A comparison of follow-up rates of women with gestational diabetes before and after the updated National Institute for Health and Care Excellence guidance advocating routine follow-up, and the association with neighbourhood deprivation

SEBASTIAN WALSH,1 MAHMOUD MAHMOUD,2 HTWE HTUN,3 SHEENA HODGETT,4 DAVID BARTON5

Abstract
Background: Gestational diabetes mellitus (GDM) occurs in one in every 23 UK pregnancies. GDM identifies the mother as high-risk for development of type 2 diabetes. The National Institute for Health and Care Excellence (NICE) published updated guidance in February 2015 recommending routine follow-up of women with GDM.

Aims: This cohort study compared follow-up rates of women with GDM before and after the updated guidance. We also investigated for an association between follow-up rates and deprivation.

Methods: Participants were identified from the database of the GDM service of two English hospitals and were organised into two cohorts: 'pre-guidance' (2012–2015) and 'post-guidance' (2015–2016). Using the recommendations of the NICE guidance as the follow-up standard, we used the hospitals’ computer system to compare follow-up rates of the two cohorts. The English Indices of Deprivation split the country into 32,844 small areas and rank them in order of deprivation such that 1 is the most deprived area and 32,844 is the least deprived. We compared the patients’ postcodes against the English Indices of Deprivation to investigate the relative levels of neighbourhood deprivation of those followed up compared with those not followed up. The Z statistic was used to test for statistical significance.

Results: 535 participants were included (pre-guidance n=306, post-guidance n=229). Baseline average age (pre-guidance 32.2 years, post-guidance 32.5 years), body mass index (30.7 kg/m², 30.9 kg/m²) and fasting glucose (4.9 mol/L, 4.8 mol/L) were all comparable between cohorts. The follow-up rate improved from 60.5% in the pre-guidance group to 69.9% in the post-guidance group. The median deprivation rank of those followed up was 14,565 compared with 13,393 in those not followed up. This difference was not found to be significant.

Conclusion: A higher proportion of women with GDM were followed up with screening for type 2 diabetes after the updated NICE guidance in 2015 recommended routine follow-up. Across the study, over a third of women were not followed up. There was no statistically significant difference in the deprivation levels of those women followed up compared with those not followed up.

Key words: diabetes, gestational, cohort studies, health services research, follow-up studies, health equity

Introduction
Gestational diabetes mellitus (GDM) occurs when women not previously known to have diabetes develop hyperglycaemia during pregnancy. The prevalence of GDM in the UK is estimated to be one in every 23 pregnancies.1 Studies suggest that women diagnosed with GDM have a 41% risk of having GDM again in subsequent pregnancies,2 and up to a 60% lifetime risk of developing type 2 diabetes.3 The presence of GDM increases the risk of fetal macrosomia (and the associated obstetric risks of shoulder dystocia and maternal morbidity from caesarean
The National Institute of Health and Care Excellence (NICE) published guidance in February 2015 entitled ‘Diabetes in pregnancy: management from preconception to the postnatal period’. It advised that women diagnosed with GDM, whose blood glucose levels returned to normal after delivery, should be followed up with either a fasting blood glucose (FBG) test after 6 weeks or by measuring glycosylated haemoglobin levels (HbA1c) after 12 weeks. Despite the prevalence and implications of GDM, no nationalised standard for how to follow up women diagnosed with GDM existed prior to the revised guidance in 2015, and follow-up was carried out according to local policies.

The primary research question of this study was: did the rate of follow-up of women diagnosed with GDM change after the updated NICE guidance in 2015? The secondary research question was: was the rate of follow-up associated with deprivation?

Methods
This was a cohort study using data collected from two district general hospitals in England. Shrewsbury and Telford Hospital NHS Trust (SaTH) operates across two distinct hospital sites: The Royal Shrewsbury Hospital and The Princess Royal Hospital Telford, but both hospitals are administered by one NHS Trust and served by a solitary staff body. The trust is the main provider of district general hospital services for nearly half a million people in Shropshire, Telford & Wrekin and mid Wales. The population served is predominantly rural with between 90% and 98% from the ‘white’ ethnic group. The GDM service at SaTH has routinely collected data on women registered with the service since 2012.

There has been a GDM policy in place at SaTH since 2010. The policy was initially based upon the 2008 NICE guidance, and then updated accordingly after the publication of the 2015 guidance. As per the NICE guidance, diagnosis was made based on the results of an oral glucose tolerance test (OGTT) with a FBG level of ≥5.6 mmol/L (≥7.0 mmol/L before the updated guidance) or 2-hour plasma glucose level of ≥7.8 mmol/L. The 2008 guidance did not contain recommendations for follow-up; however, the SaTH policy was to follow up patients treated with dietary advice only with a 6-week postnatal FBG test requested from the GP in the discharge paperwork, and those treated with oral agents or insulin with an OGTT organised by the GDM service itself. There was no change to the SaTH follow-up policy after the publication of the 2015 guidance.

We combined data collected by the GDM service (demographics, reason for having the diagnostic OGTT and treatment for GDM) with data from the hospital computer system (delivery date, results and timing of any follow-up tests performed). Women were included if they had been registered with the GDM service during or after 2012 until the end of 2016 (this was to allow for the follow-up period to have elapsed). Participants were excluded if the dataset was not sufficiently complete to be confident of whether or not follow-up had occurred (eg, hospital number not recorded, delivery date not identifiable from computer system).

Participants were split into two cohorts, those who delivered on or after 1 January 2012 until before the updated NICE guidance was implemented at SaTH on 1 April 2015 and those who gave birth on or after this date until 31 December 2016. Those who delivered before implementation of the NICE guidance were identified as the ‘pre-guidance’ cohort and those who delivered after the guidance was implemented were identified as the ‘post-guidance’ cohort.

The outcome measure was the success or failure of follow-up of women according to the 2015 NICE guidance. Participants were deemed to have been followed up appropriately if they had either a FBG test performed 6–13 weeks post-delivery or a HbA1c blood test performed 13–24 weeks post-delivery. NICE guidance does not actually specify a maximum timeframe for follow-up to occur, merely that it should. Maximum timeframes used in other studies ranged from 12 weeks to several years. We used 24 weeks as the cut-off as a balance between allowing sufficient time for follow-up to occur against the pragmatism of needing the follow-up time period to have elapsed before performing the study.

The English Indices of Deprivation are a measure of relative deprivation in small areas in England called lower-layer super output areas (LSOA) published by the UK government. The indices are a composite measure calculated by collating data on income, employment, education, health and disability, crime, barriers to housing and services, housing quality and the environment. There are 32,844 LSOAs in total and they are ranked such that ‘1’ is the most deprived area and the area ranked ‘32,844’ is the least deprived. By inputting the included women’s postcodes into the interactive web tool (published alongside the index), we were able to compare the average index rank of women who were followed up successfully with those who were lost to follow-up, and thus create a proxy correlation of follow-up rates with deprivation. A statistical comparison of the datasets was undertaken on Microsoft Excel using the Z test.

Results
Baseline characteristics
The baseline characteristics of the two cohorts are shown in Table 1. Reasons for screening were also similar between cohorts, with the most common reasons for performing an OGTT being: first degree relative with diabetes (n=109, n=75), body mass index >30 kg/m² (n=64, n=62), and ultrasound estimate of fetal growth above the 90th customised centile for gestation (n=54, n=50).

In the pre-guidance cohort, 9.2% of participants were treated with metformin as a sole agent, 2.9% were treated with insulin as a sole agent, and 3.9% were treated with a combination of both. In comparison, in the post-guidance cohort, more women were treated with insulin (17.5%) compared to only 4.8% with metformin and 4.8% with a combination of both.
In the pre-guidance cohort, 57.5% of women were followed up. Outcome cases in the post-guidance cohort. This occurred in only nine cases in the pre-guidance cohort and seven by use of a FBG test and 6.5% were followed up by HbA1c, giving a total of 69.9% of the cohort successfully treated by NICE (FBG 16.0 17.5 % treated with combination 3.9 4.8 % treated with insulin only 2.9 17.5 % treated with metformin only 9.2 4.8 Mean fasting glucose (mmol/L) 4.9 4.8 Number of women 306 229 Mean age (years) 32.2 32.5 Mean BMI (kg/m²) 30.7 30.9 % treated with any drug 16.0 27.1

Table 1 Baseline characteristics of the pre- and post-guidance cohorts

However, the average FBG reading at diagnosis did not differ significantly between the cohorts (4.9 mmol/L and 4.8 mmol/L, respectively), so it is likely that this discrepancy in treatment frequencies represents shifting therapeutic approaches to treating GDM rather than any intrinsic differences between the cohorts.

We excluded participants from the analysis if we were unable to elicit the delivery date from the computer system. This occurred in only nine cases in the pre-guidance cohort and seven cases in the post-guidance cohort.

Outcome
In the pre-guidance cohort, 57.5% of women were followed up by use of a FBG test and 6.5% were followed up by HbA1c. Accounting for women who were followed up with both methods, a total of 60.5% of the pre-guidance cohort were followed up successfully. In contrast, in the post-guidance cohort 62.0% of women were followed up with a FBG test and 17.5% with HbA1c, giving a total of 69.9% of the cohort successfully followed up.

A total of 2.8% of the women followed up were found to have FBG readings or HbA1c results in the diabetic range as stipulated by NICE (FBG ≥7 mmol/L, HbA1c ≥48 mmol/mol (6.5%)). In the pre-guidance cohort the rate of diabetes at follow-up was 2.6% compared with 3.1% in the post-guidance cohort.

The rate of follow-up was not strongly associated with baseline FBG (60.7% follow-up rate amongst women with baseline FBG ≥5.6 mmol/L compared with 57.4% follow-up rate amongst women with baseline FBG <5.6 mmol/L). There was, however, a strong association between type of GDM treatment and the incidence of follow-up: 80.2% of women treated with either metformin or insulin were followed up compared with only 60.1% of women who were treated with dietary advice alone.

Deprivation
The median LSOA deprivation rank of those women followed up successfully was 14,565 compared with 13,393 for those women not followed up successfully. Forty-six participants had to be excluded from this analysis because they lived in Wales and the Index is produced separately for England and Wales. We tested for statistical significance using the Z test, with a threshold of p<0.05 to reject the null hypothesis of no significant difference between the two datasets. We found a p value of 0.16, which meant there was insufficient evidence to reject the null hypothesis. We therefore conclude that the observed association between deprivation and being lost to follow-up was not statistically significant.

Discussion
Main findings
In this study of follow-up rates among 535 women with GDM at two district general hospitals in England, the rate of follow-up improved from 60.5% to 69.9% after the publication of updated NICE guidance. There was no statistically significant relationship between neighbourhood deprivation and follow-up rates.

Strengths and limitations
There are several strengths of this study. We included a large number of participants and, to the best of our knowledge, they represented the entire GDM population of the trust during the study period. This, coupled with the very small number of participants with missing data, limits the potential for bias.

The participants included in this study were sourced exclusively from the records of the GDM team at SaTH, who aim to record data from all women registered with the service. We were not able to externally confirm that this represented the entire population and it is possible that there are women who were missed from registration on the database – for example, because of administrative errors. However, this is likely to represent a very small number of participants, if any.

The main limitation of this study was that it was performed using data from only one NHS trust. Although women from two hospitals were included, a single GDM multidisciplinary team serves both hospitals, limiting the generalisability of the findings.

Interpretation of findings in relation to previously published work
To the best of our knowledge, this is the first study to investigate how the follow-up rate of women with GDM has changed following the updated NICE guidance. This, coupled with the limited generalisability of the study setting, limits the external validity of the results.

This is an observational study and can therefore not be used to infer a causal process, rather than to observe a correlation. We must therefore consider the possibility of confounding factors. It is possible that another factor, other than the publication of NICE guidance, was responsible for the observed improvement in the follow-up rates. For example, it is possible that the mere collection of participants’ details in the database by the GDM team (commenced in 2012) leads itself to more vigilance around follow-up. Indeed, the follow-up rate of >60% is generally higher than that reported by previous studies. However,
it should be noted that the team did not actually collect data on who had and had not been followed up.

While there was a small improvement in the proportion of women being followed up by FBG level (pre-guidance 57.5%, post-guidance 62.0%), the majority of the improvement in follow-up rates between the pre- and post-guidance cohorts was a large increase in the use of a HbA1c test to follow up women, from 6.5% to 17.5%. This is likely to be partly because of the incorporation of HbA1c in the updated NICE guidance. It is possible that the convenience of the HbA1c test (involving a one-off blood test rather than the relatively time-consuming OGTT) influenced this finding.

Across the study, women who were treated with insulin or metformin rather than only receiving dietary intervention were far more likely to be followed up (80.2% vs. 60.1%). This could be because those women had more interactions with healthcare professionals during their pregnancy, because the treatment made them more conscious of their diagnosis, or because follow-up was performed by the GDM team itself. The percentage of women treated with any agent increased by 11.1% from the pre-guidance to the post-guidance group (from 16.0% to 27.1%), predominantly due to an increased number of women using insulin. It is likely that this also accounts for some of the increase in follow-up observed between the two cohorts. The increase in women being treated with insulin or metformin in the post-guidance group is likely to be because of a change in the target blood glucose levels in the updated NICE guidance (from therapeutic targets of ≤6.0 mmol/L (fasting) and ≤8.0 mmol/L (2 hour) pre-guidance to post-guidance targets as per NICE 2015 of ≤5.3mmol/L (fasting), ≤7.8 mmol/L (1 hour) and ≤6.4 mmol/L (2 hour)).

Taking the above caveats into account, we observed a large improvement in the rate of follow-up of women with GDM after the publication of updated NICE guidance that advocated routine follow-up. There is no persuasive argument for why this observation should have occurred by chance, because of bias, or by the action of some external factor. Our interpretation of the results is that it is likely that the publication of updated NICE guidance improved the rate of follow-up of women with GDM in this population.

Another interesting finding in the study was the increased rate of diagnosis of GDM as the study progressed. This was evidenced by an increase from 7.8 diagnoses/month in the pre-guidance group to 19.1 diagnoses/month in the post-guidance group. As mean age and body mass index were comparable between the two groups, it is likely that this increase in incidence was a result of the change in diagnostic criteria from a FBG level of ≥7.0 mmol/L (pre-guidance) to ≥5.6 mmol/L (post-guidance).

Implications for future research, policy and practice
Although this study showed an improvement in the rate of follow-up of women with GDM during the study period, over one-third of the participants were not followed up. Given the high incidence of type 2 diabetes after a GDM diagnosis, this leaves a large number of high-risk individuals who are not being actively followed up.

Evidence from this study suggests that offering follow-up with a HbA1c could provide an opportunity to increase postnatal screening rates and future research could test this hypothesis more directly.

This study was performed in a focused study setting and further research should now be undertaken to investigate whether trends observed in this study are true across a wider range of settings.

Conclusion
A higher proportion of women diagnosed with GDM were followed up with screening for type 2 diabetes after the publication of updated NICE guidance in 2015 advocating routine follow-up of these women. Possible explanations for the increased follow-up rate include: the increased proportion of women receiving drug treatment owing to stricter therapeutic targets and the incorporation of HbA1c as a follow-up test in the guidance. However, over a third of women were not followed up and this potentially represents a large number of women across the country at high risk of type 2 diabetes who are not actively being followed up. In our study we found no statistically significant relationship between neighbourhood deprivation and follow-up rates.

Conflict of interest The authors declare that they have no conflicts of interest in relation to this article.

Funding None.

Contribution statement The idea for the study was conceived by DB. SW, ME and HH completed data collection. SW performed the data analysis and drafted the manuscript. DB provided feedback during the conduct of the research. SH and DB commented on the final draft.

Novelty statement To the best of our knowledge, this is the first study which investigates the effect of the publication of updated NICE guidance advocating routine follow-up of women with GDM on follow-up rates in a real-world setting. The study demonstrates that up to one-third of women at high risk of developing type 2 diabetes are not being followed up.

HRA approval Project ID: 247335.

References


---

**Semaglutide Nationwide Audit in progress**

In January 2019 ABCD launched a nationwide audit of Semaglutide in real clinical use in the UK.

**Does your centre use Semaglutide?**

If yes, REGISTER YOUR CENTRE!

http://www.diabetologists-abcd.org.uk/GLP1_Audits/Semaglutide_Audit.htm

- invited to enter your patient data into the bespoke online tool
- you can collect data on the easy-to-complete paper proformas which you can printout from the above web address
- you are able to analyse your local data easily
- the data will be automatically added to the national data in anonymised form

Please remember:
- the more data, the more complete our understanding of **Semaglutide** in real clinical practice
- all contributors will be listed in publications arising from data submission