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Translating HbA$_{1c}$ measurements into estimated average glucose values in pregnant women with diabetes

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Abstract

Aims/hypothesis This study aimed to examine the relationship between average glucose levels, assessed by continuous glucose monitoring (CGM), and HbA$_{1c}$ levels in pregnant women with diabetes to determine whether calculations of standard estimated average glucose (eAG) levels from HbA$_{1c}$ measurements are applicable to pregnant women with diabetes.

Methods CGM data from 117 pregnant women (89 women with type 1 diabetes; 28 women with type 2 diabetes) were analysed. Average glucose levels were calculated from 5–7 day CGM profiles (mean 1275 glucose values per profile) and paired with a corresponding (±1 week) HbA$_{1c}$ measure. In total, 688 average glucose–HbA$_{1c}$ pairs were obtained across pregnancy (mean six pairs per participant). Average glucose level was used as the dependent variable in a regression model. Covariates were gestational week, study centre and HbA$_{1c}$.

Results There was a strong association between HbA$_{1c}$ and average glucose values in pregnancy (coefficient 0.67 [95% CI 0.57, 0.78]), i.e. a 1% (11 mmol/mol) difference in HbA$_{1c}$ corresponded to a 0.67 mmol/l difference in average glucose. The random effects model that included gestational week as a curvilinear (quadratic) covariate fitted best, allowing calculation of a pregnancy-specific eAG (PeAG). This showed that an HbA$_{1c}$ of 8.0% (64 mmol/mol) gave a PeAG of 7.4–7.7 mmol/l (depending on gestational week), compared with a standard eAG of 10.2 mmol/l. The PeAG associated with maintaining an HbA$_{1c}$ level of 6.0% (42 mmol/mol) during pregnancy was between 6.4 and 6.7 mmol/l, depending on gestational week.

Conclusions/interpretation The HbA$_{1c}$–average glucose relationship is altered by pregnancy. Routinely generated standard eAG values do not account for this difference between pregnant and non-pregnant individuals and, thus, should not be used during pregnancy. Instead, the PeAG values deduced in the current study are recommended for antenatal clinical care.

Keywords Average glucose · Continuous glucose monitoring · Estimated average glucose · Gestation · HbA$_{1c}$ · Pregnant · Type 1 diabetes · Type 2 diabetes

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Abbreviations

ADAG A1C-Derived Average Glucose study
CGM Continuous glucose monitoring
eAG Estimated average glucose
NICE National Institute for Health and Care Excellence
PeAG Pregnancy-specific estimated average glucose

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Introduction

The relationship between HbA1c and average glucose levels has been explored in many studies, most making use of intermittent capillary blood glucose measurements [1–6]. More recently, intensive longitudinal data from continuous glucose monitoring (CGM) have been used to derive a more accurate picture of how average glucose levels compare with HbA1c over time [7–11]. The A1C-Derived Average Glucose (ADAG) study showed a linear association between CGM-measured average glucose and HbA1c levels in non-pregnant adults. However, HbA1c is considered unreliable for assessing glucose control during pregnancy owing to physiological changes that may be attributed to increased red cell production, shortened red cell life span, reduced red cell affinity for glucose, iron deficiency and iron supplementation [13–17]. This has led to uncertainty over the role of HbA1c for blood glucose assessment in pregnancy [18], with key bodies [19, 20] advising that it should not be used for diagnosing diabetes in pregnancy, and the National Institute for Health and Care Excellence (NICE) in the UK recommending that it should not be routinely used to assess glucose control in pregnancy in women with established diabetes [20]. Furthermore, the relationship between any physiological changes in HbA1c across pregnancy and average glucose levels obtained by CGM is unknown. Despite these limitations, HbA1c is widely used in clinical practice during pregnancy in the UK [21], the USA [22] and internationally [23]. Anecdotal reports also suggest that clinicians and patients are using the standard eAG value, which is reported with HbA1c levels, during pregnancy, despite it being derived from data from non-pregnant adults.

Thus, the aims of this analysis were to: (1) examine the relationship between average glucose levels assessed by CGM and HbA1c levels in pregnancy in women with type 1 and type 2 diabetes; (2) determine if this relationship changes with gestational week during pregnancy; and (3) determine whether the standard eAG calculation that is derived from HbA1c measurements is applicable to pregnant women with diabetes.

Methods

Participants This analysis used data obtained from two previously published studies: one based in the UK (East Anglia) [24] and the second in Denmark (Copenhagen) [25]. Both studies recruited pregnant women with pregestational type 1 or type 2 diabetes to prospective randomised controlled trials that explored the clinical impact of CGM on maternal, fetal and neonatal health outcomes. In the UK, pregnant participants, aged 16–45 years, were recruited from two secondary care diabetes antenatal clinics between 2003 and 2006. In Denmark, pregnant participants, aged 19–43 years, were recruited from one diabetes antenatal clinic between 2009 and 2011. Full details of clinical recruitment procedures (including the exclusion of participants with severe medical or psychological comorbidities) have been described previously [24–26]. A total of 117 participants (49 from England and 68 from Denmark), comprising 89 women with type 1 diabetes and 28 with type 2 diabetes, were included in the present analysis [26].

All participants gave written informed consent. Ethical approval was granted by the Suffolk and Norfolk Local Research Ethics Committee and the Danish National Committee on Biomedical Research Ethics. The Helsinki Declaration and Good Clinical Practice guidelines were adhered to throughout the study.

Antenatal and perinatal care All participants received routine clinical care as per national guidelines. In the UK, this involved antenatal clinic visits every 2–4 weeks, with 4–6 visits including CGM and HbA1c measurements. In Denmark, antenatal clinic visits occurred every 2 weeks, with five study visits at 8, 12, 21, 27 and 33 weeks gestation. These study visits included CGM and HbA1c measurements. CGM profiles were collected over 5–7 days. Both studies used comparable glucose targets to achieve optimum glucose control; in the UK, these were: <5.5 mmol/l before meals, <7.8 mmol/l at 60 min postprandial and <6.7 mmol/l at 120 min postprandial. In Denmark, glucose targets were set at 4.0–6.0 mmol/l before meals, 4.0–8.0 mmol/l at 90 min postprandial and 6.0–8.0 mmol/l before bed.

CGM Continuous glucose monitors were used to record electrochemically measured subcutaneous interstitial glucose concentrations every 5 min, generating 288 measurements per day. Both studies used Medtronic CGM systems (MiniMed, Medtronic, Northridge, CA, USA), with CGM Gold sensors being used in the UK and Guardian REAL-Time CGM with Sof-sensors being used in Denmark. Monitors were calibrated against capillary blood glucose measurements as per the manufacturer’s instructions.

HbA1c Blood samples for HbA1c measurements were obtained regularly throughout pregnancy at both centres. Samples
were analysed locally by assays that were DCCT-aligned and from laboratories with National Glycohemoglobin Standardization Program (NGSP) certification.

**Statistical analysis** Average glucose was calculated as the mean of all glucose values obtained in the 5–7 day CGM profile. The corresponding week of gestation was noted for glucose values and, for analysis, values were paired with the HbA1c values that had been measured within ± 1 week of the CGM profile. Each calculated average glucose value was matched to an individual HbA1c, though women contributed multiple average glucose–HbA1c pairs across their pregnancy. A mixed-effects regression model was therefore used to account for the intra-individual variation with multiple data pairs per woman. In seeking the best-fitting model for the relationship between average glucose and HbA1c, this model included the covariates: gestational age in weeks, HbA1c level and study centre (all centred to their grand mean). The models were explored for linear and curvilinear (squared) relationships, with model fit being assessed using the Akaike information criterion [27], whereby a lower score indicated a better fit of the model. All analyses were conducted in Stata 13, version 13 (StataCorp, College Station, TX, USA).

**Results**

**Relationship between average glucose levels and HbA1c in pregnancy** A total of 688 CGM profiles with a mean of 1275 (range 313–2839) glucose measures per profile were obtained for comparison with 688 HbA1c levels. Each woman contributed an average of six average glucose–HbA1c pairs across their pregnancy, at between 8 and 36 weeks gestation.

Fig. 1 shows the association between average glucose and HbA1c values obtained during pregnancy. A linear regression line, with 95% CI, is fitted to the data points ($r^2 = 19.6\%$; average glucose–HbA1c slope = 0.67 [0.57, 0.78]), showing a strong positive association. This implies that, for these women, on average a 1% (11 mmol/mol) difference in HbA1c corresponded to a 0.67 mmol/l difference in their average glucose levels.

**Determining the best-fitting model to account for how gestational changes in HbA1c influence the average glucose–HbA1c relationship** An intercept-only mixed-effects model was compared with models containing random effects for the slope of the average glucose values in relation to HbA1c levels (Table 1 and electronic supplementary material [ESM] Table 1). As the model containing the random effects of the slope coefficient provided a significant improvement in fit, the random slope was retained. The best-fitting model, with the lowest Akaike information criterion score, was model 5 (see Table 1). This model fitted average glucose to HbA1c, study centre, gestation in weeks (linear) and gestation in weeks squared (curved). Model 6 examined an interaction between HbA1c and gestation in weeks to determine whether the gradient between average glucose and HbA1c changed during pregnancy; the findings showed that it did not. There was also no interaction between HbA1c and study centre, demonstrating that the relationship between average glucose and HbA1c was consistent across the two datasets.

**Deriving a pregnancy-specific eAG** Using the best-fitting curvilinear model, Fig. 2 shows the study mean pregnancy-specific eAG (PeAG) levels changing with gestational week for a range of HbA1c levels. As an example, if the HbA1c is measured at 6.0% (42 mmol/mol) during the 12th week of gestation, the PeAG is 6.7 mmol/l, whereas if it is measured at 36 weeks gestation the PeAG is 6.4 mmol/l. To estimate PeAG at any given week during pregnancy, the following equation can be used:

\[
\text{Glucose (mmol/l)} = 6.78 + [0.04 \times (\text{HbA1c} [\%] - 6.3)] + [0.04 \times (\text{Gestation [weeks]} - 21)] - [0.001 \times (\text{Gestation [weeks]}^2 - 528)]
\]

**Comparison of PeAG with the ADAG-calculated eAG** The eAG derived using the ADAG formula and that derived by our pregnancy-specific equation for a given value of HbA1c are shown in Table 2. If we were to use the ADAG formula, an HbA1c of 6.0% (42 mmol/mol) would equate to an eAG of 7 mmol/l irrespective of the gestational week, whereas it would equate to a lower PeAG (between 6.4 and 6.7 mmol/l depending on gestational week) using our pregnancy-specific equation. This difference is more pronounced at higher levels of HbA1c, where an HbA1c of 8.0% (64 mmol/mol) equates to a PeAG of 7.7 mmol/l (at 12 weeks gestation and a PeAG of 7.4 mmol/l at 36 weeks gestation but, in contrast, using the ADAG formula, the same HbA1c value would equate to an eAG of 10.2 mmol/l throughout gestation [8].

**Discussion**

This is the first study to examine the relationship between average glucose levels obtained by CGM and HbA1c levels during pregnancy in women with diabetes. Our analysis demonstrates a positive linear relationship between average glucose and HbA1c levels, but the slope is shallower than that reported in non-pregnant adults [8]. This validates the use of HbA1c to represent average glucose levels during pregnancy, but suggests that a change in HbA1c during pregnancy reflects a smaller change in average glucose than that assumed using the ADAG model [8, 12]. In addition, while we have shown that the relationship between average glucose and HbA1c is
stable during pregnancy, the absolute mean eAG varies with gestational week. Consequently, HbA1c in pregnancy is associated with a lower eAG than that calculated by ADAG and this difference becomes more marked later in pregnancy. This means that the standard eAG reported with HbA1c is not representative of average glucose levels in pregnancy and should not be used for assessing glucose control in pregnancy. We provide an alternative pregnancy-specific calculation for PeAG based on the observed relationship of HbA1c and average glucose during pregnancy.

The work of the ADAG team has embedded the translation of eAG from HbA1c into routine clinical practice [8, 12]. However, the ADAG analysis deliberately excluded pregnant women because of pregnancy-related physiological changes in HbA1c [8] and, as a result, the routinely derived eAG may not be applicable to this population. HbA1c is known to fall with the physiological changes associated with pregnancy, particularly in early and late pregnancy [13–17]. A strength of our study is that average glucose and HbA1c data were obtained on repeated occasions (a mean of six times) in the same woman throughout pregnancy, enabling us to take account of gestational week in our data analysis. This revealed the stability of the average glucose–HbA1c relationship across pregnancy.

While our data confirm that a positive association exists between average glucose and HbA1c, the slope of the relationship was shallower than that seen in non-pregnant adults.

### Table 1

| Model | AIC      | Fixed effects          | Intercept | HbA1c | Other covariates          |
|-------|----------|------------------------|-----------|-------|--------------------------|
| 1     | 1911.58  | Intercept only         | 6.88 (6.70, 7.05) |       |                          |
| 2     | 1762.65  | + HbA1c                | 6.84 (6.69, 7.00) | 0.57 (0.37, 0.77) |                          |
| 3     | 1759.12  | + HbA1c                | 6.79 (6.63, 6.95) | 0.55 (0.35, 0.75) | −0.39 (−0.70, −0.08)    |
| 4     | 1753.49  | + HbA1c + Centre       | 6.77 (6.61, 6.93) | 0.43 (0.22, 0.64) | −0.43 (−0.75, −0.12)    |
|       |          | + Gestation            |           |       | −0.01 (−0.02, −0.00)     |
| 5     | 1750.17  | + HbA1c + Centre + Gestation | 6.78 (6.62, 6.94) | 0.50 (0.28, 0.72) | −0.39 (−0.70, −0.07)    |
|       |          | + Centre + Gestation   |           |       | 0.04 (−0.01, 0.08)       |
|       |          | + Gestation^2          |           |       | −0.001 (−0.002, −0.000)  |
| 6     | 1752.15  | + HbA1c + Centre + Gestation + Gestation^2 | 6.78 (6.62, 6.94) | 0.50 (0.28, 0.72) | −0.39 (−0.70, −0.07)    |
|       |          | + Centre + Gestation + Gestation^2 | |       | 0.03 (−0.01, 0.08)       |
|       |          | + HbA1c × Gestation    |           |       | −0.001 (−0.002, −0.000)  |
|       |          |                        |           |       | 0.00 (−0.01, 0.02)       |

Data shown as regression coefficient (95% CI)

The mixed-effects models were fit between average glucose as the outcome and explanatory variables, using time nested within each mother.

AIC, Akaike information criterion
[7–9], and this gradient remained stable during the last two trimesters of pregnancy. For example, previous data indicate that a 1% (11 mmol/mol) difference in HbA1c is equivalent to a 1.0–2.0 mmol/l difference in average glucose [7–9, 11], whereas our data show that in pregnancy a much smaller difference in average glucose, 0.67 mmol/l, equates to a 1% (11 mmol/mol) difference in HbA1c. This suggests that a change in HbA1c during pregnancy reflects a smaller change in average glucose compared with that seen outside of pregnancy.

Having established that the gradient between the average glucose–HbA1c relationship is stable from the first trimester in pregnancy, there are nevertheless challenges when translating HbA1c levels to eAG values, given the physiological changes that occur in pregnancy. The implication of a fall in HbA1c or average glucose levels as pregnancy progresses means that any fluctuations in either become more sensitive to the gradient relationship. We have shown that gestational week is an important factor to account for when calculating an eAG from the average glucose–HbA1c relationship during pregnancy, since mean levels of average glucose vary throughout pregnancy. Using our best-fitting model, the HbA1c during pregnancy translates to a PeAG that is of a magnitude of 0.5–2.8 mmol/l difference compared with the eAG obtained using the ADAG formula that laboratories report [8, 12], and this difference is more pronounced at higher levels of HbA1c (Table 2). This means that pregnant women and their clinicians could be misled by the standard eAG readings currently generated for laboratory reports and by automated online calculators. Furthermore, many glucose-monitoring devices generate an estimated HbA1c from average glucose data. It is likely that this HbA1c estimation is currently based on the ADAG formula, which may also be unintentionally misleading during pregnancy.

We consider that our analysis performed in pregnant women builds substantially on the ADAG team’s work. It is important, however, to note that while there are similarities, there are also several differences between our analysis and that of the ADAG. In contrast to the prospectively designed ADAG study [8], ours and other studies [9, 11] were pragmatic and made use of existing clinical data obtained from other studies. Compared with the ADAG study, which used 507 participants, of whom 427 had diabetes [8], our study is relatively small and we recognise that a larger study would help to improve the precision of our model to more confidently ascertain the relationship between average glucose and HbA1c levels. In addition, the ADAG study included participants with a greater range of HbA1c levels, including many with far higher HbA1c values than the participants in our study.

The ADAG study paired the average glucose measures obtained by intermittent CGM readings taken for 2–3 days, every 4 weeks over a period of 3 months (giving ~2500 glucose values per participant) to an HbA1c taken at the end of the 3 months measurement period [8], resulting in one average glucose–HbA1c pair per participant. In contrast, to address the complex issue of gestational physiological changes in HbA1c, the average glucose values obtained in our study were derived from an individual CGM session of 5–7 days, (giving a mean of 1275 glucose values), and were compared with an HbA1c value taken within ± 1 week of the CGM profile, yielding a mean of six average glucose–HbA1c pairs per participant. Previous small studies conducted in the 1980s used capillary blood glucose testing to calculate average glucose in pregnancy and showed a strong positive correlation between HbA1c and the preceding 8–12 weeks’ average glucose values [28, 29]. We obtained the strongest relationship between average glucose and HbA1c when both were measured within a few weeks of each other (ESM Table 2), suggesting that in pregnancy an HbA1c value is more reflective of current average glucose readings (obtained by CGM) than those obtained

| HbA1c % (mmol/mol) | ADAG eAG mmol/l | PeAG mmol/l |
|-------------------|-----------------|-------------|
|                   | 12 weeks gestation | 24 weeks gestation | 36 weeks gestation |
| 5.0 (31)         | 5.4             | 6.2         | 6.2         | 5.9         |
| 6.0 (42)         | 7.0             | 6.7         | 6.7         | 6.4         |
| 7.0 (53)         | 8.6             | 7.2         | 7.2         | 6.9         |
| 8.0 (64)         | 10.2            | 7.7         | 7.7         | 7.4         |
over the preceding 3 months. Anecdotally, it is very common to see dramatic reductions in HbA1c over very short periods of time (<4 weeks) at the start of pregnancy as women are motivated to rapidly optimise their glucose control upon finding out that they are pregnant; this may account for this more proximal relationship.

The ADAG study chose to weight their analysis with intermittent daytime capillary glucose readings but we did not. The rationale for weighting their analysis is unclear as: (1) CGM is already calibrated with regular capillary glucose readings; and (2) intermittent capillary glucose readings do not represent the ‘true’ average glucose value across the 24 h day since they are intermittent, ignore overnight glucose levels and may be skewed by postprandial glucose excursions. Since the ADAG analysis found that the average glucose–HbA1c relationship was unchanged if only CGM readings were used for analysis, we decided to adopt this approach and not weight our analysis [8].

Our data have some further limitations; the women in our study were predominantly white European, which may limit applicability of our findings to women from other cultures and backgrounds. Our analysis did not include any women with gestational diabetes, so care needs to be taken with regard to its applicability in this context. We did not have data on haematocrit levels or iron deficiency/supplementation in our participants but, given that these are factors in the physiological changes of HbA1c during pregnancy, this information might be useful to include in any future analysis of HbA1c and average glucose levels in pregnancy.

We know from population-based studies that HbA1c in pregnancy is a useful guide for pregnancy outcome and risk stratification [30] and is recommended by NICE for this purpose [20]. The ADA recommends regular assessment of glucose control during pregnancy, using monthly HbA1c, to maintain a level of 6.0–6.5% (42–48 mmol/mol) [22]; however, NICE was unable to make this recommendation because of a lack of data for validation of the relationship between HbA1c to average glucose levels during pregnancy [20]. Our current analysis now provides this validation.

CGM is increasingly being used in clinical practice. The average glucose level calculated from the intensive longitudinal glucose data on these devices is far superior to that obtained by capillary glucose meters. Increasing the accessibility and use of CGM as an alternative to capillary glucose testing may significantly improve glucose management during pregnancy [24–26]. One of the difficulties of using CGM in pregnancy is determining exactly which aspects of glucose control to target. Targeting weekly PeAG could be a simple way to help women achieve the glucose control necessary to maintain their HbA1c at ‘low risk’ levels across pregnancy. Our data would suggest that maintaining a PeAG of 6.4–6.7 mmol/l throughout pregnancy should achieve an HbA1c of 6.0% (42 mmol/mol), which is necessary for reducing the risk of adverse pregnancy outcomes.

In summary, HbA1c can be translated to eAG values in pregnant women with diabetes, but these are not the same as those commonly reported. Therefore, pregnancy-specific values, PeAG, are recommended for use in antenatal clinical care.

Data availability Data from the English study is available on request from HRM (Helen.Murphy@uae.ac.uk). Data from the Danish study is available on request from ERM (elisabeth.reinhardt.mathiesen@regionh.dk).

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Duality of interest The authors confirm that there is no duality of interest associated with this manuscript.

Contribution statement RB, EMS and HRM conceived and designed the study, HRM and RT designed the English study and contributed to data acquisition. ERM and ALS designed the Danish study and contributed to data acquisition. GRL, MSG and EMS analysed the data. All authors interpreted the data. GRL and EMS drafted the initial paper prior to it being critically revised by all authors. All authors approved the final version of the article to be published GRL had full access to all of the data in the study, HRM to the English data and ERM to the Danish data. GRL, HRM and ERM are the guarantors of this work and take responsibility for the integrity of the data and the accuracy of the data analysis.

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