DOI: http://dx.doi.org/10.18203/2320-1770.ijrcog20174693

Original Research Article

Does mild hyperglycemia in 75gm Glucose Tolerance Test (GTT) affect outcome in pregnant women?

Karthiga Prabhu J.*, Muthulakshmi M.

Department of Obstetrics and Gynecology, SRM Medical College Hospital and Research Centre, Potheri, Kanchipuram, Tamil Nadu, India

Received: 24 September 2017 **Accepted:** 07 October 2017

*Correspondence:

Dr. Karthiga Prabhu J.,

E-mail: j.karthigaprabhu@gmail.com

Copyright: © the author(s), publisher and licensee Medip Academy. This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Background: Glucose tolerance in pregnancy is fundamentally linked to fetal growth. The relationship between maternal glycemia and adverse outcomes is a continuous process, with no distinct cut off point for increased risk. The objective of the study is to find out whether mild hyperglycemia in 2 hour 75 gm Glucose Tolerance Test (GTT) affects maternal and perinatal outcome in pregnant women.

Methods: This case control study was conducted in SRM Medical College during a 10-month period. Mild hyperglycemia was diagnosed when the - 2hour non- fasting 75gm GTT was between 120-139 mg/dl and Controls were women with 2hour nonfasting 75gm GTT <120mg/dl. Maternal and neonatal parameters were noted and the results were compared.

Results: During the study period 142 delivered women had mild hyperglycemia, of which 10 patients on subsequent blood sugar monitoring required insulin for blood sugar control. There was significant family history of diabetes in women with mild hyperglycemia when compared to controls. There was no significant difference in incidence of hypertension, hypothyroidism, preterm delivery and caesarean section between the two groups. LGA (Large for gestational age babies) (p=0.001) and serum triglyceride levels (p=0.04) were significantly more in women with mild hyperglycemia when compared to controls.

Conclusions: Mild hyperglycemia during pregnancy should not be ignored and periodic blood sugar monitoring should be done to improve maternal and fetal outcome.

Keywords: GDM, Glucose tolerance test, Hyperglycemia in pregnancy, Impaired glucose tolerance, Large for gestational age (LGA) babies

INTRODUCTION

Increasing carbohydrate intolerance in pregnant women is associated with a graded increase in adverse maternal and fetal outcomes.¹ Untreated GDM is associated with a higher incidence of complications during pregnancy and increased perinatal mortality and morbidity.

Glucose tolerance in pregnancy is fundamentally linked to fetal growth. Glucose is the main energy substrate for intrauterine growth and is transmitted steadily from mother to fetus. Pedersen hypothesized that in maternal diabetes, hyperglycemia leads to fetal hyperinsulinemia and increased utilization of glucose and, hence, increased fetal growth.² This is supported by observations in gestational and pregestational diabetes where higher glucose concentrations, particularly post meal, predict greater infant birth weight.³

A relation between maternal glucose and fetal growth also exist in women who do not have diabetes.⁴ In India, Gestational diabetes is diagnosed and treated when 2hour non-fasting 75gm GTT was ≥140mg/dl.⁵ The relationship between maternal glycemia and adverse outcomes is a continuous process, with no distinct cut off point for increased risk.⁶

The objective of the study is to find out whether mild hyperglycemia in 2 hour 75 gm GTT (120-139 mg/dl) affects maternal and perinatal outcome in pregnant women.

METHODS

This case - control study was conducted in SRM Medical College Hospital and Research Centre during a 10 month period. All patients delivered during the study period were divided into 3 groups based on 2hour non- fasting 75gm GTT - Gestational diabetes (≥140 mg/dl), Mild hyperglycemia (120-139 mg/dl) and Controls (<120mg/dl). Women with gestational diabetes and pregestational diabetes were excluded from the study. Maternal and neonatal parameters were noted and the results were compared between women with mild hyperglycemia and controls.

Statistical Analysis was done using Graph pad Instat 3 and SPSS 12 Software. P value <0.05 was considered significant.

RESULTS

Total number of deliveries during the ten month study period was 1027. Among the delivered mothers 142 women had mild hyperglycemia, of which 10 patients required insulin for blood sugar control on subsequent blood sugar monitoring.

Table 1: Maternal characteristics between women with mild hyperglycemia and controls.

	Cases (142)	Controls (827)	p
Mean age (yrs)	26.38±3.92	25.51±3.67	0.14
<20yrs	4 (3.1%)	13 (1.5%)	
20-29yrs	113 (79.6%)	711(85.9%)	
30-34 yrs	22 (15.6%)	84 (10.1%)	
>35yrs	2 (1.5%)	19 (2.3%)	0.38
Parity			
Primipara	60 (42%)	299 (36%)	0.18
Previous history	15 (10.5%)	65 (7.85%)	0.32
of pregnancy loss			
History of GDM in	6 (4.2%)	2 (0.02%)	-
previous pregnancy			
Family history of	42 (29.5%)	36 (4.3%)	0.03*
diabetes			
BMI			
25-29.9kg/m ²	32 (22.5%)	144 (17.4%)	0.15
(overweight)			
>30kg/m ² (obesity)	18 (12.6%)	75 (9.06%)	0.21
Hypothyroidism	24 (16.9%)	54 (6.5%)	0.33

^{*}statistically significant

Mild hyperglycemia was noted in 2 patients in first trimester, 30 patients in second tyrimester and remaining in third trimester. There was significant family history of diabetes in women with mild hyperglycemia (29.5%) when compared to controls (4.3%), p=0.04 (Table 1).

Table 2: Maternal and neonatal outcome between women with mild hyperglycemia and controls.

	Cases (142)	Controls (827)	p	
Hypertensive	12 (8.4%)	69 (8.3%)	0.31	
disorders				
Urinary tract	14 (9.8%)	51 (6.2%)	0.1	
infection				
Mean serum	218.8±68.4	179.25±55.25	0.04*	
triglyceride levels		mg/dl		
Caesarean section	62 (43.6%)	290 (35%)	0.05	
Indications for CS				
Fetal distress	14 (30%)	30 (35.2%)		
CPD	17 (36.9%)	29 (34.11%)		
Doubtful scar	7 (15.2%)	16 (18.8%)		
Malpresentations	2 (4.3%)	2 (2.35%)		
Failed induction	3 (6.5%)	7 (8.23%)		
Cervical dystocia	3 (6.5%)	5 (5.88%)		
Preterm labour	9 (6.3%)	50 (6%)	0.8	
Sex of baby				
Male	75 (53.1%)	433 (52.3%)	1	
Female	67 (46.9%)	394 (47.6%)		
Weight of the baby				
<2500g	6 (9.3%)	15 (12.09%)		
2500-3500g	47 (73.4%)	105 (82%)		
>3500g	22 (15.5%)	55 (6.6%)	0.001^{*}	
NICU admission	11 (7.7%)	60 (7.2%)	0.8	
Indications – NICU				
Birth asphyxia	4 (36.3%)	16 (26.7%)		
Neonatal	5 (45.4%)	32 (53.3%)		
jaundice				
Prematurity	2 (18.1%)	12 (20%)		

^{*}statistically significant

There was no significant difference in incidence of hypertension, hypothyroidism, preterm delivery and caesarean section between the two groups. LGA (large for gestational age babies $\geq 3500 \, \mathrm{g}$) (p=0.001) and serum triglyceride levels (p=0.04) were significantly more in women with mild hyperglycemia when compared to controls (Table 2).

DISCUSSION

GDM is becoming an increasing health problem worldwide. GDM is associated with increased maternal – fetal morbidity as well as long-term complications in both mother as well as the offsprings. Recent studies have reported that maternal glucose intolerance less severe than GDM is associated with an increased risk of adverse pregnancy outcomes.⁶

In the present study, though there were no significant difference in hypertensive disorders, preterm labour, urinary tract infections and NICU infections, Large for gestational age babies (Birthweight \geq 3500gm) was significantly more in women with mild hyperglycemia when compared to controls.

Scholl et al has shown that increased maternal glucose concentration was associated with an increased risk of large-for-gestation fetuses (p<0.001) and a decreased risk of fetal growth restriction (p<0.05) in non diabetic mothers.⁴ Maternal complications including cesarean section and chorioamnionitis were increased twofold or more with high maternal glucose concentrations. A observational study from the Toronto Tri-Hospital Gestational Diabetes Project has shown that in women without GDM increasing carbohydrate intolerance is associated with a graded increase in adverse fetomaternal outcomes. Macrosomia is an important factor in high cesarean delivery rates for women with untreated borderline GDM.¹

Several studies suggested that maternal circulating fatty acids play important roles in fetal growth and development. Elevated Fatty acids in the maternal circulation are associated with increased insulin resistance and β -cell dysfunction, which play a role in the development of gestational diabetes mellitus (GDM) and increases the risk of adverse perinatal outcomes, including preterm delivery.

In the present study serum triglyceride levels was also significantly more in women with mild hyperglycemia when compared to controls. Wei et al, has shown that blood serum Triglyceride, total cholesterol and LDL-Cholesterol concentrations were higher in the Gestational impaired glucose tolerance (GIGT) and GDM groups (p<0.05) which is supported by Chen et al. who noted that abnormalities in fat metabolism are present in both GDM and GIGT women.^{9,10}

CONCLUSION

Mild hyperglycemia during pregnancy should not be ignored and periodic blood sugar monitoring should be done to improve maternal and fetal outcome.

Funding: No funding sources Conflict of interest: None declared

Ethical approval: The study was approved by the

Institutional Ethics Committee

REFERENCES

- 1. Sermer M, Naylor CD, Farine D, Kenshole AB, Ritchie JW, Gare DJ et al. The Toronto Tri-Hospital gestational diabetes project: A preliminary review. Diabetes Care. 1998 Aug;21 Suppl 2:B33-42.
- 2. Pedersen J. Weight and length at birth of infants of diabetic mothers. Acta Endocrinol. 1954;16:330-42.
- 3. Jovanovic-Peterson L, Peterson CM, Reed GF, Metzger BE, Mills JL, Knopp RH, et al. Maternal postprandial glucose levels and infant birth weight: the Diabetes in Early Pregnancy Study. The National Institute of Child Health and Human Development—Diabetes in Early Pregnancy Study. Am J Obstet Gynecol. 1991;164:103-11.
- 4. Scholl TO, Sowers M, Chen X, Lenders C. Maternal glucose concentration influences fetal growth, gestation, and pregnancy complications. Am J Epidemiol. 2001;154 (6): 514-520.
- Seshiah V, Balaji V, Balaji S, Sekar A, Sanjeevi CB, Green A. One step screening procedure for screening and diagnosis of gestational diabetes mellitus. J Obstet Gynecol India. 2005;55(6):525-9.
- HAPO Study Cooperative Research Group. The hyperglycemia and adverse pregnancy outcome (HAPO) study. Intl J Gynaecol Obstet. 2002;78:69-77
- 7. Berghaus TM, Demmelmair H, Koletzko B. Fatty acid composition of lipid classes in maternal and cord plasma at birth. Eur J Pediatr. 1998;157:763-8.
- 8. Schaefer-Graf UM, Graf K, Kulbacka I, Kjos SL, Dudenhausen J, Vetter K, et al. Maternal lipids as strong determinants of fetal environment and growth in pregnancies with gestational diabetes mellitus. Diabetes Care. 2008;31:1858-63.
- Chen X, Scholl TO, Leskiw M, Savaille J, Stein TP. Differences in maternal circulating fatty acid composition and dietary fat intake in women with gestational diabetes mellitus or mild gestational hyperglycemia. Diabetes Care. 2010;33(9):2049-54.
- 10. Wei J, Gao J, Cheng J. Gestational diabetes mellitus and impaired glucose tolerance pregnant women. Pak J Med Sci. 2014 Nov-Dec;30(6):1203-8.

Cite this article as: Prabhu JK, Muthulakshmi M. Does mild hyperglycemia in 75gm Glucose Tolerance Test (GTT) affect outcome in pregnant women?. Int J Reprod Contracept Obstet Gynecol 2017;6:4866-8.