






## RESEARCH ARTICLE

# Hyperglycemia in pregnancy diagnosed using glycated hemoglobin (HbA<sub>1c</sub>) in Uganda: a preliminary cross-sectional report [version 1; peer review: 1 approved]

Felix Bongomin <sup>1,2</sup>, Andrew P. Kyazze <sup>2</sup>, Sandra Ninsiima<sup>3</sup>, Ronald Olum <sup>1</sup>, Gloria Nattabi<sup>1</sup>, Winnie Nabakka<sup>3</sup>, Rebecca Kukunda<sup>3</sup>, Charles Batte <sup>4</sup>, Phillip Ssekamatte<sup>3,5</sup>, Joseph Baruch Baluku<sup>6,7</sup>, Davis Kibirige<sup>3,8</sup>, Stephen Cose <sup>8,9</sup>, Irene Andia-Biraro<sup>1,8,9</sup>

<sup>1</sup>Department of Medicine, Makerere University, Kampala, Uganda

<sup>2</sup>Department of Medical Microbiology & Immunology, Gulu University, Gulu, Uganda

<sup>3</sup>Department of Medicine, Uganda Martyrs Hospital Lubaga, Kampala, Uganda

<sup>4</sup>Department of Medicine, Makerere Lung Institute, Makerere University College of Health Sciences, Kampala, Uganda

<sup>5</sup>Department of Immunology and Molecular Biology, School of Biomedical Sciences, Makerere University College of Health Sciences, Kampala, Uganda

<sup>6</sup>Division of Pulmonology, Mulago National Referral Hospital, Kampala, Uganda

<sup>7</sup>Directorate of Programs, Mildmay Uganda, Wakiso, Uganda

<sup>8</sup>Medical Research Council/Uganda Virus Research-Institute Uganda Research Unit, Entebbe, Uganda

<sup>9</sup>Department of Clinical Research, Faculty of Infectious and Tropical Disease (ITD), London School of Hygiene and Tropical Medicine, London, United Kingdom

**V1** First published: 15 Dec 2020, 2:70  
<https://doi.org/10.35241/emeraldopenres.14014.1>

Latest published: 15 Dec 2020, 2:70  
<https://doi.org/10.35241/emeraldopenres.14014.1>

## Abstract

**Background:** Hyperglycemia in pregnancy (HIP) is a common medical complication during pregnancy and is associated with several short and long-term maternal-fetal consequences. We aimed to determine the prevalence and factors associated with HIP among Ugandan women.

**Methods:** We consecutively enrolled eligible pregnant women attending antenatal care at Kawempe National Referral Hospital, Kampala, Uganda in September 2020. Mothers known to be living with diabetes mellitus or haemoglobinopathies and those with anemia (hemoglobin <11g/dl) were excluded. Random blood sugar (RBS) and glycated hemoglobin A1c (HbA<sub>1c</sub>) were measured on peripheral venous blood samples. HIP was defined as an HbA<sub>1c</sub> ≥5.7% with its subsets of diabetes in pregnancy (DIP) and prediabetes defined as HbA<sub>1c</sub> of ≥6.5% and 5.7-6.4% respectively. ROC curve analysis was performed to determine the optimum cutoff of RBS to screen for HIP.

**Results:** A total of 224 mothers with a mean (± SD) age 26±5 years were enrolled, most of whom were in the 2<sup>nd</sup> or 3<sup>rd</sup> trimester (94.6%, n=212) with a mean gestation age of 26.6±7.3 weeks. Prevalence of


## Open Peer Review

Approval Status 

1

### version 1

15 Dec 2020

1. **Ankia Coetzee** , Stellenbosch University and Tygerberg Hospital, Stellenbosch, South Africa

Any reports and responses or comments on the article can be found at the end of the article.

HIP was 11.2% (n=25) (95% CI: 7.7-16.0). Among the mothers with HIP, 2.2% (n=5) had DIP and 8.9% (n=20) prediabetes. Patients with HIP were older (28 years vs. 26 years,  $p=0.027$ ), had previous tuberculosis (TB) contact (24% vs. 6.5%,  $p=0.003$ ) and had a bigger hip circumference (107.8 ( $\pm 10.4$ ) vs. 103.3 ( $\pm 9.7$ ) cm,  $p = 0.032$ ). However only previous TB contact was predictive of HIP (odds ratio: 4.4, 95% CI: 1.2-14.0;  $p=0.022$ ). Using HbA<sub>1c</sub> as a reference variable, we derived an optimum RBS cutoff of 4.75 mmol/L as predictive of HIP with a sensitivity and specificity of 90.7% and 56.4% (area under the curve = 0.75 (95% CI: 0.70-0.80,  $p<0.001$ )), respectively.

**Conclusions:** HIP is common among young Ugandan women, the majority of whom are without identifiable risk factors.

### Keywords

Hyperglycemia in pregnancy, prediabetes, Hemoglobin A1c, Uganda



This article is included in the [Healthier Lives](#) gateway.

**Corresponding author:** Felix Bongomin ([f.bongomin@gu.ac.ug](mailto:f.bongomin@gu.ac.ug))

**Author roles:** **Bongomin F:** Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Kyazze AP:** Investigation, Project Administration, Resources, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Ninsiima S:** Investigation, Project Administration, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Olum R:** Data Curation, Formal Analysis, Investigation, Methodology, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Nattabi G:** Methodology, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Nabakka W:** Investigation, Methodology, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Kukunda R:** Investigation, Methodology, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Batte C:** Funding Acquisition, Project Administration, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Ssekamatte P:** Investigation, Methodology, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Baruch Baluku J:** Conceptualization, Investigation, Methodology, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Kibirige D:** Investigation, Methodology, Project Administration, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Cose S:** Funding Acquisition, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; **Andia-Biraro I:** Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing

**Competing interests:** No competing interests were disclosed.

**Grant information:** Research reported in this publication was supported by the Forgarty International Centre of the National Institutes of Health under Award Number D43 TW0011401. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health. Also, this work was partially supported by the African Career Accelerator Award grant held by Dr. Irene Andia-Biraro, which is funded by the CRICK African Network and is hosted at the MRC/UVRI & LSHTM Uganda Research Unit in Entebbe, Uganda.

*The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.*

**Copyright:** © 2020 Bongomin F *et al.* This is an open access article distributed under the terms of the [Creative Commons Attribution License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**How to cite this article:** Bongomin F, Kyazze AP, Ninsiima S *et al.* **Hyperglycemia in pregnancy diagnosed using glycated hemoglobin (HbA<sub>1c</sub>) in Uganda: a preliminary cross-sectional report [version 1; peer review: 1 approved]** Emerald Open Research 2020, 2:70 <https://doi.org/10.35241/emeraldopenres.14014.1>

**First published:** 15 Dec 2020, 2:70 <https://doi.org/10.35241/emeraldopenres.14014.1>

## Introduction

Pregnancy is naturally characterized by insulin resistance and hyperinsulinemia leading to hyperglycemia, the most common endocrinopathy during the gestation period (Farrar, 2016; Saravanan *et al.*, 2020). Previously, any hyperglycemia detected in pregnancy was termed gestational diabetes (GDM) (WHO, 2013). However, recently, the term hyperglycemia in pregnancy (HIP) has been proposed by the World Health Organization (WHO) (Diabetes Canada Clinical Practice Guidelines Expert Committee *et al.*, 2018; WHO, 2018). HIP classifies hyperglycemia based on both onset and severity (Diabetes Canada Clinical Practice Guidelines Expert Committee *et al.*, 2018; WHO, 2018). It includes the more severe manifestations of total diabetes in pregnancy (comprising of known and previously undiagnosed diabetes in pregnancy [DIP]), which may persist in the post-partum period and a more benign form, GDM (WHO, 2013). DIP and GDM are together termed hyperglycemia first detected in pregnancy (WHO, 2013).

HIP is a growing public health concern and adversely affects maternal and child health, and is likely to contribute to the growing global diabetes epidemic (Bianco & Josefson 2019; Guariguata *et al.*, 2014). Depending on the diagnostic criteria used and the population of pregnant women studied, the resulting prevalence of HIP can vary widely. Results from a recent survey on HIP prevalence in 173 countries found country-specific prevalence estimates ranging from <1% in Germany up to 28% for a study in Nepal, using a variety of criteria (Jiwani *et al.*, 2012). Globally, HIP has been estimated to affect nearly 16.9%, or 21.4 million, live births among women of reproductive age, with total diabetes in pregnancy accounting for an estimated 16.0% of these cases (Guariguata *et al.*, 2014). In this report, more than 90% of cases of HIP were estimated to occur in low- and middle-income countries (LMICs), with South-East Asian and African regions having the highest number of live births affected with HIP at over 6.0 (23.2%) and 4.3 (16.0%) million cases, respectively (Guariguata *et al.*, 2014).

Previous studies assessing the prevalence of HIP have mostly concentrated on high risk mothers, such as those with advanced age, high gravidity or in a specific period of gestation age, which may not reflect the true prevalence in the general population of pregnant women (Adefisan *et al.*, 2020; Cosson *et al.*, 2019; Mukuve *et al.*, 2020). Moreover, screening and intervention on HIP during antenatal care (ANC) are not routine in most LMICs, making an accurate estimation of the burden of this treatable condition largely impossible. This could be due to the several caveats associated with current tests, which requires overnight fasts, multiple clinic visits and the oral glucose tolerance test (OGTT) which is labor intensive.

Despite the serious public health implications of HIP, there has been no universal definition and no universal standards for screening and diagnosis, and a wide variety of methods are applied (Guariguata *et al.*, 2014). However, fasting plasma glucose (FPG), 1-hour, 2-hour or 3-hour plasma glucose following a 75g OGTT, interpreted according to the American Diabetes Association (ADA), WHO or the International

Association of the Diabetes and Pregnancy Study Group (IADPSG) criteria are the most commonly used methods (Guariguata *et al.*, 2014; Meek *et al.*, 2020). Glycated hemoglobin (HbA<sub>1c</sub>) can also be used to screen for and diagnose HIP, especially in early pregnancy (Goyal *et al.*, 2020). Trimester specific cutoff values for HbA<sub>1c</sub> have recently been proposed (Sánchez-González *et al.*, 2018), though not widely validated or adopted by international guidelines (Sánchez-González *et al.*, 2018).

The aim of this study was to determine the prevalence of HIP and its associated risk factors among pregnant women attending ANC in Uganda, irrespective of their gestation age, using HbA<sub>1c</sub>. Secondly, we sought to determine an appropriate random blood sugar (RBS) cutoff value for screening for HIP in our setting.

## Methods

### Study design and setting

We conducted a cross-sectional study in a large specialized obstetrics and gynecology national referral hospital in Kampala, Uganda in September 2020.

This study enrolled pregnant women attending the ANC clinic at the directorate of Obstetrics and Gynecology at Kawempe National Referral Hospital (KNRH), which also serves as the academic teaching hospital for Makerere University College of Health Sciences. KNRH was purposively selected due to its central location attracting a large number of patients from Kampala and its surrounding districts, thus representing the demographics of both urban and peri-urban patient populations. On average, about 50-60 mothers attend the ANC clinic at KNRH from Tuesday to Thursday every week.

### Study population

Eligible participants were all pregnant women attending ANC (regardless of gestation age) during the study period who provided informed consent to participate in the study. Pregnant women known to be living with diabetes or haemoglobinopathies were excluded. In addition, patients with anemia (hemoglobin <11g/dl) were excluded at analysis.

### Sample size and sampling procedure

With an estimated prevalence of HIP at 15.6% in Uganda (Kiiza *et al.*, 2020), using the formula for the determination of sample size for prevalence studies (Kish-Leslie) (Kish, 1965), with an assumed non-response rate of 10%, precision of 5%, and a Z-score of 1.96 at 95% confidence interval, a sample size of 217 was anticipated. Eligible participants were identified, and consecutively sampled with the assistance of a senior nurse at the ANC clinic and two other trained study nurses.

### Data collection

Study clinicians administered a semi-structured study questionnaire through a face-to-face interview to collect information regarding risk factors and symptoms for HIP and maternal characteristics: age, gravidity, education level, occupation, marital status, HIV status, tuberculosis contact, gestation age, history of abortion, smoking and alcohol usage, and the number of ANC

visits in the current pregnancy. Gestation age was estimated using the date of the last normal menstrual period. Polyuria, polydipsia, and polyphagia were considered classic symptoms of diabetes.

**Diagnosis of hyperglycemia in pregnancy.** All consenting mothers were subjected to a RBS and HbA<sub>1c</sub> tests. RBS was performed at the point of care on venous blood samples using the On-Call™ Plus Glucometer (ACON Biotech, China) according to manufacturer's instruction. HbA<sub>1c</sub> was estimated using Cobas® 6000 analyzer series (Roche Diagnostics) at the Central Diagnostic Laboratory Services (CDLS), Medical Research Council/Uganda Virus Research Institute and London School of Hygiene and Tropical Medicine Uganda Research Unit laboratory within 24 hours of sample collection. Prediabetes was defined as HbA<sub>1c</sub> of 5.7% to 6.4% and overt diabetes (diabetes in pregnancy, DIP) as HbA<sub>1c</sub> of 6.5% or higher, according to the American Diabetes Association (Goyal *et al.*, 2020) and consistent with guidelines of the IADPSG (Meek *et al.*, 2020; Saravanan *et al.*, 2020), and the WHO classification of hyperglycemia first diagnosed in pregnancy (WHO, 2018). Patients with HIP were referred for the appropriate clinical care according to the national guidelines.

**Complete blood count.** A complete blood count was performed using the HumaCount 5D Hematology System (Wiesbaden, Germany) using 3 mL of blood samples collected in EDTA tubes within 12 hours from the time of sample collection. Patients with hemoglobin concentration of 11g/dl or lower were regarded as having anemia according to the WHO definition (World Health Organization, 2011) of anemia in pregnancy and were subsequently excluded from the study at analysis. Anemic patients were also referred for appropriate management by the clinical team.

**Anthropometrical assessment.** Weight was measured with minimal clothing and without shoes using a digital bathroom weighing scale (SECA-Germany), placed on a flat surface and recorded to the nearest 0.1kg. Height was measured using a calibrated stadiometer. Waist and hip circumferences were measured using a tailor's measuring tape to the nearest 0.1 cm. Body mass index and Waist-Hip Ratios were calculated accordingly. Brachial blood pressure was measured from both arms while mothers were sitting down with their feet flat on the ground using an automated machine with an appropriate adult cuff size. The average of the two measurements was considered as the participant's blood pressure. Women were classified as hypertensive if systolic and diastolic blood pressure were  $\geq 140$ mmHg and  $\geq 90$ mmHg, respectively, and normal if blood pressure is less than 140/90mmHg.

#### Statistical analysis

Statistical analyses were performed using STATA version 16 (StataCorp LLC, College Station, TX, USA) and GraphPad Prism version 8.0.4 (GraphPad Software, La Jolla, CA, USA). The data were expressed as absolute numbers and percentages for categorical variables, and as means and standard deviations (mean $\pm$ SDs) for continuous variables. Shapiro-Wilk normality test was applied to evaluate all quantitative variables

to select the appropriate test. Welch's One-Way ANOVA was used to compare continuous data across groups. Chi-square or Fisher's Exact tests were used to assess associations between HIP across demographic and clinical characteristics of participants. Variables with a p value  $< 0.2$  were fitted into a multivariable logistic regression model to adjust for confounders. Receiver operating characteristics (ROC) curve analysis was performed to determine the optimum cutoff of RBS in those patients who met criteria for HIP in relation to HbA<sub>1c</sub> test. Area under the ROC curve (AUC) was shown with 95% Wilson confidence intervals (CI). Optimal diagnostic cut-off value for RBS were calculated using Youden's J statistic (sensitivity+specificity-1). For this analysis, the hypothesis was that at the optimal RBS cut off, the AUC = 0.7. In all analyses,  $P < 0.05$  was considered significant at 95% CI.

#### Ethical considerations

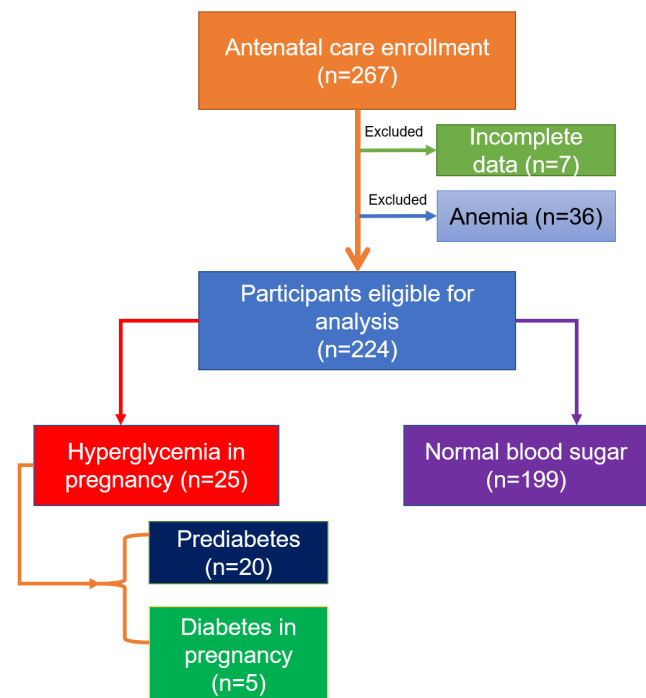
This study was approved by the Makerere University School of Medicine Ethics and Research Committee (reference number #REC REF 2020-113). All mothers provided informed written consent to participate after the study procedure, risks and benefits were explained to them.

#### Results

In September 2020, a total of 267 pregnant women participated in the study. However, 43 participants were excluded due to either incomplete data or presence of anemia (Figure 1).

#### Baseline characteristics of study participants

Of the 224 eligible participants, most were married (91.1%, n=204), and attending ANC for the first time in the current



**Figure 1. Study flow diagram.**

pregnancy (63.4%, n=142). Over three quarters had attended post-primary education (77.7%, n=174) and were business-women (42.2%, n=99). The mean age of the women was 26 years (SD± 5), of which 138 (61.6%) were ≥25 years old. Just over one-third of the participants were primigravida (34.8%, n=78), and the majority of the mothers were in their 2<sup>nd</sup> or 3<sup>rd</sup> trimester of pregnancy (94.6%, n=212), with a mean gestation age of 26.6 weeks (SD ± 7.3) (Table 1). In total, 43 (19.2%) women had at least one of the classic symptoms of diabetes.

**Table 1. Sociodemographic and maternal characteristics of the study participants.**

Participant variable	N (%) or Mean ± SD
<b>Antenatal care visit at enrollment</b>	
First	142 (63.4)
Second	27 (12.1)
Third	19 (8.4)
Fourth and more	36 (16.1)
<b>Age (years)</b>	26 ± 5.0
<25 years	86 (38.4)
≥25 years	138 (61.6)
<b>Marital status</b>	
Married	204 (91.1)
Single	12 (5.4)
Widowed	8 (3.6)
<b>Education level</b>	
Informal	4 (1.8)
Primary	46 (20.5)
Secondary	117 (52.2)
Tertiary	57 (25.5)
<b>Occupational status</b>	
Business	99 (42.2)
Professional	46 (20.5)
Unemployed	79 (35.3)
<b>Smoking status</b>	
Former	2 (0.9)
Never	222 (99.1)
<b>Alcohol usage</b>	
Current	12 (5.4)
Former	38 (17.0)
Never	174 (77.7)

Participant variable	N (%) or Mean ± SD
<b>Family history of diabetes</b>	
No	185 (82.6)
Yes	39 (17.4)
<b>District of residence</b>	
Kampala	156 (69.6)
Wakiso	62 (27.7)
Mpigi	1 (0.5)
Mukono	4 (1.8)
Entebbe	1 (0.5)
<b>Residence</b>	
Peri-Urban	63 (28.1)
Urban	161 (71.9)
<b>HIV status</b>	
Positive	6 (2.7)
Negative	218 (97.3)
<b>BCG scar</b>	
Yes	162 (72.3)
No	62 (27.7)
<b>Family history of tuberculosis</b>	
Yes	22 (9.8)
No	202 (90.2)
<b>Tuberculosis contact</b>	
Yes	19 (8.5)
No	205 (91.5)
<b>Family size</b>	
≤4	177 (79.0)
≥5	47 (21.0)
<b>Symptoms of diabetes</b>	
Yes	43 (19.2)
No	181 (80.8)
<b>Gravidity</b>	
Primigravida	78 (34.8)
Multigravida	120 (53.6)
Grand multigravida	19 (8.5)
Great grand multigravida	7 (3.1)
<b>Previous abortion</b>	
Yes	36 (16.1)
No	188 (83.9)

Participant variable	N (%) or Mean $\pm$ SD
<b>Gestation age at enrollment (weeks)</b>	26.6 $\pm$ 7.3
<b>Trimester at enrollment</b>	
1	12 (5.4)
2	101 (45.1)
3	111 (49.6)
<b>Anthropometry</b>	
Weight (kilograms)	68.9 $\pm$ 12.4
Height (meters)	1.6 $\pm$ 0.06
Body mass index (kg/m <sup>2</sup> )	27.3 $\pm$ 4.8
Waist circumference (centimeters)	95.2 $\pm$ 10.4
Hip circumference (centimeters)	103.8 $\pm$ 9.8
Waist-hip circumference	0.92 $\pm$ 0.08
<b>Blood pressure at enrollment</b>	
Systolic blood pressure (mmHg)*	125 $\pm$ 18
Diastolic blood pressure (mmHg)*	78 $\pm$ 12.8
Normal	188 (91.5)
Hypertensive	19 (8.5)

\*Average of two measurements taken.

### Prevalence of HIP

RBS and HbA<sub>1c</sub> was performed for all the 224 participants. The median (range) RBS was 4.6 (2.8-8.0) mmol/l. The median (range) of HbA<sub>1c</sub> was 5.2 (3.6-14.9).

The overall prevalence of HIP was 11.2% (n=25) (95% CI: 7.7-16.0); 2.2% (n=5) (95% CI: 1.0-5.1) had DIP, and 8.9% (n=20) (95% CI: 5.9-13.4) prediabetes using the WHO criteria. Patients with HIP were slightly older than those without (28 years vs. 26 years, p=0.027), had previous tuberculosis contact (24% vs. 6.5%, p=0.003), had a bigger hip circumference (107.8 ( $\pm$ 10.4) vs. 103.3 ( $\pm$ 9.7) cm, p = 0.032) and a higher proportion of urban dwellers had HIP compared to their rural counterparts (88% vs.69.8%), though this was not statistically significant (p=0.062) (Table 2). However, after accounting for important confounders in a multivariable logistic regression models, none of these factors showed a statistically significant association (Table 3).

The mean RBS was slightly higher in those with HIP compared to those with normal HbA<sub>1c</sub>; however this was not to statistical significance (Figure 2).

Using HbA<sub>1c</sub> as a reference variable, ROC curve and the AUC for RBS as a predictor of HIP was 0.75 (95% CI: 0.70-0.80,

**Table 2. Prevalence of hyperglycemia among pregnant women at Kawempe National Referral Hospital.**

Participant variable	HIP (n=25)	No HIP (n=199)	P-value
	N (%)	N (%)	
<b>ANC visit at enrollment</b>	2 (1)	2 (1)	0.857
First	16 (64.0)	126 (63.3)	0.277
Second	5 (20)	22 (11.1)	
Third	0 (0)	19 (9.5)	
Fourth and more	4 (16)	32 (16.1)	
<b>Age, mean (<math>\pm</math>SD)</b>	28.4 $\pm$ 4.8	26.0 $\pm$ 5.0	<b>0.027</b>
<25 years	7 (28)	79 (39.7)	0.257
$\geq$ 25 years	18 (72)	120 (60.3)	
<b>Marital status</b>			
Married	25 (100)	179 (89.9)	0.252
Single	0 (0)	12 (6.0)	
Widowed	0 (0)	8 (4.0)	
<b>Education level</b>			
Informal	0 (0)	4 (2.0)	0.661
Primary	7 (28)	39 (19.6)	
Secondary	13 (52)	104 (52.3)	
Tertiary	5 (20)	52 (26.1)	
<b>Occupational status</b>			
Business	13 (52)	86 (43.2)	0.666
Professional	5 (20)	41 (20.6)	
Unemployed	7 (28)	72 (36.2)	
<b>Smoking status</b>			
Former	0 (0)	2 (1.0)	>0.999
Never	25 (100)	197 (99.0)	
<b>Alcohol usage</b>			
Current	2 (8)	10 (5.0)	0.324
Former	6 (24)	32 (16.1)	
Never	17 (68)	157 (78.9)	
<b>Family history of diabetes</b>			
No	20 (80)	165 (82.9)	0.717
Yes	5 (20)	34 (17.1)	
<b>District of residence</b>			
Kampala	17 (68)	139 (69.8)	0.061
Wakiso	6 (24)	56 (28.1)	
Mpigi	1 (4)	3 (1.5)	
Mukono	0 (0)	1 (0.5)	
Entebbe	1 (4)	0 (0.0)	

Participant variable	HIP (n=25)	No HIP (n=199)	P-value
	N (%)	N (%)	
<b>Residence</b>			
Peri-Urban	3 (12)	60 (30.2)	0.062
Urban	22 (88)	139 (69.8)	
<b>HIV status</b>			
Positive	1 (4)	5 (2.5)	0.513
Negative	24 (96)	194 (97.5)	
<b>BCG scar</b>			
Yes	10 (40)	52 (26.1)	0.144
No	15 (60)	147 (73.9)	
<b>Family history of tuberculosis</b>			
Yes	4 (16)	18 (9.0)	0.282
No	21 (84)	181 (91.0)	
<b>Tuberculosis contact</b>			
Yes	6 (24)	13 (6.5)	<b>0.003</b>
No	19 (76)	186 (93.5)	
<b>Family size</b>			
≤4	19 (76)	158 (79.4)	0.694
≥5	6 (24)	41 (20.6)	
<b>Symptoms of diabetes</b>			
Yes	4 (16)	39 (19.6)	0.793
No	21 (84)	160 (80.4)	
<b>Gravidity</b>			
Primigravida	4 (16)	74 (37.2)	0.112
Multigravida	16 (64)	104 (52.3)	
Grand multigravida	3 (12)	16 (8.0)	
Great grand multigravida	2 (8)	5 (2.5)	
<b>Previous abortion</b>			
Yes	7 (28)	29 (14.6)	0.085
No	18 (72)	170 (85.4)	
<b>Gestation age at enrollment (weeks; mean ± SD)</b>	25.8 ± 7.7	26.7 ± 7.3	0.558
<b>Trimester at enrollment</b>			
1	1 (4)	11 (5.5)	0.922
2	12 (48)	89 (44.7)	
3	12 (48)	99 (49.7)	

Participant variable	HIP (n=25)	No HIP (n=199)	P-value
	N (%)	N (%)	
<b>Anthropometry, mean ± SD</b>			
Weight (kilograms)	71.4 ± 15.0	68.6 ± 12.0	0.274
Height (meters)	1.6 ± 0.07	1.6 ± 0.06	0.408
BMI (kg/m <sup>2</sup> )	28.7 ± 5.9	27.1 ± 4.6	0.113
Waist circumference (centimeters)	97.5 ± 13.4	94.9 ± 10.0	0.227
Hip circumference (centimeters)	107.8 ± 10.4	103.3 ± 9.7	<b>0.032</b>
Waist-hip circumference	0.92 ± 0.08	0.92 ± 0.08	0.327
<b>Blood pressure at enrollment, Mean ± (SD)</b>			
Systolic blood pressure (mmHg)*	122 ± 12	125 ± 18	0.394
Diastolic blood pressure (mmHg)*	77 ± 8	78 ± 13	0.759
Normal	23 (92)	165 (82.9)	0.386
Hypertension	2 (8)	34 (17.1)	

HIP, hyperglycemia in pregnancy. \*Average of two measurements taken.

**Table 3. A multivariable logistic regression model showing factors associated with hyperglycemia among pregnant women at Kawempe National Referral Hospital.**

Demographic and clinical characteristics	Adjusted Odds Ratio	95% CI	P-value
<b>Age</b>	1.04	0.92 - 1.16	0.549
<b>District of residence</b>			
Entebbe	1.00		
Kampala	1.55	0.49 - 4.92	0.458
Mukono	9.36	0.6 - 146.93	0.112
<b>Residence</b>			
Rural	1.00		
Urban	3.80	0.92 - 15.63	0.064
<b>BCG scar</b>			
Yes	1.00		
No	1.72	0.64 - 4.58	0.280

Demographic and clinical characteristics	Adjusted Odds Ratio	95% CI	P-value
<b>Tuberculosis contact</b>			
No	1.00		
Yes	4.14	1.23 - 13.98	0.022
<b>Gravidity</b>			
Primigravida	1.00		
Multigravida	1.67	0.43 - 6.43	0.458
Grand multigravida	2.08	0.3 - 14.55	0.462
Great grand multigravida	4.01	0.28 - 56.64	0.304
<b>Previous abortion</b>			
No	1.00		
Yes	1.24	0.37 - 4.21	0.728
<b>Body mass index</b>	0.98	0.84 - 1.15	0.826
<b>Hip circumference</b>	1.06	0.98 - 1.14	0.132

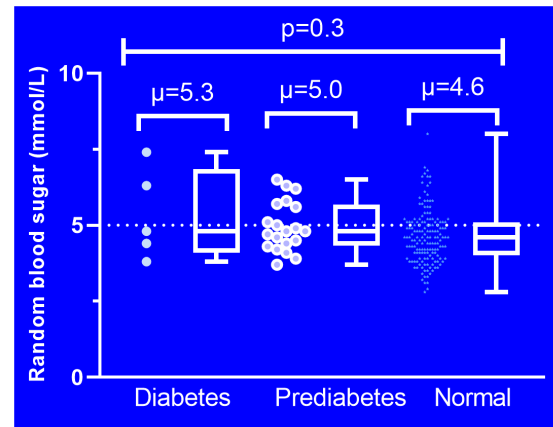
CI, confidence interval.

$p < 0.001$ ) (Figure 3). We derived optimum cutoffs for RBS of 4.75 mmol/L with a sensitivity and specificity of 90.7% and 56.4%, respectively. At a lower RBS cutoff of 4.0 mmol/L, the sensitivity and specificity was 99.5% and 23.6%, respectively, and at a higher RBS cutoff of 5.5 mmol/L, the sensitivity and specificity was 26.9% and 83.5%, respectively.

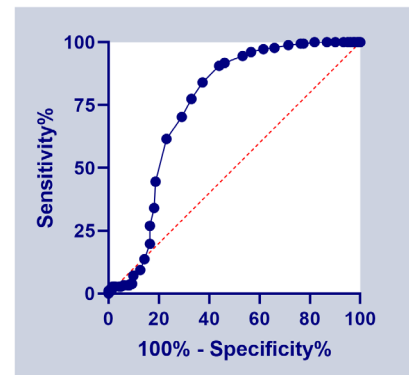
## Discussion

The use of HbA<sub>1c</sub> for screening, diagnosis and monitoring of diabetes and prediabetes in pregnancy remains a work in progress with several unanswered questions (Hughes *et al.*, 2016). In the present study, we aimed to determine the prevalence and factors associated with HIP using HbA<sub>1c</sub> in Uganda. To our knowledge, this is the first study to report on the use of HbA<sub>1c</sub> to screen for HIP in Uganda. In our study, the prevalence of HIP ranged between 7.5 and 16.0%. This is consistent with the estimated prevalence of HIP of 16% reported in the Africa region (Guariguata *et al.*, 2014). In two previously published studies from Uganda, the prevalence of HIP was 15.6% using FPG criteria (Kiiza *et al.*, 2020) and 31.9% using OGTT criteria (Nakabuye *et al.*, 2017). The observed differences in the prevalence of HIP across these studies could be due to the difference in the diagnostic criteria used. It is well established that due to physiological changes in pregnancy, HbA<sub>1c</sub> level decreases as gestation age increases (Kumpatla *et al.*, 2013; Rafat & Ahmad, 2012; Schaible *et al.*, 2018). This could explain the low prevalence observed in our study.

HIP is typically diagnosed between 24<sup>th</sup> and 28<sup>th</sup> weeks of gestation (WHO, 2018). However, evidence from the metacentric landmark trial, hyperglycemia and adverse pregnancy outcome (HAPO) showed that continued exposure to hyperglycemia



**Figure 2.** Random blood sugar levels across different glycemic status.



**Figure 3.** Receiver operating characteristic curve.

non-diagnostic for diabetes was associated with adverse maternal and fetal outcomes (Catalano *et al.*, 2012; HAPO Study Cooperative Research Group *et al.*, 2008). Based on this finding, current guidelines recommend early screening and appropriate management of HIP to improve maternal and fetal outcomes (WHO, 2018). OGTT is generally considered the gold standard for screening for HIP (Coetzee *et al.*, 2020). However, FPG and HbA<sub>1c</sub> can also be used. HbA<sub>1c</sub> has been shown to correlate with poor maternal-fetal outcomes (Ho *et al.*, 2017). Studies to establish normal HbA<sub>1c</sub> reference ranges in pregnancy are scarce. Among healthy non-diabetic pregnant women, a recent study from Mexico has shown that the upper limit of HbA<sub>1c</sub> increases with gestation age (Sánchez-González *et al.*, 2018). In this population, the cutoff for the diagnosis of HIP was nearly identical to the American Diabetes Association criteria for the diagnosis of diabetes and prediabetes in non-pregnant population and in early pregnancy. However, given the ease of HbA<sub>1c</sub> compared to OGTT, testing may improve follow-up rates and combining HbA<sub>1c</sub> analysis with FPG or waist circumference may improve detection rates (Hughes *et al.*, 2016).

Risk factors for HIP include advancing age; obesity; excessive weight gain during pregnancy; a family history of diabetes; HIP during a previous pregnancy; a history of stillbirth or infant with congenital abnormality; and glycosuria during pregnancy significantly overlap with those of type 2 diabetes mellitus (Diabetes Canada Clinical Practice Guidelines Expert Committee *et al.*, 2018; Farrar, 2016; Saravanan *et al.*, 2020). In our study, the majority of the patients were young, and family history of diabetes was only elicited in 20% of the patients. This is consistent with published studies in Uganda and elsewhere that have reported that over one-third to half of mothers do not have known risk factors (Nakabuye *et al.*, 2017; Thacker & Petkewicz, 2009). This has implications for the selection of patients for screening. RBS has been studied as a possible screening tool. It is interesting that none of the patients with DIP in our study had RBS above 11.1 mmol/L. In one study conducted in Nigeria, using OGTT as a reference standard, the best threshold for screening was 5.4 mmol/L for RBS, which had a sensitivity of 45% and a specificity of 90.0% (Adefisan *et al.*, 2020). In our study, with a cutoff of 4.75mmol/L, we found the reverse, a higher sensitivity (90.7%) and a lower specificity (56.4%). However, at a cutoff of 5.5 mmol/L, we found a similar diagnostic performance (sensitivity of 27% and specificity of 84%). Given the high sensitivity of RBS at a relatively lower RBS cutoff and the cost of performing HbA<sub>1c</sub> especially in LMICs, RBS – a cheap and readily available modality – may be used alongside HbA<sub>1c</sub> for screening for HIP in our setting.

Our study is not without limitations. Firstly, we had a small sample size derived from a single center and thus our findings may not be generalizable to the general population of pregnant women in Uganda. Secondly, there is no established HbA<sub>1c</sub> reference ranges among Ugandan women stratified by gestation age. It is therefore likely that we may have missed some cases of HIP since we used HbA<sub>1c</sub> cutoff for non-/early pregnancy population. However, we excluded anemic patients by performing hemoglobin estimation for all mothers. Lastly, being a pilot study, we were unable to retrieve key risk factors, such as

pre-pregnancy weight, and birth weights and perinatal outcomes of previous pregnancies. However, over 60% of the mothers were primigravida. However, the strength of this study lies in its inclusiveness of pregnant mothers of different gestation ages from both urban and peri-urban communities. We report for the first time the feasibility of screening for HIP using HbA<sub>1c</sub> in a resource limited setting and the utility of RBS as an adjunct to HbA<sub>1c</sub> to aid in identifying of mothers who are likely to have HIP.

## Conclusions

In conclusion, we found a slightly over 10% prevalence of HIP among Uganda women all ages of gestation. The majority of those diagnosed with HIP were young without identifiable risk factors for hyperglycemia. RBS and HbA<sub>1c</sub> may be used complementarily to diagnose HIP in resource constrained settings. We recommend a larger, multicenter study using different diagnostic modalities to confirm of findings.

## Data availability

### Underlying data

Figshare: Hyperglycemia in pregnancy diagnosed using glycated hemoglobin (HbA<sub>1c</sub>) in Uganda: a preliminary cross-sectional report dataset, <https://doi.org/10.6084/m9.figshare.13292690.v1> (Bongomin, 2020).

Data are available under the terms of the [Creative Commons Zero “No rights reserved” data waiver](#) (CC0 1.0 Public domain dedication).

## Acknowledgements

We appreciate the support of Prof. Ally Prebtani through the Rainier Arnhold Senior House Officer Teaching Support (RASHOTS) Project to Dr. Felix Bongomin. We would like to thank Sarah Apoto for assisting with data entry.

## References

- Adefisan AS, Olagbuji BN, Adeniyi AA, *et al.*: **Diagnostic accuracy of random plasma glucose and random blood capillary glucose in detecting international association of diabetes and pregnancy study groups- defined hyperglycemia in early pregnancy.** *Niger J Clin Pract.* 2020; **23**(8): 1087–1094. [PubMed Abstract](#) | [Publisher Full Text](#)
- Bianco ME, Josefson JL: **Hyperglycemia During Pregnancy and Long-Term Offspring Outcomes.** *Curr Diab Rep.* 2019; **19**(12): 143. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Bongomin F: **Hyperglycemia in pregnancy diagnosed using glycated hemoglobin (HbA<sub>1c</sub>) in Uganda: a preliminary cross-sectional report dataset.** *figshare.* Dataset. 2020. <http://www.doi.org/10.6084/m9.figshare.13292690.v1>
- Catalano PM, McIntyre HD, Cruickshank JK, *et al.*: **The hyperglycemia and adverse pregnancy outcome study: associations of GDM and obesity with pregnancy outcomes.** *Diabetes Care.* 2012; **35**(4): 780–6. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Coetzee A, Sadhai N, Mason D, *et al.*: **Evidence to support the classification of hyperglycemia first detected in pregnancy to predict diabetes 6–12 weeks postpartum: A single center cohort study.** *Diabetes Res Clin Pract.* 2020; **169**: 108421. [PubMed Abstract](#) | [Publisher Full Text](#)
- Cosson E, Vicaut E, Sandre-Banon D, *et al.*: **Performance of a selective screening strategy for diagnosis of hyperglycaemia in pregnancy as defined by IADPSG/WHO criteria.** *Diabetes Metab.* 2019; **46**(4): 311–18. [PubMed Abstract](#) | [Publisher Full Text](#)
- Diabetes Canada Clinical Practice Guidelines Expert Committee, Feig DS, Berger H, *et al.*: **Diabetes and Pregnancy.** *Can J Diabetes.* 2018; **42** Suppl 1: S255–282. [PubMed Abstract](#) | [Publisher Full Text](#)
- Farrar D: **Hyperglycemia in pregnancy: prevalence, impact, and management challenges.** *Int J Womens Health.* 2016; **8**: 519–27. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Goyal A, Gupta Y, Singla R, *et al.*: **American Diabetes Association “Standards of Medical Care-2020 for Gestational Diabetes Mellitus”: A Critical Appraisal.** *Diabetes Ther.* 2020; **11**(8): 1639–44. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Guariguata L, Linnenkamp U, Beagley J, *et al.*: **Global estimates of the prevalence of hyperglycaemia in pregnancy.** *Diabetes Res Clin Pract.* 2014; **103**(2): 176–85.

[PubMed Abstract](#) | [Publisher Full Text](#)

HAPO Study Cooperative Research Group, Metzger BE, Lowe LP, *et al.*: **Hyperglycemia and adverse pregnancy outcomes.** *N Engl J Med.* 2008; **358**(19): 1991–2002.

[PubMed Abstract](#) | [Publisher Full Text](#)

Ho YR, Wang P, Lu MC, *et al.*: **Associations of mid-pregnancy HbA1c with gestational diabetes and risk of adverse pregnancy outcomes in high-risk Taiwanese women.** *PLoS One.* 2017; **12**(5): e0177563.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Hughes RCE, Rowan J, Florkowski CM: **Is There a Role for HbA1c in Pregnancy?** *Curr Diab Rep.* 2016; **16**(1): 5.

[PubMed Abstract](#) | [Publisher Full Text](#)

Jiwani A, Marseille E, Lohse N, *et al.*: **Gestational diabetes mellitus: results from a survey of country prevalence and practices.** *J Matern Fetal Neonatal Med.* 2012; **25**(6): 600–610.

[PubMed Abstract](#) | [Publisher Full Text](#)

Kiiza F, Kayibanda D, Tumushabe P, *et al.*: **Frequency and Factors Associated with Hyperglycaemia First Detected during Pregnancy at Itojo General Hospital, South Western Uganda: A Cross-Sectional Study.** *J Diabetes Res.* 2020; **2020**: 4860958.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Kish L: **Survey Sampling.** In: *Syst Biol.* 1965; **46**: 643.

Kumpatla S, Aravindalochanan V, Rajan R, *et al.*: **Evaluation of performance of A1c and FPG tests for screening newly diagnosed diabetes defined by an OGTT among tuberculosis patients—a study from India.** *Diabetes Res Clin Pract.* 2013; **102**(1): 60–64.

[PubMed Abstract](#) | [Publisher Full Text](#)

Meek CL, Lindsay RS, Scott EM, *et al.*: **Approaches to screening for hyperglycaemia in pregnant women during and after the COVID-19 pandemic.** *Diabet Med.* 2020; **4**: e14380.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Mukuve A, Noorani M, Sendagire I, *et al.*: **Magnitude of screening for gestational**

**diabetes mellitus in an urban setting in Tanzania; a cross-sectional analytic study.** *BMC Pregnancy Childbirth.* 2020; **20**(1): 418.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Nakabuye B, Bahendeka S, Byaruhanga R: **Prevalence of hyperglycaemia first detected during pregnancy and subsequent obstetric outcomes at St. Francis Hospital Nsambya.** *BMC Res Notes.* 2017; **10**(1): 174.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Rafat D, Ahmad J: **HbA1c in pregnancy.** *Diabetes Metab Syndr.* 2012; **6**(1): 59–64.

[PubMed Abstract](#) | [Publisher Full Text](#)

Sánchez-González CM, Castillo-Mora A, Alvarado-Maldonado IN, *et al.*: **Reference intervals for hemoglobin A1c (HbA1c) in healthy Mexican pregnant women: a cross-sectional study.** *BMC Pregnancy Childbirth.* 2018; **18**(1): 424.

[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

Saravanan P, Diabetes in Pregnancy Working Group, Maternal Medicine Clinical Study Group, *et al.*: **Gestational diabetes: opportunities for improving maternal and child health.** *Lancet Diabetes Endocrinol.* 2020; **8**(9): 793–800.

[PubMed Abstract](#) | [Publisher Full Text](#)

Schaible B, Calhoun BC, Bush S, *et al.*: **Hemoglobin A1c as a Screening Strategy for Gestational Diabetes.** *Medical and Dental Research.* 2018; **1**(1): 1–4.

[Publisher Full Text](#)

Thacker SM, Petkewicz KA: **Gestational Diabetes Mellitus.** *US Pharm.* 2009; **34**(9): 43–48.

[Reference Source](#)

WHO: **WHO Recommendation on The Diagnosis of Gestational Diabetes in Pregnancy.** Who. 2018.

[Reference Source](#)

World Health Organisation: **Diagnostic Criteria and Classification of Hyperglycaemia First Detected in Pregnancy.** WHO/NHM/NM.13.2 Ed. Geneva: World Health Organization." World Health Organisation, Geneva, Switzerland. 2013.

[Reference Source](#)

World Health Organization: **Haemoglobin Concentrations for the Diagnosis of Anaemia and Assessment of Severity.** World Health Organisation, Geneva, Switzerland. 2011.

[Reference Source](#)

# Open Peer Review

Current Peer Review Status: 

---

## Version 1

Reviewer Report 07 June 2021

<https://doi.org/10.21956/emeraldopenres.15099.r27510>

© 2021 Coetzee A. This is an open access peer review report distributed under the terms of the [Creative Commons Attribution License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

 **Ankia Coetzee** 

Division of Endocrinology, Stellenbosch University and Tygerberg Hospital, Stellenbosch, South Africa

Thanks for the invitation to review the article entitled "Hyperglycemia in pregnancy diagnosed using glycated hemoglobin in Uganda: a preliminary cross-sectional report."

The title is representative of the study and the abstract satisfactory. The introduction is scientifically accurate and meaningful. Study aims are clearly communicated and addressed appropriately with good quality methods. The authors use the HbA1c as the gold standard for the diagnosis of hyperglycemia in pregnancy and derive the RBS as a predictor of HIP with HbA1c as the reference. HIP is diagnosed in 20 pregnant women, using WHO criteria (validated for prediabetes outside of pregnancy). Five women have pre-existing/overt Diabetes Mellitus (HbA1c>6.5%) and the 20 "prediabetes" and 5 women with diabetes are categorized as HIP.

### Comments:

It is unconventional to use HbA1c as the gold standard considering that, compared to an OGTT, HbA<sub>1c</sub> has less evidence to support its use. This should be clearly stated and added to the limitations section.

The rationale for the inclusion of women in the first trimester in this context is also unclear. It has been suggested that RBS (plasma) performs surprisingly well as a first-trimester predictive tool for later GDM diagnosis, but this was not the aim of this study.

Diagnosis of hyperglycemia in pregnancy:

- The authors state that venous blood was used for the glucometer whereas capillary blood is usually used for point of care testing on a glucometer. There is a small but significant difference in the blood glucose results analyzed on a bedside glucometer when the samples are taken from capillary or venous sources. Can the authors please elaborate on the sampling and on the interchangeability of venous and capillary samples using this device?

Accuracy of RBS:

- How does the specific glucometer used, perform at the RBS levels obtained in this study?

Has the glucometer method been standardized against isotope dilution mass spectrometry (ID/MS)? It has been proposed that a total error <5% with an imprecision [CV] <2% has to be achieved for a glucometer to be used in pregnancy, in view of this, is the glucometer used in this study fit for purpose?

Accuracy of HbA1c

- Was the HbA1C determination an approved International Federation of Clinical Chemistry and Laboratory Medicine method traceable to the DCCT trial?

Can the authors comment on the relationship between HbA1c and RBS in the "prediabetes" in pregnancy category (n=20) >24 weeks gestation? This would be a very worthwhile addition to the current study considering this is the group that one might forego OGTT on the basis of hbA1c and RBS in the future.

**Is the work clearly and accurately presented and does it cite the current literature?**

Yes

**Is the study design appropriate and is the work technically sound?**

Yes

**Are sufficient details of methods and analysis provided to allow replication by others?**

Yes

**If applicable, is the statistical analysis and its interpretation appropriate?**

Yes

**Are all the source data underlying the results available to ensure full reproducibility?**

No source data required

**Are the conclusions drawn adequately supported by the results?**

Yes

**Is the argument information presented in such a way that it can be understood by a non-academic audience?**

Yes

**Does the piece present solutions to actual real world challenges?**

Yes

**Is real-world evidence provided to support any conclusions made?**

Yes

**Could any solutions being offered be effectively implemented in practice?**

Yes

**Competing Interests:** No competing interests were disclosed.

**Reviewer Expertise:** Hyperglycemia first detected in pregnancy

**I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.**

-----