.001
	HbA1c	0.51	0.41 to 0.60	0.867

*WHO and †NICE diagnostic criteria. ‡Area under the curve under the null hypothesis=0.5. Test rated excellent for AUC 0.9–1.0, very good for AUC 0.8–0.9, good for AUC 0.70–0.80, sufficient for AUC 0.60–0.70, poor for AUC 0.50–0.60 and invaluable for AUC <0.5 if $p < 0.05$.^{23 24}

§The lower the p value, the higher the clinical relevance of the test.

HbA1c, glycated hemoglobin; NICE, National Institute for Health and Care Excellence; OGTT, oral glucose tolerance test; ROC, receiver operating characteristic.

risk factor for GDM had the least specificity, it was a more sensitive screening tool for the condition compared with glycosuria and random blood glucose testing. Regarding diagnostic tests, FPG had more clinically useful sensitivity and specificity, while 1-hour OGTT had the highest specificity, DOR and AUC.

Short-term and long-term adverse effects of gestational hyperglycemia are well documented.^{1–3} Prevention and management start with timely and accurate diagnosing. Apart from maternal risk evaluation through medical/obstetric history taking and physical assessment, reagent-strip glycosuria and RBG are used for screening purposes particularly in resource-constraint settings and serve as the basis for diagnostic testing.⁹ It is therefore crucial that these screening tests are clinically useful and adequately sensitive (at least 70%) to rule out the condition. Highly specific tests are known to be vital for disease confirmation after a positive screening.²⁵

Like in most low-income and middle-income health-care settings, reagent-strip glycosuria testing is a routine ANC practice in Ghana, and diagnostic decision is made based on its outcome. NICE recommends glycosuria of ≥1+ on more than one occasion or ≥2+ on one occasion as indication to conduct the 'one-step' OGTT.⁹ In Ghana, glycosuria values of 1+/2+ on two occasions or 3+/4+ on one occasion are indications for OGTT.¹⁴ In this study, women with glycosuria of 1+ and above at any one point during pregnancy were less than 1%. The test sensitivity was <5% and only 3% of clients had positive glycosuria.

With the prevalence of GDM based on the 2-hour OGTT and fasting blood glucose being much higher, it is not surprising that NICE recommends a re-evaluation of use of glycosuria in pregnancy,⁹ while others have discredited its use.^{17 26}

It has been established that during pregnancy, renal glucose threshold is highly variable with a reduction in glycemic thresholds needed for the diagnosis of gestational diabetes,²⁶ while there is possibility of asymptomatic hyperglycemia without detectable glycosuria.¹⁸ Often, glycosuria and proteinuria are checked simultaneously using a single test strip and provide vital basis for detecting pre-eclampsia as well. Hence, the test is cost-effective despite its associated limitations. In primary healthcare settings where glycosuria testing is predominant, clients with trace results should be closely monitored and possibly booked for diagnostic testing if trace result is obtained on more than one occasion. But it is important that the diagnostic decision for GDM does not rely on the presence of glycosuria because it is highly insensitive and has a very low DOR.

Maternal risk factors are increasingly becoming a significant predictor for GDM.^{3 8} We found that a third of the pregnant women had at least one risk factor for GDM. In a similar study in South Africa, almost half (45.8%) had at least one risk factor.⁷ However, 26.0% of those 'at risk' in the South African study developed GDM compared with 54.3% in our study. We observed that the use of risk factors is a better screening tool compared

with glycosuria and RBG because whereas risk factors would miss approximately half of the true positive rate, glycosuria and RBG would miss approximately 90%. This supports the ongoing call for universal screening where all pregnant women are tested using a recommended diagnostic criterion instead of selective screening based on screen-positive results.^{7 10 12}

HbA1c test had the highest number of false negative and least DOR, and the test performance was diagnostically invaluable. As pregnancy progresses, HbA1c levels decrease²⁷ as a result of factors such as increased red cell turnover, hemoglobin variations and iron deficiency anemia,²⁸ which is a major public health problem in Ghana. This might explain why our mean HbA1c was 5.5%, similar to the 5.3% reported in South African.⁷ HbA1c provides information about 3-month glycemic control. Values >6.5% at first prenatal visit are suggestive of overt diabetes.⁹ However, in deprived populations where over 90% of pregnant women are unaware of their prepregnancy glycemic status and half make their first ANC visit after the first trimester, it is difficult to delineate pre-existing hyperglycemia from pregnancy-induced diabetes. Considering the limited availability of HbA1c testing, its high cost (approximately \$15 equivalent in Ghana) and the lack of correlation between HbA1c and average blood glucose due to gestational metabolic alterations,²⁹ the use of HbA1c as a routine GDM detection tool in resource-constraint settings is unnecessary at the population level.

Although the diagnostic accuracy of random blood glucose was poor, its accuracy was better than glycosuria and HbA1c. Interestingly, the mean RBG was only 0.3 mmol/L-point higher than the mean FPG. In our study setting, care seekers typically attend healthcare facilities in fasting state in anticipation of service providers unexpectedly requesting a biochemical test that might require overnight fasting. Also, long waiting times and high clientele turnout often compel clients to skip breakfast or eat meals typically smaller than usual in order to arrive early at the hospital and thus minimize delays. Meanwhile, the effect of pregnancy-related physiologic changes altering dietary patterns and preferences cannot be ruled out.²⁷ No wonder the highest sensitivity and specificity for RBG were at the 5.2 mmol/L threshold. RBG using capillary finger prick checked on a glucose meter is cheap and can be done by low-cadre health workers with minimum training, thereby making it the test of choice in primary healthcare care settings. However, none of the health regulating bodies have established a diagnostic threshold for this test, although NICE recommends RBG >7.0 mmol/L as an indication for OGTT, while WHO/IADPSG recommend >11.1 mmol/L as diagnostic of clinical (overt) diabetes. This >11.1 mmol/L cut-off would have yielded only one positive case in this study. Due to non-consensus on diagnostic threshold for RBG, and the test not being sufficiently sensitive to detect GDM, the effectiveness of RBG needs to be further explored with emphasis on the reference threshold and optimum

postprandial conditions under which the test would be most accurate.

FPG had the highest true positive rate and the test performance was very good, thereby making it useful for ruling out GDM and minimizing missed cases. In the USA, FPG reduced 7% missed cases when used as a screening tool.³⁰ The test has an added advantage of being readily available, easy to perform, relatively inexpensive, requiring minimal client preparation and is applicable for both GDM screening and diagnostic purposes. When assessing diagnostic accuracy, tests with high discriminatory measures are helpful in screening and making health policy decisions, whereas highly predictive tests are useful in predicting the probability of the disease at the individual level in order to facilitate client-centered case management.²³ The NICE cut-off (≥ 5.6 mmol/L) had higher PPV, thus making it the test and cut-off option for predicting the probability of GDM in an individual. The WHO cut-off (≥ 5.1 mmol/L) was the most sensitive but has the potential for overdiagnoses due to the high number of false positives. Nonetheless, its high sensitivity makes it the test of choice if prevention of missed cases is the focus.

Even though WHO,¹⁰ IADPSG,¹² and NICE⁹ have established FPG as a diagnostic test for GDM, the preferred test is the 'gold standard' 2-hour OGTT. The criteria for diagnosing diabetes in pregnancy based on the 1-hour postload value have not been fully established,¹⁰ and we have found its sensitivity to be <50%. Therefore, in settings where 2-hour postprandial glucose-load testing is unavailable for confirmation, GDM diagnosis and decision to commence therapy should be established based on preprandial glucose values ≥ 5.6 mmol/L as substitute for an OGTT. This should be complemented by the presence of risk factors.

In the wake of updated guidelines for detecting hyperglycemia in pregnancy whereby diagnostic thresholds have reduced,⁹⁻¹² there are widespread concerns about overdiagnosing and the associated implications particularly for fragile health systems. Nonetheless, evidence of increasing prevalence of pregnancy-triggered diabetes from negligible rates to almost 30% in certain settings in African⁵⁻⁷ necessitates a critical review of diagnostic strategies by health systems if adverse outcomes such as newborn macrosomia, hypoglycemia, hyperinsulinemia, respiratory distress, perinatal mortality, maternal pre-eclampsia, cesarean delivery¹ and risk for long-term metabolic conditions³ are to be reduced.

Our study is not without limitations. Although HbA1c was performed in the fasting state (n=445), few non-fasting pregnant women (n=35) were included. Nonetheless prandial state has been shown not to affect HbA1c levels. Whereas random blood glucose was checked from capillary finger prick before 20 gestational weeks, FPG was checked from venous blood from 20 to 34 weeks. Variability in sampling collection procedure and testing times poses a challenge when equating these two glucose values. Because pregnant women also receive ANC in primary facilities where laboratory services could

be unavailable, accuracy of point-of-care testing using fasting whole blood glucose obtained from a capillary finger prick should have been investigated as well.

Despite these limitations, unlike in most diagnostic accuracy studies where retrospective data sources were used, our study was conducted prospectively using the blind comparison with a gold standard design, a type of randomized controlled trial design that compares the accuracy of diagnostic procedures in the same individual. This design increases the probability of the test outcomes being close to the true values. In low-income and middle-income countries, studies of this nature are mostly concentrated in tertiary hospitals. Our study shows higher glycemic values among primary healthcare users. This provides evidence of obstetric transition even in rural communities and highlights the need to enhance access to quality maternal healthcare at primary healthcare levels.

Implications for practice

Gestational diabetes is common in the study population and was high among primary healthcare users. In as much as overdiagnoses (false positive) will increase costs, underdiagnoses (false negative) have adverse public health implications. Selective screening using risk factors is better compared with glycosuria and random blood glucose because these tests missed majority of cases, thereby decreasing opportunities for diagnostic testing. Because FPG was the most sensitive test with minimal false negatives diagnoses, its integration into all GDM detection procedures through universal testing of all pregnant women is recommended as it is easier to accomplish for many women compared with OGTT. At the population level, the WHO diagnostic criteria ≥ 5.1 mmol/L could be useful for screening purposes because of its discriminatory properties, a key consideration for making health policy decisions. At the individual level, the NICE diagnostic criteria ≥ 5.6 mmol/L, which has higher GDM prediction properties, could be useful in making therapeutic decisions. In settings where 2-hour postprandial OGTT is difficult to obtain, fasting glucose value ≥ 5.6 mmol/L should be complemented with risk factor assessment and collectively serve as the basis to commence treatment.

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Data sharing statement This study is part of a larger prospective cohort study aimed at assessing the effects of hyperglycemia in pregnancy on maternal, fetal and neonatal outcomes and the extent of normoglycemia in the short-term period after delivery. Both data collection and data analysis are still ongoing. Hence the authors are unable to publicly share the data sets generated.

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REFERENCES

1. Metzger BE, Lowe LP, Dyer AR, *et al.* Hyperglycemia and adverse pregnancy outcomes. *N Engl J Med* 2008;358:1991–2002.
2. Wendland EM, Torloni MR, Falavigna M, *et al.* Gestational diabetes and pregnancy outcomes—a systematic review of the World Health Organization (WHO) and the International Association of Diabetes in Pregnancy Study Groups (IADPSG) diagnostic criteria. *BMC Pregnancy Childbirth* 2012;12:23.
3. Damm P, Houshmand-Oeregaard A, Kelstrup L, *et al.* Gestational diabetes mellitus and long-term consequences for mother and offspring: a view from Denmark. *Diabetologia* 2016;59:1396–9.
4. Guariguata L, Linnenkamp U, Beagley J, *et al.* Global estimates of the prevalence of hyperglycaemia in pregnancy. *Diabetes Res Clin Pract* 2014;103:176–85.
5. Macaulay S, Dunger DB, Norris SA. Gestational diabetes mellitus in Africa: a systematic review. *PLoS One* 2014;9:e97871.
6. Hall V, Thomsen RW, Henriksen O, *et al.* Diabetes in Sub Saharan Africa 1999–2011: epidemiology and public health implications. A systematic review. *BMC Public Health* 2011;11:564.
7. Adam S, Rheeder P. Screening for gestational diabetes mellitus in a South African population: prevalence, comparison of diagnostic criteria and the role of risk factors. *S Afr Med J* 2017;107:523–7.
8. Anand M, Anand M, Mahajan DS. To study the incidence of gestational diabetes mellitus and risk factors associated with GDM. *International Journal of Advances in Medicine* 2017;4:112–6.
9. NICE. *Diabetes in pregnancy. Management of diabetes and its complications from pre-conception to the postnatal period. NICE guideline 3 Methods, evidence and recommendations.* London, UK: National Institute for Health and Care Excellence and National Collaborating Centre for Women's and Children's Health, 2015.
10. WHO. *Diagnostic criteria and classification of hyperglycaemia first detected in pregnancy.* Geneva, Switzerland: World Health Organization, 2013.
11. American Diabetes Association. (2) Classification and diagnosis of diabetes. *Diabetes Care* 2015;38(Suppl 1):S8–S16.
12. 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: response to weinert. *Diabetes Care* 2010;33:e98–82.
13. Metzger BE, Gabbe SG, Persson B, *et al.* International association of diabetes and pregnancy study groups Consensus Panel. International association of diabetes and pregnancy study groups recommendations on the diagnosis and classification of hyperglycemia in pregnancy. *Diabetes Care* 2010;33:676–82.
14. Ministry of Health. *Standard treatment guidelines. Ghana National Drugs Programme (GNDP).* 6th edn. Accra, Ghana, West Africa: Yamens Press. Ltd, 2010.
15. Agbozo F, Abubakari A, Der J, *et al.* Prevalence of low birth weight, macrosomia and stillbirth and their relationship to associated maternal risk factors in Hohoe Municipality, Ghana. *Midwifery* 2016;40:200–6.
16. Oppong SA, Ntumu MY, Amoakoh-Coleman M, *et al.* Gestational diabetes mellitus among women attending prenatal care at

- Korle-Bu Teaching Hospital, Accra, Ghana. *Int J Gynaecol Obstet* 2015;131:246–50.
17. Alto WA. No need for glycosuria/proteinuria screen in pregnant women. *J Fam Pract* 2005;54:978.
 18. Cersosimo E, Ajmal M, Naukam RJ, *et al.* Role of the kidney in plasma glucose regulation during hyperglycemia. *Am J Physiol* 1997;272:E756–E761.
 19. Bossuyt PM, Reitsma JB, Bruns DE, *et al.* STARD 2015: an updated list of essential items for reporting diagnostic accuracy studies. *BMJ* 2015;351:h5527.
 20. Ghana Statistical Service. Population and housing census. regional analytical report, volta region. *Accra, Ghana Ghana Statistical Service* 2010;2013:1–153.
 21. Cochran WG. *Sampling techniques*. US: John Wiley & Sons, 2007.
 22. Faul F, Erdfelder E, Lang AG, *et al.* G*Power 3: a flexible statistical power analysis program for the social, behavioral, and biomedical sciences. *Behav Res Methods* 2007;39:175–91.
 23. Raslich MA, Markert RJ, Stutes SA. Selecting and interpreting diagnostic tests. *Biochem Med* 2007;17:151–61.
 24. Mallett S, Halligan S, Thompson M, *et al.* Interpreting diagnostic accuracy studies for patient care. *BMJ* 2012;345:e3999.
 25. Glas AS, Lijmer JG, Prins MH, *et al.* The diagnostic odds ratio: a single indicator of test performance. *J Clin Epidemiol* 2003;56:1129–35.
 26. Hanna FW, Peters JR. Screening for gestational diabetes: past, present and future. *Diabet Med* 2002;19:351–8.
 27. Moya J, Phillips L, Sanford J, *et al.* A review of physiological and behavioral changes during pregnancy and lactation: potential exposure factors and data gaps. *J Expo Sci Environ Epidemiol* 2014;24:449–58.
 28. Narayan KM, Pettitt DJ. Glycosylated hemoglobin in pregnancy. *Diabetes Care* 1996;19:1455–55.
 29. Berggren EK, Boggess KA, Mathew L, *et al.* First Trimester Maternal Glycated Hemoglobin and Sex Hormone-Binding Globulin Do Not Predict Third Trimester Glucose Intolerance of Pregnancy. *Reprod Sci* 2017;24:613–8.
 30. 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;28:825–8.