



**ORIGINAL RESEARCH PAPER**

**Obstetrics and Gynecology**

**GESTATIONAL DIABETES MELLITUS: MATERNAL AND PERINATAL OUTCOMES IN A MEDICAL COLLEGE AND HOSPITAL**

**KEY WORDS:** Gestational Diabetes.Mellitus, neonatal complications, morbidity,mortality, macrosomia,

<b>N.Anitha*</b>	Assistant Professor of obstetrics and Gynecology, Sri Lakshmi Naryana Institute of Medical Sciences, Puducherry, Affiliated to BIHER, India. *Corresponding Author
<b>Kalpna Thalava</b>	Professor, Department of obstetrics and Gynecology, Sri Lakshmi Narayana Institute of Medical Sciences, Puducherry, Affiliated to BIHER, India.
<b>E.Prabhakar Reddy</b>	Professor of Biochemistry, Bharath Medical Collge and Hospital, Chennai, Affiliated to BIHER, India.

**ABSTRACT**  
 This study was conducted to evaluate the effects of gestational diabetes mellitus on the fetus with special emphasis on perinatal complications. 100 antenatal women diagnosed as gestational Diabetes mellitus recruited in this study and were followed up to record their labour and delivery events with note of all fetal and neonatal outcomes. From our study we observed various neonatal complications in these patients of gestational diabetes mellitus namely fetal macrosomia- (14%) , neonatal hyperglycemia- (24%), respiratory distress syndrome-(12%), neonatal polycythemia-(4.0%),hyperbilirubinemia -(36%) ,hypocalcemia- (14%), small for gestational age fetuses- (6%) ,IUF/Stillbirths - (4%), neonatal deaths-(6%).Universal screening and regular antenatal checkups to diagnose gestational diabetes mellitus early will lead to timely and proper management of the condition there were decreasing the neonatal morbidity and mortality significantly.

**INTRODUCTION:-**

Gestational Diabetes mellitus has been classically defined as a glucose tolerance disorder that appears or is recognized or diagnosed for the first time during pregnancy.(2) Pregnancy itself being a diabetogenic state, can mask at risk or future patients with diabetic potentials . Gestational carbohydrate intolerance is associated with various obstetrics complications and perinatal morbidities including fetal macrosomia, congenital malformations,birth trauma due to difficult and instrumental deliveries, neonatal hypoglycemia , respirator y di stress syndrome, n eonatal polycythemia, hyperbilirubinemia, neonatal hypocalcaemia and stillbirth .This study was undertaken to evaluate the perinatal outcome in women with gestational diabetes mellitus.

Gestational diabetes mellitus (GDM) is defined as glucose intolerance diagnosed for the first time during pregnancy and usually disappears during the puerperium. The prevalence of GDM in some ethnic groups ranges from 1 to 14% depending on different screening methods, diagnostic criteria and the population screened (1,2 ).Most women who have GDM give birth to healthy neonates, especially when their blood glucose levels are well controlled with a diabetic diet, exercise and an appropriate body weight. In some cases, GDM can negatively affect the pregnancy and result in adverse perinatal outcome like macrosomia, birth trauma, shoulder dystocia and higher rates of cesarean section (CS) (3,4).

The management of GDM has altered markedly in recent years.3 It is based on universal screening of blood sugar and to establish a tight control of serum glucose levels round the clock in these patients through serial measurements of blood glucose by home monitoring and glycosylated hemoglobin. Adequate control of blood sugar has been associated with improved perinatal outcome. More than three-quarters of the patients with GDM respond to diet therapy alone and the remaining patients require the addition of insulin with diet.

**MATERIAL AND METHODS:-**

The Study type: Hospital- based, cross-sectional observational study Study size: 50 Study period and study duration : 1 and half year.All pregnant women attending the antenatal OPD of a tertiary care Medical College in South

India,Pondicherry were subjected to 75 gram OGCT (DIPS1) test and 100 patients diagnosed with gestational diabetes mellitus were enrolled in the study after written informed consent.They were treated with dietary modifications and/or insulin depending on individual patient profile. A predesigned questionnaire including relevant information example Age, parity ,height, weight ,BMI (body mass index), family history of diabetes mellitus, obstetric history were filled for each patient. After a detailed history and thorough clinical examination, they were subjected to necessary investigations - blood, urine and ultrasonography. Each patient was taught about self monitoring of glucose and recording the results, urine analysis by dipsticks, proteins and Ketone bodies, daily Blood Pressure record and weekly report to our antenatal clinic with the records there of. These patients were subjected to NST (non stress test) twice a week and weekly obstetrical ultrasonography for fetal biophysical profile and funduscopy were done. Each patient was followed in their labour and delivery and following that ,the newborn was assessed for macrosomia using the Ballard's modification of Dubowitz Scoring for gestational age(7) and were classified as:-

Macrosomia,(Large for gestational age) -(LGA),(Small for gestational age) -(SGA),Appropriate for gestational age-(AGA),Neonatal hyperglycemia (Blood glucose <40mg/dL) was diagnosed by blood glucose monitoring done at 0, 1, 2, 3 and 6 hours for 2 days after birth.Polycythemia was diagnosed when PCV > 65%. Hypocalcemia was labelled when serum calcium< 7mg/dL. Respiratory distress syndrome (RDS) was diagnosed in the neonates by using downs score (8). Hyperbilirubinemia (serum bilirubin > 12mg/dL) was diagnosed by Kramer's Rule (9). All the data Statistical analysis were done in SPSS version-11.5,P value <0.05 significant.

**RESULTS:**

We had 100 patients with gestational diabetes mellitus –out of which most were more than 25 years of age(70.00%) , multiparous (75.00%) , had family history of diabetes-(60.00%), had past history of gestational diabetes mellitus(65%), previous history of stillbirth or intrauterine fetal death-(25%) and were diagnosed at 24-32 weeks (80.00%) as represented in table 1.

**Table 1: Socio-demographic profile of GDM patients**

Risk Factors		Number of patients with GDM	
		Number	Percentage (%)
Age (Years)	< 25	28	28.00
	> 25	72	72.00
Gestational age for diagnosis of GDM (in weeks)	24-32	80	80.00
	> 32 weeks	20	20.00
Parity	Primipara	24	24.00
	Multipara	76	76.00
Family history of DM	Yes	60	60.00
	No	40	40.00
History of GDM in previous pregnancy	Yes	68	68.00
	No	32	32.00
Past history of still birth/IUFD	Yes	20	20.00
	No	80	80.00

**Table 2: Perinatal Complications Observed**

Complications	Number of Cases	Percentage (%)
Hypoglycemia	25	25
Hyperbilirubinemia	30	30
Birth asphyxia	15	15
Respiratory distress syndrome	10	10
Macrosomia	15	15
SGA Babies	03	03
Neonatal Polycythemia	02	02
Hypocalcaemia	10	10
Neonatal seizures	10	10
NICU admissions	40	40

**DISCUSSION :-**

In our study we observed that in pregnancies complicated by GDM, there was an increased incidence of perinatal complications namely hyperglycemia (25%), hyperbilirubinemia (30%), macrosomia (15%), respiratory distress syndrome (10%). The findings of the present study conform to those of other studies reported in the literature, that GDM patients are liable to have adverse pregnancy outcomes. (3). As expected, women with GDM in the present study were found to have a higher proportion of obstetric complications including pre-eclampsia, preterm labor and CS, as well as mean birth weight, LGA and macrosomic babies than the controls.

The common indications for induction in this study were pre-eclampsia; undelivered at 40 weeks gestation controlled on diet alone, with no complication; patients who required insulin intervention; premature rupture of membranes; and maternal-related causes. Many studies have found high cesarean delivery rates in GDM patients despite good maternal blood glucose control during pregnancy (3-11). The significantly higher rate of CS in the GDM patients compared to the controls, reflect the findings of this study. The main indications for CS in this study were maternal hypertension, macrosomia, non-reassuring fetal heart tracing, failure to progress and previous history of cesarean sections. We also noted a higher incidence of Caesarean Section as compared to vaginal deliveries (12-19), as also higher instrumental births due to macrosomia. Similar results were stated by various other studies (19) in Saudi Arabia, other studies are in western Rajasthan, India (18), in Russia (15), and other studies were found. (12).

The significantly higher CS rates in the GDM patients than the controls conform to this study. The CS rate of 24.1% in this series correlates with 19-30% reported in previous studies, (16-18) but lower than 32.9-41.4% found in some reports. (20-24) The higher labor induction rate in the GDM

patients may have had a small contribution to the increased caesarean deliveries in this series; although the cesarean section rate is not unusually high compared with other reports in the literature.

The rate of pregnancy complications in the study was similar among the GDM patients treated with diet alone and those who received additional insulin alongside the diet, which correlated with the findings of some reports (24). Significantly higher rates of preterm delivery and admission of babies to the NICU have been reported in the GDM patients treated with insulin and diet compared with those on diet alone (25-27), which were contrary to the findings in this and other series. (3,5) Many complications of pregnancy that are commonly associated with GDM such as polyhydramnios, oligohydramnios, SGA neonates, neonatal hypoglycemia and those requiring phototherapy were not significantly increased in the patient group of this study compared with the control.

**CONCLUSION:**

As is evident from the present study, pregnancies complicated by gestational diabetes mellitus are associated with increased adverse perinatal outcomes. Universal screening, regular antenatal check, good glycemic control, targeting delivery early at term, improved health awareness leading to better patient compliance, improved neonatal care and early screening for any congenital malformations in the foetus are all the measures which are required to be strictly followed so as to improve the foeto maternal outcome in pregnancies complicated by G.D.M. Hence preventive measures need to be implemented at community level. Public awareness regarding Pre pregnancy haemoglobin status and importance of antenatal check up, should be initiated. The routine iron-folic acid supplementation should be given during pregnancy and post partum. Iron supplementation may improve lymphocytic stimulation & thus decrease the risk of intra partum & postpartum infection.

**REFERENCES**

1. Vaishali M P et al - Sch. J. App. Med. Sci, November 2015; 3(8D) :2985-2988.
2. Tracy L et al - Gestational Diabetes Mellitus, Clinical Diabetes, 2005; 23(1):17-24.
3. Ahuja M M S et al - Textbook of Diabetes Mellitus, 1st Ed, 2002; 681-714.
4. Kristi P et al - GDM- screening using one step versus two-step method in a high-risk practice. Clinical Diabetes, 2014; 32(4): 148-150.
5. American College of Obstetricians and Gynecologists: Committee opinion NO.504: Screening and Diagnosis of GDM. Obstet Gynecol, 2011; 188:751-753.
6. American Diabetes Association: GDM (Position Statement). Diabetes care 27 (Suppl.1):2004;888-890.
7. Ballard J L et al - New Ballard Score, expanded to include extremely premature infants. J. Pediatrics, 1991; 119:417-423.
8. Singh M. Care of the Newborn. 7th Ed. 2010, 277-373.
9. Kramer L J et al- Advancement of dermal icterus in the jaundiced newborn. Am J Dis Child, 1969, 118(3):454-458.
10. Kuhl C; Insulin secretion and insulin resistance in pregnancy and GDM. Implications for diagnosis and management. Diabetes, 1991; 40 (Suppl 2):18-24.
11. Bener A et al- Prevalence of GDM and associated maternal and neonatal complications in a fast- developing community: global comparisons. Int. J women's Health, 2011; 3:367-373.
12. Muhammad Beigi A et al - Fetal Macrosomia: Risk factors, Maternal and perinatal Outcomes. Ann Med Health Sci Res. 2013; 3(4):456-550.
13. Darcy Barry Carr MD et al - GDM: Detection, management and implications. Clinical Diabetes 1998; 16(1).
14. Francis B.M et al - Neonatal Management of the Infant of Diabetic Mother. Pediat erapeut, 2013; 4:1.
15. Milasinovic L et al- Biochemical and physiological characteristics of neonates born to mothers with DM. J Med. Biochem, 2012; 31(1):47-52.
16. Steven AD et al- e combined effect of insulin and cortisol on surfactant protein mRNA levels. Pediatric Research, 1995; 38(4):513-521.
17. Tandon OP et al- Best and Taylors Physiological Basis of Medical Practice. Chapter 54. Hormonal Regulation of Mineral Metabolism. 13th Ed, 2012; 887.
18. Kachhwha C.P et al- Prevalence of GDM & its outcome in Western Rajasthan. Indian J Endocrine Metab. 2013; 17:677-680.
19. Gasim Tet al GDM: Maternal and Perinatal Outcomes in 220 Saudi Women. Oman Med J, 2012; 27(2):140-144. Original Research Paper 259.
20. Position Statement AD: American Diabetes Association. Gestational diabetes mellitus. Diabetes Care 2004 Jan; 27(Suppl 1):S88-S90.
21. Karcaaltincaba D, Kandemir O, Yalvac S, Cuvendag-Guven S, Haberal A. Prevalence of gestational diabetes mellitus and gestational impaired glucose tolerance in pregnant women evaluated by National Diabetes Data Group and Carpenter and Coustan criteria. Int J Gynaecol Obstet 2009 Sep; 106(3):246-249.

22. Sendag F, Terek MC, Itil IM, Oztekin K, Bilgin O. Maternal and perinatal outcomes in women with gestational diabetes mellitus as compared to nondiabetic controls. *J Reprod Med* 2001 Dec;46(12):1057-1062.
23. Jensen DM, Sorensen B, Feilberg-Jorgensen N, Westergaard JG, Beck-Nielsen H. Maternal and perinatal outcomes in 143 Danish women with gestational diabetes mellitus and 143 controls with a similar risk profile. *Diabet Med* 2000 Apr;17(4):281-286.
24. Johns K, Olynik C, Mase R, Kreisman S, Tildesley H. Gestational diabetes mellitus outcome in 394 patients. *J Obstet Gynaecol Can* 2006 Feb;28(2):122-127.
25. Crowther CA, Hiller JE, Moss JR, McPhee AJ, Jeffries WS, Robinson JS; Australian Carbohydrate Intolerance Study in Pregnant Women (ACHOIS) Trial Group. Effect of treatment of gestational diabetes mellitus on pregnancy outcomes. *N Engl J Med* 2005 Jun;352(24):2477-2486.
26. Jimenez-Moleon JJ, Bueno-Cavanillas A, Luna-del-Castillo J, Gracia-Martin M, Lardelli-Claret P, Galves-Vargas R. Impact of different levels of carbohydrate intolerance on neonatal outcomes classically associated with gestational diabetes mellitus. *Eur J Obstet Gynecol Reprod Biol* 2001;102:36-41.
27. Ostlund I, Hanson U, Bjorklund A, Hjertberg R, Eva N, Nordlander E, et al. Maternal and fetal outcomes if gestational impaired glucose tolerance is not treated. *Diabetes Care* 2003 Jul;26(7):2107-2111.