

Frequent Monitoring of A1C During Pregnancy as a Treatment Tool to Guide Therapy

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OBJECTIVE— No guidelines for A1C measurement exist for women with gestational diabetes mellitus (GDM). The aim of this study was to document the rate of A1C decline in women with GDM.

RESEARCH DESIGN AND METHODS— Women with GDM in the Santa Barbara County Endocrine Clinic are managed with a carbohydrate-restricted diet and self-monitored blood glucose before and 1-h postprandial. Insulin is started if the preprandial glucose concentration is ≥ 90 mg/dl and/or a 1-h postprandial glucose concentration is ≥ 120 mg/dl. Capillary A1C was tested weekly using the DCA2000+ analyzer.

RESULTS— Twenty-four women with GDM (aged 29.0 ± 7.3 years) with initial A1C $\geq 7.0\%$ were recruited. Baseline A1C was $8.8 \pm 1.8\%$. Mean A1C decline was 0.47% per week (range 0.10–1.15%); the maximum was 4.3% in 4 weeks.

CONCLUSIONS— This study documents rapid decline in A1C during pregnancy and the utility of weekly A1C to guide therapy.

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A1C is routinely measured approximately every 3 months in individuals with diabetes to assess the mean glucose concentration. The erythrocyte life span is ~ 120 days (1). Thus, the 3-month interval between tests of A1C reflects the mean blood glucose over the preceding weeks to months. There are no clear guidelines for the frequency of testing A1C during pregnancy (2). In pregnancies complicated by type 1 or type 2 diabetes, most studies report the relationship between the first trimester A1C and the risk of spontaneous abortion and/or congenital malformations (3,4). The goal for therapy in pregestational diabetes is to sustain the A1C at $< 6.0\%$, although this level of A1C assumes that the measurement is performed only once each trimester (5–9). In addition, A1C is not recommended routinely in women with gestational diabetes mellitus (GDM). The

life span of the erythrocyte during pregnancy is shortened to ~ 90 days, and thus the test measures the mean glucose over a shorter time interval than in the nonpregnant state (10). Hence, the rate of change of A1C in pregnancy reflects the glycemic control over the past few weeks. Therefore, the measurement of A1C more frequently during pregnancy may be used to guide therapeutic decisions in all pregnancies complicated by diabetes including GDM. The aim of this study was to document the rate of A1C decline during the first 4 weeks after the initiation of treatment in women with GDM.

RESEARCH DESIGN AND METHODS

This was an observational study of pregnant women attending the Santa Barbara County Health Care Services Obstetrics Clinic for care who have the diagnosis of GDM (diabetes first

diagnosed during pregnancy [2]) and who were referred to the Prenatal Diabetes/Endocrine Clinic for management. Because of the high rates of diabetes in this largely Latino population and because many of the women only seek medical attention during pregnancy, women are screened for GDM very early in pregnancy. Many women found to have diabetes early in pregnancy undoubtedly have preexisting type 2 diabetes, which falls under the classification of GDM when first identified during pregnancy. Treatment consists of a carbohydrate-restricted meal plan (11) and fingerstick blood glucose monitoring before and 1-h after each meal (11). Insulin is initiated if the diet does not achieve premeal glucose concentrations < 90 mg/dl and/or 1-h postprandial glucose concentrations < 120 mg/dl within 1 week of the carbohydrate restriction prescription (12,13). For simplicity in the clinic and to impress upon the diabetic women the importance of tight glucose control, a point of care A1C by fingerstick is routinely obtained at every weekly visit and analyzed immediately using the DCA2000+ analyzer (14,15). The DCA A1C is a Clinical Laboratory Improvement Amendments (CLIA)-waived test with a coefficient of variation of 2.3–3.3% at a normal A1C concentration of 5.2 and of 2.8–3.7% at an elevated A1C concentration of 11.9%.

RESULTS— Twenty-four Latina women with the diagnosis of GDM whose initial A1C was $\geq 7.0\%$ and followed during the first 1–4 weeks of treatment were included in the analysis. The mean \pm SD age was 29.0 ± 7.3 years, A1C at enrollment was 8.80 ± 1.83 , and the duration of follow-up for this study was 3.2 ± 1.0 weeks. Mean gestational age at diagnosis of GDM was 12.2 ± 7.4 weeks.

Figure 1 shows the A1C decline during the first 4 weeks of treatment (range 1.0–4.0 weeks) for all 24 women. The mean decline was $0.47 \pm 0.30\%$ per week, and the maximum decline over the 4 weeks was 4.3%. The decline was greatest among women with the highest A1C at baseline. Among the 20 women with an initial A1C $< 10.0\%$, the decline was

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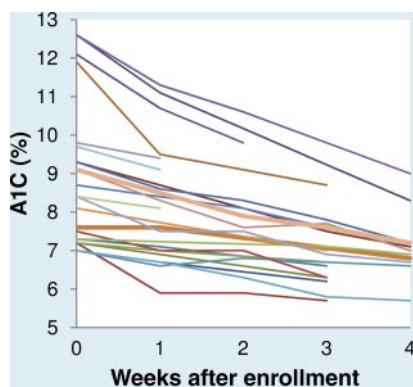


Figure 1—A1C (%) at enrollment and weekly for up to 4 weeks of follow-up in 24 women with GDM.

$0.36 \pm 0.15\%$ per week. After the 1st month of decline, the A1C stabilized with a small drop of only 0.05% per week to the end of gestation (data not shown).

CONCLUSIONS— This study documents that a rapid decline in A1C can be achieved during pregnancy when normoglycemia is vigorously instituted and achieved and thereafter sustained for 4 weeks. Home glucose monitoring on a regular basis is certainly the cornerstone of management of GDM and was initiated in the women enrolled in this study from the beginning. However, many high glucose values can be missed with the usual six or seven fingersticks per day. When A1C is measured at every weekly visit and the result is compared with the woman's previous value, then the rate of change of A1C (decline or rise) can be used to assess the glycemic control and guide therapeutic decisions. A randomized trial in a larger group of women will be needed to determine whether weekly A1C measurements will lead to an improvement in outcome.

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