Impact of introducing the International Association of Diabetes and Pregnancy Study Groups (IADPSG) criteria on pregnancy outcomes in Japan

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Abstract. To estimate pregnancy complications in women newly diagnosed with gestational diabetes mellitus (GDM) according to the new International Association of Diabetes and Pregnancy Study Groups (IADPSG) criteria and verify the validity of introducing the IADPSG criteria in Japan. This retrospective study included data from women with singleton delivery at 22 weeks or later at a tertiary hospital during 2000–2009. We compared pregnancy outcomes between women who would now be diagnosed with GDM according to the IADPSG criteria but not by the old JSOG criteria (IGT group, n = 503) and women with normal glucose tolerance according to both the criteria (NGT group, n = 2,789). Multivariate analysis was performed and adjusted for background factors. Maternal age at delivery and pre-pregnancy BMI were significantly higher in the IGT group than in the NGT group, while gestational weeks at delivery did not differ between the groups. No difference was observed in the rates of GDM-related composite complications (defined as cases with at least one of the following: macrosomia, shoulder dystocia, neonatal hypoglycemia, neonatal hyperbilirubinemia, or neonatal respiratory distress syndrome) at 11.9% and 8.8% (adjusted odds ratio (OR) 1.30, 95% confidence interval (CI) 0.90–1.87, \( p = 0.16 \)). Pregnancy outcomes did not differ significantly between the IGT and NGT groups, except for frequencies of total neonatal admissions at 10.5% and 7.1%, respectively (adjusted OR 1.55, 95% CI 1.12–2.13, \( p < 0.01 \)).

Key words: Gestational diabetes mellitus (GDM), International Association of the Diabetes and Pregnancy Study Group (IADPSG), Pregnancy, Japanese

Based on the results of the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) study reported in 2008 [1], the International Association of Diabetes and Pregnancy Study Groups (IADPSG) defines the diagnosis of gestational diabetes mellitus (GDM) when a 75-g oral glucose tolerance test (OGTT) performed between 24 and 28 weeks of gestation exceeds one of the following: 92 mg/dL (5.1 mmol/L) fasting plasma glucose (FPG), 180 mg/dL (10 mmol/L) 1-hour post-glucose load, or 153 mg/dL (8.5 mmol/L) 2-hour post-glucose load [2]. While many countries have adopted these international diagnostic criteria, the increase in the number of pregnant women diagnosed with GDM [3-9] has placed a heavy financial burden on medical care systems [10, 11]. Moreover, there is a lack of clear evidence that the new criteria reduce the incidence of perinatal adverse events. The American College of Obstetricians and Gynecologists (ACOG) does not support the IADPSG criteria [12].

Prior to the adoption of the IADPSG criteria, Japan had its own diagnostic criteria for GDM from the Japan Society of Obstetrics and Gynecology (JSOG), which required two or more of the following: fasting plasma glucose (FPG) concentration of 100 mg/dL (5.5 mmol/L), 1-h post-glucose load of 180 mg/dL (10 mmol/L), or 2-h post-glucose load exceeding 150 mg/dL (8.3 mmol/L) in a 75-g OGTT. However, the old criteria were based on the National Diabetes Data Group (NDDG) criteria [13] and Carpenter and Coustan (C&C) criteria [14], which means they were intended to predict the onset of postpartum type-2 diabetes. It thus became evident that criteria aimed at preventing perinatal adverse events would be preferable. To achieve this, the diagnostic criteria used in Japan for GDM was changed in 2010 by introducing the IADPSG criteria [15]. As has
been seen in other countries, the adoption of these criteria in Japan has led to a 2.7- to 5.7-fold increase in the prevalence of GDM. However, it remains unclear whether it has contributed to improving maternal and fetal prognoses [9, 16-19].

The objective of this study was to verify the validity of introducing the IADPSG criteria by examining pregnancy outcomes among women who were diagnosed with GDM based on the IADPSG criteria but not with the old JSOG criteria.

**Materials and Methods**

*Study design and subjects*

This single-center retrospective study used our center’s database. Data were collected from an institutional perinatal database and the women’s medical records. The study was approved by the ethics committee of Yokohama City University (No. B180700037).

The subjects were pregnant women who underwent a 75-g OGTT during pregnancy and had a singleton delivery at 22 weeks’ gestation or later at the Perinatal Center for Maternity and Neonate of Yokohama City University Medical Center from January 1, 2000, to December 31, 2009. Following Japanese clinical guidelines [15], women considered to be at high risk of GDM undergo a 75-g OGTT at any time during pregnancy.

Basically, all pregnant women undergo a 2-step screening. Specifically, before 20 weeks of gestation, a 75-g OGTT is performed in women with certain risk factors (family history of diabetes, history of GDM, history of macrosomia [fetal weight ≥4,000 g], history of shoulder dystocia, age ≥35 years, obesity [pre-pregnant BMI ≥30]) and with casual blood glucose concentration ≥95 mg/dL. Then, all women except those with pre-pregnancy DM, overt DM; defined as FPG ≥126 mg/dL (7 mmol/L) or HbA1c ≥6.5%, or already diagnosed with GDM in early pregnancy undergo universal screening with 50-g glucose challenge test (50 g GCT) from 24 to 28 weeks’ gestation. If the glucose challenge test result exceeds an hourly value of 140 mg/dL, a 75-g OGTT is performed.

A 75-g OGTT is also performed in other cases, such as when urinary sugar level ≥2+ is observed in a qualitative urine examination as part of a pregnancy health check, or in the presence of hydramnios or macrosomia (estimated fetal weight ≥4,000 g).

Women with multiple pregnancies, pre-pregnancy DM, or overt DM were excluded. Finally, as subjects of the present study, we chose pregnant women treated as NGT before the introduction of the IADPSG criteria.

We allotted the subjects into two groups. Women previously not diagnosed with GDM under the old JSOG criteria, but who received a diagnosis under the new IADPSG criteria, were placed in the impaired glucose tolerance (IGT) group, while those with normal glucose tolerance (NGT) under both the old JSOG and new IADPSG criteria were placed in the NGT group. Since these two groups had not been diagnosed with GDM under the old JSOG criteria at the time, neither received any intervention for GDM.

Maternal characteristics and pregnancy outcomes were compared between the groups. Data on the following characteristics was gathered from our department database and patient medical records: age (years), height (cm), pre-pregnancy weight (kg), pre-pregnancy body mass index (BMI) (kg/m²), weight at delivery (kg), BMI at delivery (kg/m²), multiparity, gestational age at delivery (weeks), gestational weight gain (GWG) (kg), gestational age at GDM diagnosis (weeks).

The outcomes examined were: infant birth weight (g) and large-for-gestational-age (LGA), small-for-gestational-age (SGA), macrosomia, shoulder dystocia, preeclampsia (PE), total cesarean deliveries, emergency caesarean deliveries, low Apgar score (APS), low umbilical arterial pH (UApH), neonatal hypoglycemia, neonatal hyperbilirubinemia, respiratory distress syndrome (RDS), neonatal intensive care unit (NICU) admission, total neonatal admissions, stillbirth, neonatal mortality within 1 week, and diabetes-related composite complications (at least one of the following: macrosomia, shoulder dystocia, neonatal hypoglycemia, neonatal hyperbilirubinemia, RDS).

Pre-pregnancy weight was the self-reported prepregnancy body weight at the initial examination. Babies were defined as LGA and SGA when their birth weights were above the 90th percentile and below the 10th percentile, respectively, for their gestation age. Macrosomia was defined as birth weight ≥4,000 g. Shoulder dystocia was defined as when some kind of manipulation (McRoberts maneuver, suprapubic pressure, or intravaginal manipulation) was required to expulse the fetal shoulder. The definition of PE covered cases of hypertension (reproducible blood pressure ≥140/90 mmHg) and urinary protein creatinine ratio >0.27 between 20 weeks’ gestation. Emergency caesarean delivery did not include scheduled caesarean delivery, even if it was advanced due to the onset of labor or rupture of membranes. Low APS was defined as Apgar score <7 at 5 minute and low UApH as <7.1. Neonatal hypoglycemia was defined as a plasma glucose level below 40 mg/dL. Hyperbilirubinemia was defined as when phototherapy was required because infant birth weight- and age-based reference values were exceeded. RDS was defined by characteristic findings on chest radiographic examinations and requiring oxygen within 24 h after birth. NICU
admission covered neonates who required intensive monitoring of their general condition. Total neonatal admission covered NICU admissions as well as cases that did not need intensive management but required interventions such as blood tests or phototherapy.

Presentation of data and statistical analysis

Statistical analyses were performed using JMP Pro version 12 (SAS Institute Inc., Cary, NC). Continuous variables were expressed as median (interquartile range) and categorical variables as frequencies and proportions. Comparison of the characteristics was performed with the Wilcoxon test for continuous variables and with the chi-square test for categorical variables.

Multiple logistic regression analysis was performed for categorical variables to calculate adjusted odds ratios (ORs) and 95% CI. Multiple linear regression analysis was performed for gestational age (weeks) at delivery and infant birth weight to calculate adjusted regression coefficients (RCs) and 95% CI. These multivariate analyses were adjusted for maternal age, height, pre-pregnancy weight, and multiparity. Variables with \( p < 0.05 \) were considered to be statistically significant.

Results

There was a total of 8,916 births during the study period. Among them, 3,772 women underwent 75-g OGGT during pregnancy. Of these, 286 women had multiple gestation and 22 women had FPG \( \geq 126 \) mg/dL, yielding a total of 305 women (some overlap) who were excluded. Further, 175 women who were diagnosed with GDM under the old JSOG criteria were excluded, leaving 3,292 subjects for the analysis (Fig. 1).

There were 503 women in the IGT group and 2,789 women in the NGT group. Table 1 shows the maternal characteristics of both groups. Maternal ages at delivery were 34 and 33 years, respectively, significantly higher in the IGT group (\( p < 0.01 \)). Gestational ages at delivery did not differ between the groups (\( p = 0.07 \)), though pre-pregnancy BMI and BMI at delivery were both significantly higher in the IGT group (\( p < 0.01, p < 0.01 \)). Gestational weight gain was 8.8 kg and 9.4 kg, respectively, showing significantly less weight gain in the IGT group (\( p < 0.01 \)).

Table 2 shows the pregnant outcomes. No difference was observed in the rates of GDM-related composite complications at 11.9% and 8.8%, respectively (adjusted OR 1.30, 95% CI 0.90–1.89, \( p = 0.16 \)). Infant birth weight was 3,094 g and 3,038 g, respectively, heavier in the IGT group but not significantly different (adjusted RC 10.0, 95% CI –14–34, \( p = 0.41 \)). Similarly, the frequencies of LGA did not differ significantly at 10.3% and 8.7%, respectively (adjusted OR 1.11, 95% CI 0.79–1.53, \( p = 0.53 \)).

Rates of PE and caesarian delivery did not differ between the groups. The rate of NICU admission did not differ between the groups, however total neonatal admission, at 10.5% and 7.1%, respectively, was significantly higher in the IGT group after adjustment (adjusted OR 1.55, 95% CI 1.12–2.13, \( p < 0.01 \)).

Discussion

In the present study, we compared the mild IGT pregnant women with NGT pregnant women. The former had significantly higher frequencies of total neonatal admissions; however, there were no differences regarding other pregnancy outcomes between the two groups.

A 2016 secondary analysis of the HAPO study showed that pregnant women with mild IGT who were diagnosed with GDM under the IAGPSG criteria but not the C&C criteria exhibited more frequent LGA, caesarian deliveries, neonatal hypoglycemia, and hyperbilirubinemia compared to pregnant women with NGT [20]. Further, a 2015 Canadian study found that pregnant women with mild IGT diagnosed under the IAGPSG criteria but not the old Canadian criteria exhibited more frequent hyper-
Tensive disorder of pregnancy (HDP), caesarian delivery, and LGA, and higher birth weights than pregnant women with NGT [21]. A 2017 Australian study reported higher rates of macrosomia/LGA, nursery admission, labor intervention, HDP, and other issues among pregnant women with mild IGT diagnosed under the IAGP/SG criteria but not the old Australian criteria, compared to pregnant women with NGT [22]. This shows that preg-

### Table 1 Maternal characteristics

<table>
<thead>
<tr>
<th></th>
<th>IGT group</th>
<th>NGT group</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, y.o.: median (IQR)</td>
<td>34 (30–36)</td>
<td>33 (29–36)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Height, cm: median (IQR)</td>
<td>158 (155–162)</td>
<td>158 (155–162)</td>
<td>0.26</td>
</tr>
<tr>
<td>Prepregnancy weight, kg: median (IQR)</td>
<td>54 (50–62)</td>
<td>52 (48–58)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Prepregnancy BMI, kg/m²: median (IQR)</td>
<td>21.4 (19.7–24.8)</td>
<td>20.7 (19.3–23.0)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Weight at delivery, kg: median (IQR)</td>
<td>64.5 (58.4–71.8)</td>
<td>62.4 (57.5–68.4)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Gestational weight gain (kg), median (IQR)</td>
<td>8.8 (6.2–11.9)</td>
<td>9.4 (7.1–12.0)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Multiparous, n (%)</td>
<td>260 (51.7%)</td>
<td>1,319 (47.3%)</td>
<td>0.07</td>
</tr>
</tbody>
</table>

IGT, Impaired Glucose Tolerance; NGT, Normal Glucose Tolerance

### Table 2 Pregnancy outcomes

<table>
<thead>
<tr>
<th></th>
<th>IGT group n = 503</th>
<th>NGT group n = 2,789</th>
<th>crude OR (95% CI)</th>
<th>p value</th>
<th>adjusted*1 OR (95% CI)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diabetes related composite complications*1, n (%)</td>
<td>60 (11.9)</td>
<td>246 (8.8)</td>
<td>1.40 (1.04–1.89)</td>
<td>0.03</td>
<td>1.30 (0.90–1.87)</td>
<td>0.16</td>
</tr>
<tr>
<td>Macrosomia, n (%)</td>
<td>7 (1.4)</td>
<td>39 (1.4)</td>
<td>1.00 (0.44–2.24)</td>
<td>0.99</td>
<td>0.80 (0.32–1.71)</td>
<td>0.58</td>
</tr>
<tr>
<td>Shoulder dystocia, n (%)</td>
<td>8 (1.6)</td>
<td>36 (1.3)</td>
<td>1.24 (0.57–2.67)</td>
<td>0.59</td>
<td>1.08 (0.46–2.26)</td>
<td>0.84</td>
</tr>
<tr>
<td>Neonatal hypoglycemia, n (%)</td>
<td>7 (1.4)</td>
<td>21 (0.8)</td>
<td>1.86 (0.79–4.40)</td>
<td>0.15</td>
<td>1.84 (0.71–4.20)</td>
<td>0.19</td>
</tr>
<tr>
<td>Neonatal hyperbilirubinemia, n (%)</td>
<td>37 (7.4)</td>
<td>144 (5.2)</td>
<td>1.46 (1.00–2.12)</td>
<td>&lt;0.05</td>
<td>1.48 (0.98–2.10)</td>
<td>0.06</td>
</tr>
<tr>
<td>RDS, n (%)</td>
<td>6 (1.2)</td>
<td>28 (1.0)</td>
<td>1.19 (0.49–2.89)</td>
<td>0.7</td>
<td>1.14 (0.42–2.62)</td>
<td>0.78</td>
</tr>
<tr>
<td>Pre-eclampsia, n (%)</td>
<td>10 (2.0)</td>
<td>50 (1.8)</td>
<td>1.11 (0.56–2.21)</td>
<td>0.76</td>
<td>0.89 (0.42–1.71)</td>
<td>0.73</td>
</tr>
<tr>
<td>LGA, n (%)</td>
<td>52 (10.3)</td>
<td>243 (8.7)</td>
<td>1.21 (0.88–1.66)</td>
<td>0.24</td>
<td>1.11 (0.79–1.53)</td>
<td>0.53</td>
</tr>
<tr>
<td>SGA, n (%)</td>
<td>40 (8.0)</td>
<td>266 (9.5)</td>
<td>0.82 (0.58–1.16)</td>
<td>0.26</td>
<td>0.85 (0.59–1.20)</td>
<td>0.36</td>
</tr>
<tr>
<td>Low APS, n (%)</td>
<td>39 (7.8)</td>
<td>195 (7.0)</td>
<td>1.12 (0.78–1.60)</td>
<td>0.54</td>
<td>1.14 (0.78–1.62)</td>
<td>0.49</td>
</tr>
<tr>
<td>Low UApH, n (%)</td>
<td>13 (2.6)</td>
<td>36 (1.3)</td>
<td>2.03 (1.07–3.85)</td>
<td>0.03</td>
<td>2.00 (1.00–3.75)</td>
<td>0.05</td>
</tr>
<tr>
<td>Emergency CS, n (%)</td>
<td>58 (11.5)</td>
<td>291 (10.4)</td>
<td>1.12 (0.83–2.01)</td>
<td>0.27</td>
<td>1.03 (0.75–1.39)</td>
<td>0.87</td>
</tr>
<tr>
<td>Total CS, n (%)</td>
<td>138 (27.4)</td>
<td>662 (23.7)</td>
<td>1.21 (0.98–1.50)</td>
<td>0.08</td>
<td>1.06 (0.85–1.33)</td>
<td>0.58</td>
</tr>
<tr>
<td>NICU admission, n (%)</td>
<td>25 (5.0)</td>
<td>109 (3.9)</td>
<td>1.29 (0.82–2.00)</td>
<td>0.27</td>
<td>1.32 (0.83–2.04)</td>
<td>0.24</td>
</tr>
<tr>
<td>Neonatal admission (include NICU), n (%)</td>
<td>53 (10.5)</td>
<td>198 (7.1)</td>
<td>1.54 (1.12–2.12)</td>
<td>&lt;0.01</td>
<td>1.55 (1.12–2.13)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Stillbirth, n (%)</td>
<td>1 (0.2)</td>
<td>7 (0.3)</td>
<td>0.79 (0.10–6.45)</td>
<td>0.83</td>
<td>0.98 (0.05–5.60)</td>
<td>0.98</td>
</tr>
</tbody>
</table>

OR, Odds ratio; RC, Regression Coefficient; IQR, Interquartile range; IGT, impaired glucose tolerance; NGT, normal glucose tolerance

*1 Diabetes-related composite complications defined as cases with at least one of the following: macrosomia, shoulder dystocia, neonatal hypoglycemia, neonatal hyperbilirubinemia, or RDS.

*2 Adjusted Odds Ratio adjusted for NGT group, adjusted for maternal age, height, pre-pregnancy weight, multiparous

*3 Adjusted Regression Coefficient for NGT group, adjusted for maternal age, height, pre-pregnancy weight, multiparous

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nant women who were not diagnosed under their nations’ old criteria but were diagnosed with mild IGT under the IADPSG criteria are at higher risk than women with NGT. The present study did not observe an increase in perinatal adverse events that previous research has shown except total neonatal admission. One possible reason for this is that the timing for performing 75-g OGTT in Japan does not match the IADPSG recommendation. The JSOG places no limits on the gestational age when a 75-g OGTT can be performed on pregnant women suspected of having IGT [15], meaning that many women are tested during a period other than 24–28 weeks gestation. In fact, about 60% (1,991/3,292) of the subjects had received 75-g OGTT outside the range of 24 to 28 weeks. In previous research into issues unique to Japan, we did not observe an improvement in perinatal adverse events in a group that was diagnosed with GDM before 20 weeks’ gestation and received interventions early [23], though this result could have been influenced by the fact that many of the subjects were diagnosed at a time not recommended by the IADPSG criteria. That said, the present study also found an increase in neonatal admissions, which suggests that the introduction of the IADPSG criteria has enabled the detection of cases of mild IGT.

In this study, we did not examine GDM morbidity rates before and after introducing the IADPSG criteria, although some previous Japanese studies have reported 2.7–5.7-fold increases [9, 16–19]. The cost-effectiveness of introducing the IADPSG criteria has been studied in other countries [10, 11], but remains controversial. This is because the introduction of these criteria would lead to an increase in the number of patients diagnosed with GDM, which would in turn cause an increase in overall medical costs. However, if a substantial improvement in pregnancy and delivery outcomes can be achieved, then the introduction of these criteria can be considered appropriate and the drawbacks can be ignored. There is a need for further investigations to examine this topic from a cost-benefit standpoint.

This study had the following limitations. First, it was a single-center, retrospective study of a small number of cases. Second, treatment of GDM has two aims, to improve the perinatal outcome of the index pregnancy and to improve the long-term health of both the mother and the offspring. However, the latter was not examined in the present study.

In conclusion, pregnant women with IGT, who were within the normal range under the old JSOG criteria but to be newly diagnosed under the IADPSG criteria, were at higher risk of neonatal admission than were pregnant women with NGT; however, there were no differences in other pregnancy outcomes between the two groups. We recommend that the appropriateness of introducing the IADPSG criteria in Japan should be reassessed, including at the timing of 75-g OGTT.

Acknowledgments

None.

Declaration of Interest

None.

References


