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Original Research Article

Pregnancy outcome in pre-gestational and gestational diabetic women: a prospective observational study

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ABSTRACT

Background: India is considered the world capital of diabetes, proper care and management of the same is the demand of society. The present study is carried out to identify the disease burden of GDM/ overt DM among antenatal cases. The main objective was to study the maternal and fetal outcome of diabetes complicating pregnancy.

Methods: The present study was conducted at Dr. Rajendra Prasad Government Medical College at Tanda, District KangraHP from October 2015 to September 2016. All antenatal cases were screened for diabetes by OGTT 75 gm, 2-hour blood glucose level as per DIPSI guidelines and labelled as GDM/ overt DM. Those who fulfilled selection criteria were enrolled in the study. A total of 6452 cases who attended antenatal clinic during the study duration and were screened were selected for the study. 116 cases were found to have GDM / overt DM. Seventy-nine had GDM with OGTT>140 mg/dl (DIPSI guidelines) and 37 had overt DM with 2 hours PP >200 mg/dl (WHO criteria). Total 100 cases comprised of study group were followed till delivery to study maternal and fetal outcome.

Results: The prevalence of diabetes in pregnancy was found to be low 1.79%. GDM was found to be more prevalent than overt diabetes in pregnant women (66% versus 34%). Among the antenatal maternal complications observed missed abortion (11.8% versus 1.55; p=0.026), polyhydraminos (26.4% versus 10.6%; p=0.04) and preterm labour (17.6% versus 4.5%; p=0.003) were significantly more common in overt diabetics than GDM cases. IUFD (8.8% versus 0; p=0.014) was also significantly more common in overt diabetics than GDM case. RDS was found significantly higher in neonates of overt diabetics as compared to GDM mothers (14.7% versus 1.5%; p=0.009).

Conclusions: Early detection and good glycemia control by MNT and insulin, regular antenatal check-ups, patient counselling and compliance, intrapartum fetal monitoring and early neonatal care are keys to improved outcome.

Keywords: Gestational, Diabetes, Hyperglycemia, Pre-gestational women, Gestational diabetic women

INTRODUCTION

A global estimate for hyperglycemia in pregnancy (HIP) have brought out that 16.9% of total pregnancies globally were affected by some form of hyperglycemia in 2013.¹ This translates to 21.4 million live births at risk of being exposed to a hyperglycemia intrauterine milieu. Southeast Asia region had the maximum prevalence of HIPof 25% and India is standing tall at 27.5% against the global average of 16.9%. Country specific estimates have further brought out that India had the highest number of women

affected by the HIP with an estimated 5.7 million cases in 2013. Diabetes in pregnancy can be classified into two major types: Gestational diabetes (GDM) –women who have carbohydrate intolerance of variable severity with onset or first recognition during pregnancy.^{1,2} Pregestational or Overt DM (PGDM) - women who were known to have diabetes before pregnancy.

The risk of adverse pregnancy outcome increases continuously with increasing maternal glucose even below the traditional cut-offs for gestational diabetes, with no threshold effect found. The findings of the Hyperglycaemia and adverse pregnancy outcomes (HAPO study) have shown continuous relationship between maternal glucose level and adverse outcomes.³

Pregnancy aggravates the diabetic state of the mother due to the diabetogenic effect of pregnancy hormones. However, the impact is more severe in the overt diabetic group. Pregnancy in diabetic women leads to disturbed glucose homeostasis, increased insulin requirement, and increased risk of nephropathy and retinopathy in mother. There are several factors that are associated with a greater risk of developing GDM/ overt DM during pregnancy. Women with age >25 years, high BMI or obesity, family history of DM, or belonging to ethnic groups like Asian, Pacific Islander, Native American, Hispanic, or black racial have a higher prevalence of DM. Women with past history of GDM or with a previous history of macrosomic baby/fetal loss/ congenital anomalies/ polyhydramnios/ still-birth or persistent glycosuria are found to be at increased risk.⁴ They demand more careful assessment, timely screening, early recognition, intense monitoring, and appropriate management. GDM accounts for 90% of all cases of diabetes in pregnancy, and if unrecognized and untreated, threatens the lives of both mother and baby. Women with GDM often give birth to macrosomic or large gestational age (LGA) infants. Pregnancies for complicated by GDM also have a 4-fold increased risk of perinatal mortality. Almost 3 million babies are stillborn every year, and GDM is a major contributor to this unacceptable loss of life.⁵ Although GDM is a temporary phenomenon for pregnant women, more than 50% of women with GDM develop type 2 diabetes within 5-10 years of delivery.⁶ Moreover, infants of women with GDM have a higher prevalence of overweight or obesity as young children and adolescents, and a higher risk of developing type 2 DM later in life. Studies have revealed that women with GDM usually have better maternal and neonatal outcome as compared to overt diabetes group and most of the women with GDM revert back to euglycemic state in postpartum period.^{7,8} Thus, diabetes complicating pregnancy act as a two-way sword- diabetes affecting pregnancy state of the mother and pregnancy affecting the diabetic state of the mother. Since India is considered the world capital of diabetes, proper care, and management of the same is the demand of society. The present study is carried out to identify the disease burden of GDM/ overt DM among antenatal cases in a tertiary care centre in the hilly terrain of North India. The main objective was to study the maternal and fetal outcome of diabetes complicating pregnancy with an idea to fill the knowledge gap.

METHODS

The present study was a prospective observational study, carried out in the Department of Obstetrics and Gynaecology and Department of Paediatrics, at Dr. Rajendra Prasad Government Medical College at Tanda, District Kangra, Himachal Pradesh. This was a one-year study conducted from October 2015 to September 2016.

Sample size

All the pregnant women attending the antenatal clinic or admitted in the antenatal ward and labor room with: Oral glucose tolerance test (OGTT) value >140 mg/dl. Known case of DM or 2 hr postprandial (PP) value >200 mg/dl were eligible to be enrolled for study, after taking written informed consent.

Inclusion criteria

All the pregnant women attending the antenatal clinic with singleton pregnancy, pregnancy with GDM and pregnancy with overt DM were included in the study.

Exclusion criteria

Women with multi-fetal gestation, antepartum eclampsia, jaundice complicating pregnancy, placenta praevia, chronic hypertension, known case of heart disease, seizure disorder on antiepileptics, tuberculosis/ bronchial asthma were excluded from the study group.

Methodology

After initial registration of all the antenatal cases at antenatal clinic, the details of obstetrics/menstrual and medical history were noted and complete general and obstetrics examination was done. This was followed by routine antenatal tests and screening for diabetes by Oral glucose tolerance test (OGTT) 75 gm glucose load irrespective of their last meal timing and blood glucose testing at 2 hours as per Diabetes in Pregnancy Study Group (DIPSI) guidelines at their first, visit. Women with normal values were again given glucose challenge in a similar manner between 24-28 weeks of gestation. Those women attending the antenatal clinic in our hospital for the first time in a later trimester were also screened with the help of DIPSI guidelines. The plasma glucose was estimated by the glucose oxidase-peroxidase (GOD-POD) method. Women with value ≥ 140 mg/dl and < 200 mg/dl were labelled as GDM and those with 2 hr PP \geq 200 mg/dl were labelled as overt diabetes (WHO criteria)

All GDM and overt DM cases who were ready to participate in the study and fulfilling selection criteria were enrolled in the study. The participants were admitted to the antenatal ward and were treated with diet and/or insulin therapy. GDM women were advised medical nutrition therapy (MNT) for 2 weeks in consultation with the dietician of the hospital. Those who achieved desired target levels while maintaining fasting plasma glucose (FPG) \leq 90 mg/dl and peak post-meal glucose \leq 120 mg/dl were discharged with weekly follow-up. Those who did not respond by maintaining FPG \leq 90 mg/dl and 2 hr PP \leq 120 mg/dl were advised insulin in consultation with the

medical department of the hospital. All of them were followed until delivery Figure 1.

Maternal complications like missed or threatened abortion, polyhydramnios, pregnancy induced hypertension (PIH), pre-eclampsia (PE), preterm labor, premature rupture of membrane (PROM) and any other morbidity was recorded in predesigned proforma. The period of gestation at the time of delivery and mode of delivery including instrumental delivery or cesarean section were recorded.

A detailed examination of the new born was done. The APGAR score at 1 minute and 5 minutes of birth and the resuscitation required were noted. Fetal parameters such as birth weight, any features of macrosomia, congenital malformation, blood sugar levels for incidence of hypoglycemia (at 0,1,2,3 hour of life followed by 6 hourly blood glucose monitoring till 48 hours of life: as per standard protocol followed in the pediatrics department of college), respiratory distress syndrome, hypocalcemia, polycythemia, neonatal jaundice, early neonatal death etc. were recorded in consultation with Paediatrician. The management and investigations of all the new-borns were done till their discharge and were recorded on predesigned proforma.

Statistical analysis

The complete data was collected in a Microsoft excel sheet and statistical analysis of data was done with the help of Epi info software. Mean values and percentages were collected to analyze the study.

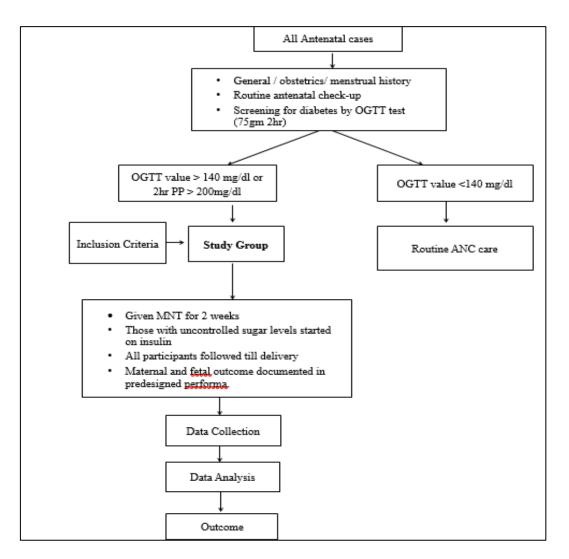


Figure 1: Methodology flow chart.

Ethical issues

There was no drug trial or human /animal experiment involved. All the women were treated in their best interest

in consultation with a consultant on duty. All the deliveries and cesarean sections were attended by a paediatrician. The Investigators and supervisors were well aware of the guidelines for ethics in the biomedical research document of ICMR (2006), Helsinki Declaration (modified 2000), and the policy of the Institutional Ethics Committee of Dr. RPGMC Tanda.

RESULTS

The present study was conducted to study the maternal and fetal outcome of diabetes in pregnancy in Department of Obstetrics and Gynaecology and Department of Paediatrics, at Dr Rajendra Prasad Government Medical College at Tanda, District Kangra, Himachal Pradesh from October 2015 to September 2016 after the due approval of the 'Institutional Ethics Committee'.

A total of 6452 antenatal cases who were screened as per DIPSI guidelines were selected. 116 cases were found to have GDM / overt DM. Thus, prevalence of diabetes complicating pregnancy was found to be 1.79%. Out of 116, 79 were labelled as GDM (OGTT >140 mg/dl) and 37 had overt DM (2 hr PP >200 mg/dl, WHO criteria). 2 cases did not consent for participation in study and 14 cases were lost to follow up. Total 100 cases comprised of study group.

Table 1: Characteristics of singleton women with gestational and pregestational diabetes mellitus as well age and neonatal birth weight at delivery.

Characteristic	Gestational diabetes (n=66)	Pregestational diabetes (n=34)	Statistical comparison between groups		
Mean Maternal Age (years)	29.5	31.4	p=0.010		
No. (%) Primigravidae	25 (37.9)	8 (23.5)	p =0.148		
Mean Pre pregnancy BMI * (Kg/m2) No. (%)	24.72 ± 3.75	25.27 ± 5.18	p=0.588		
Lean (18.5-22.9)	20 (30.3)	10 (29.4)	NA		
Overweight (23.0-27.5)	14 (21.2)	6 (17.6)	NA		
Obese (>27.5)	32 (48.5)	15 (44.2)	NA		
No. (%) Previous still birth	10 (15.1)	9 (26.5)	p=0.07		
Mean HbA1C level	5.81 ±0.305%	8.188±1.71%	p=0.001		
Mean gestational age at current delivery (Weeks) No. (%)	37.4±2.3	34.2±3	p=0.006		
Gestational age at delivery					
<34 weeks	1 (1.5)	2 (5.8)			
34-36+6	9 (13.6)	11 (32.3)			
37-42	4 (81.8)	1 (44.1)			
Mean birth weight at current delivery (gms.)	3205±073	2803±098	p=0.019		

* WHO classification for Asian population

Table 2: Distribution of cases according to antenatal maternal complications.

Type of complication	Total	GDM	GDM (66)		DM (34)	Devolues	
	Total	Ν	%	n	%	P value	
Missed abortion	5	1	1.5	4	11.8	0.026#	
UTI	7	3	4.5	4	11.8	0.180	
Candidiasis	9	6	9	3	8.8	0.245	
DKA	1	0	0	1	2.9	0.160	
D. retinopathy	2	0	0	2	5.8	0.216	
D. nephropathy	1	0	0	1	2.9	0.734	
Preterm labour	9	3	4.5	6	17.6	0.030#	
PPROM	8	3	4.5	5	14.7	0.076	
Polyhydraminos	16	7	10.5	9	26.4	0.040#	
Pre-eclampsia	12	6	9	6	17.6	0.212	
PROM	9	7	10.5	2	5.8	0.434	
Total	79	37		42			
Uneventful	37/100	30	45	7	20.5	0.004#	

p<0.05 statistically significant</pre>

Table 3: Distribution of cases according to antenatal fetal complications.

Type of complication	Total	GDM	GDM (66)		DM (34)	P value	
		Ν	%	n	%	r value	
Congenital malformation	4	1	1.5	3	8.8	0.067	
IUFD	3	0	0	3	8.8	0.014#	
Macrosomia	15	11	16.7	4	11.8	0.884	
IUGR	2	0	0	2	5.8	0.216	
Uteroplacental insufficiency	2	0	0	2	5.8	0.216	
Total	26	12	14				
Uneventful	76/100	53	80.3	23	67.6	0.034#	

p<0.05 statistically significant

Table 4: Mode of delivery and intrapartum complications.

Type of complication	Total	GDM	(66)	Overt DM (34)			
	(100)	n	%	Overt DM II	Overt DM I	%	P value
NVD	44	33	50	10	1	32.3	0.12
LSCS	34	21	31.8	11	2	38.2	0.521
Instrumental Delivery	14	10	15.2	4	0	11.7	0.644
Intrapartum Hypoglycemia	1	0	0	0	1	2.9	0.738
Acute Fetal Distress	9	6	9	3	0	8.8	0.965
Meconium Stained Liqour	9	6	9	3	0	8.8	0.965
Stillbirth	2	2	3	0	0	0	0.305
Shoulder Dystocia	2	2	3	0	0	0	0.305

Table 5: Neonatal complications.

Neonatal complication	Total (92)	GDN	1	OVERT DM			
		Ν	%	OVERT DM II	OVERT DM I	%	P value
RDS	6	1	1.5	5	0	14.7	0.009#
Hypoglycemia	13	8	12	5	0	14.7	0.716
NNJ	15	9	13.6	5	1	17.6	0.595
LGA	8	4	6.06	3	1	11.7	0.602
Macrosomia (>4000gm)	15	12	18.2	3	0	8.8	0.2
LBW	10	4	6.06	5	1	17.6	0.067
Neonatal Period Uneventful	44	36	54.5	8		23.5	0.003#

p<0.05 statistically significant.</pre>

Of 100 cases, 66 cases had GDM and 34 cases had overt DM. Of 34 overt DM cases, 31 had Type II DM and 3 had Type I DM (IDDM) Figure 2. Characteristics of women in both the groups are shown in Table 1.

In present study around 88% cases were with age >25 years. The mean age was found to be higher in overt diabetics as compared to GDM women, 31.4 years versus 27.5 years respectively with p=0.010 (Table 1).

Women with GDM had lower BMI compared to overt diabetics, 24.72 ± 3.75 kg/m2 and 27 ± 5.18 kg/m2 respectively with p value 0.588 (Table 1).

In present study 31 cases (31%) had positive family history of diabetes. Positive family history in mother was significantly more commonly associated with GDM (21,

31.8%) than overt DM group (4, 11.7%) with p value 0.028.

Past history of infertility, early pregnancy failures and term IUFD were enquired, the difference was statistically significant only in relation with history of term IUFD, significantly higher in overt diabetics with p value 0.03 (Table 1).

HbA1c was found higher in overt diabetics compared to women with GDM, $8.18\pm 1.7\%$ and $5.89\pm 0.305\%$ respectively with p=0.001(Table 1).

Total 47 cases, 45 in GDM (68.2%) and 2 in overt DM group (5.9%) were managed with MNT alone while insulin therapy was supplemented in 41 cases, 16 in GDM group (24.2%) and 25 in overt DM group (73.5%) to achieve desired glycemic control.(Fig3)12 cases did not receive any treatment and constituted those cases who

were unbooked unsupervised cases and came in labour (2) or had some adverse pregnancy outcome like missed abortion (5)/CMF(4)/IUFD(1), where pregnancy needed to be terminated before instituting any treatment.

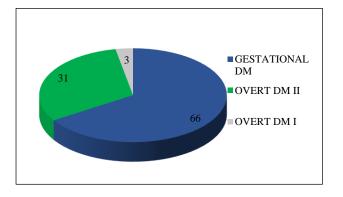


Figure 2: Distribution of cases according to type of diabetes.

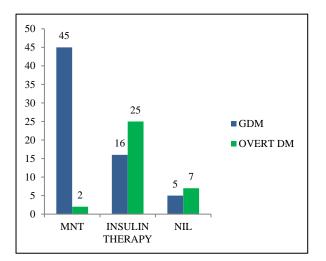


Figure 3: Distribution of cases according to treatment instituted.

Antenatal maternal complications developed in 63 cases (63%) (Table 2). Antenatal maternal complications noted were missed abortion (5, 5%), UTI (7, 7%), candidiasis (9, 9%), Diabetic Ketoacidosis (1, 1%), Diabetic Retinopathy (2, 2%), Diabetic Nephropathy (1, 1%), preterm labour (9, 9%), preterm PROM (8, 8%), polyhydraminos (16, 16%), pre-eclampsia (12, 12%) and PROM (9, 9%). Antenatal period was uneventful in significantly higher number of GDM cases (30, 45%) compared to overt diabetics (7, 20.5%) with p=0.004.

Among fetal complications observed were CMF in 4 cases, IUFD in 3, macrosomia in 15 cases, IUGR with Uteroplacental insufficiency in 2 cases. (Table 3).

CMF (4) in fetus were observed in 1case of GDM (1.5%) and 3 cases (8.8%) of overt diabetics. IUFD (3) was observed in 3 cases of overt diabetics (8.8%). Macrosomia (15) was observed in 11 cases of GDM (16.7%) and 4 cases of overt DM (11.8%). IUGR with Utero-placental

insufficiency (2) were observed in 2 cases of overt diabetics (5.8%).

Of total cases, 8 had 1st and 2nd trimester losses, 3 had early preterm delivery (POG <34 weeks), 20 had late preterm delivery (POG 34 weeks - 36+6weeks) and 69 had term delivery (POG 37 weeks - 42 weeks). The mean gestational age at termination of pregnancy was significantly earlier in overt DM group as compared to GDM group, 34.2 ± 3.3 weeks and 37.4 ± 2.3 weeks respectively with p value 0.006 (Table 1).

34 cases were delivered by LSCS, 14 cases had instrumental delivery while 44 cases had normal vaginal delivery. Suction and evacuation were done in 5 cases for missed abortion. LSCS rate were higher in women with overt diabetes (13, 38.2%) as compared to GDM (21, 31.8%) with p value 0.521. Instrumental delivery was found higher in GDM than overt diabetics 15.2 versus 11.8%, p=0.644 (Table 4)

Among intrapartum fetal complications observed, 9 fetuses developed acute fetal distress (9%), 9 had meconium-stained liquor (9%), 2 were stillborn (2%) and 2 had shoulder dystocia (2%) (Table 4).

Shoulder dystocia (2) were observed in 2 cases of GDM (3%) and none in overt diabetics with p value 0.305. Still birth (2) was observed in 2 cases of GDM group (3%) and none in overt diabetics (0%) with p=0.305.

In present study out of total 92 new-borns, 10 had birth weight <2500 gm (10%), 67 had birth weight between 2500 to 4000 gm (67%) and 15 had birth weight >4000 gm (15%). The mean birth weight of new-borns was 3035 ± 855 gm. New-borns born to GDM mothers were found significantly heavier than overt diabetics, with mean birth weight 3250 ± 730 gm and 2820 ± 980 gm respectively with p=0.019. (Table 1).

Macrosomia (birth weight >4000 gm) (15) was observed in 12 neonates of GDM mothers (18.2%) and 3 neonates of overt diabetes (8.8%) with p value 0.214. LGA (birth weight greater than 90th centile) (8) was noted in 4 newborns of GDM mothers (6.06%) and 4 new-borns of overt diabetics (11.7%) with p=0.602. LBW (birth weight <2500 gm) (10) was detected in 4new-borns of born to GDM mothers (6.06%) and 6 new-borns born to mothers with overt diabetes (17.6%) with p value 0.067 (Table 5).

RDS (6) was noted in significantly higher in neonates of overt diabetics (5, 14.7%) compared to GDM women (1, 1.5%) with p=0.009. Neonatal hypoglycemia (13) was observed in 8 new-borns of GDM mothers (12%) and 5 new-borns of overt diabetics (14.7%) with p=0.716. NNJ (15) was observed in 9 new-borns of GDM mothers (13.6%) and 6 new-borns of overt diabetics (17.6%) with p value 0.595 (Table 5).

Out of 92 cases, one had multiple congenital anomalies incompatible with life, 3 had intrauterine fetal demise and 2 were still born. Rest 86 neonates were normal and euglycemic at the time of discharge from the hospital.

DISCUSSION

The present study prevalence of diabetes complicating pregnancy was found to be 1.79% (116/6452). Kavyashree et al 2014 in Mysore, Karnataka reported similar prevalence rate of 1.64% (50/3044) of which 90% had GDM, 6% were overt diabetics type I and 4% were overt diabetics type II.⁹ In studies conducted in various other parts of country, prevalence of GDM was found to be 2.6% by Gajjar et al 2005 in Baroda, Gujarat; 3.6% by Sawant et al 2012 in Loni, Maharashtra.^{10,11}

The mean age was found to be higher in overt diabetics as compared to GDM women, 31.4 years versus 27.5 years respectively with statistically significant difference with p=0.010. Age >25 years is a recognised risk factor for developing GDM in pregnancy. In present study around 88% cases were with age >25 years. In a study by Shefali et al 2006 mean age in overt diabetics was 29 ± 6 years and in GDM was 29 ± 5 years which was comparable to our study.¹² In present study 31 cases were found to have positive family history of diabetes (31%). 23 cases of GDM (34.8%) and 8 cases of overt diabetes (23.4%) had associated positive family history of DM. It was close to that found by study by Patil et al 2014 and, Kalra et al 2013.^{13,14}

In the present study, LSCS rate were higher in women with overt diabetes (13, 38.2%) as compared to GDM (21, 31.8%) with p value 0.521. In studies by Wahabi et al 2014 and Abu Heijaet al 2015 caesarean rate in overt diabetics and GDM was 46.6% versus 26% and 60.3% versus 27.9% respectively.^{15,16} While in a study by Patil et al 2014 caesarean rate was reported higher in GDM than overt diabetics (48% versus 30%).¹⁷ Also, in several studies conducted in various parts of country on GDM, very higher rate of caesarean section were reported than our study and the rate has increased dramatically in recent past.

In present study instrumental delivery rate were found higher in GDM than overt diabetics 15.2% versus 11.8%, p=0.644. In studies by Gajjaret al 2005, Kalyani et al20 2014 and Swaroop et al 2015 rate of instrumental delivery was 11%, 12% and 18% in GDM respectively which was comparable to our study.^{10,19} In a study by Kalraet al 2013 reported rate of instrumental delivery was 3% which was lower than our study.¹⁴

In present study pre-eclampsia was observed in 12 cases, 6 in GDM group (9%) and 6 in overt DM group (17.6%). Thus, pre-eclampsia was found higher in mothers with overt DM with p=0.212. Studies by Sugiyama et al 2014 and Abu Heijaet al 2015 reported rate of pre-eclampsia of 6.1% versus 10.1% and 7.8% versus 17.2% in GDM and

overt diabetics respectively which was comparable to our study. $^{16,19} \,$

In present study polyhydraminos was observed in 16 cases, 7 in GDM (10.6%) and 9 in overt diabetics (26.4%). This was significantly higher in overt diabetics with p value 0.040. In studies by Patil et al 2014 and Abu Heijaet al 2015 incidence was 5.2% versus 7.6% and 6.8% versus 4.9% in overt diabetics and GDM respectively.^{16,17} Sawant et al 2012 and Kavyashreeet al 2014 reported incidence of polyhydraminos in GDM 16.6% which was higher than our study.^{9,11} Poor glycemic control and poor compliance to treatment especially in third trimester increases the rate of polyhydraminos. In present study preterm labour (PTL) was observed in 9 cases, 3 in GDM (4.5%) and 6 in overt diabetic cases (17.6%) significantly higher in overt diabetics with p value 0.03. This finding was comparable with other studies done by Wahabi et al 2014 (18% versus 8.5 %) and Heijaet al 2015 (25.9% versus 9.5 %).^{15,16}

In present study CMF in fetus was observed in 4 cases (4%), 1 in GDM (1.5%) and in 3 fetuses of overt diabetic mothers (8.8%) with p=0.067. While Sawant et al 2012 and Swaroop et al 2015 reported CMF rate of 11.11% and 13.6% respectively in GDM which was higher than our study.11,17 CMF in DM is associated with severity of hyperglycemia during period of organogenesis. So, CMF are found higher in overt diabetics. Pinter et al studied the mechanism by which hyperglycemia produced dysmorphogenesis and proposed it to be hyperglycemia induced yolk sac failure and resulting compromise of nutrient transfer and oxygenation to the early embryo.¹⁸ Good glycemic control in preconception and early pregnancy has major role in prevention of CMF in pregnancies complicated by diabetes.

In present study new-borns born to GDM mothers were heavier than overt diabetics, the difference was found to be statistically significant with p=0.019. In present study macrosomia (birth weight >4000 gm) was detected in 15 neonates (92). 12 neonates having macrosomia were born to GDM mothers (18.2%) and 3 neonates with macrosomia were born to mothers with overt diabetes (8.8%). Kalra et al 2013 reported comparable rate of 18% in their study in GDM.¹⁴ In present study 8 new-borns were LGA (birth weight greater than 90th centile) (8%). This was noted in 4 new-borns of GDM mothers (6.06%) and 4 new-borns of overt diabetics (11.7%) with p=0.602. It is very important to find large for gestational age neonates as they remain at a higher risk of hypoglycemia in neonatal period and also at a higher risk of developing childhood obesity and early onset diabetes mellitus later in life. In a study Shefali et al 2006 reported the incidence of 27.6% and 19.2% in GDM and overt diabetics respectively which was higher than our study.12

In present study RDS was observed significantly higher in neonates of overt diabetics with p=0.009. Heija et al 2015 also reported higher rate of RDS in overt diabetics (8.5 versus 2.6%) while Patil et al 2014 reported higher rate in GDM compared to overt diabetics (11.6 versus 5.1%).^{16,17} In similar study by Kavyashree et al 2014 incidence of RDS was 12.5% which was comparable to our study.⁹

Limitations

The present study was an institutional based study. Thus the prevalence of diabetes in pregnancy found in this study may not be exact representation of the study population. There was no control group taken in the study thus comparison with non-diabetic data lacking in the present study.

CONCLUSION

Timely screening, early detection of GDM and overt DM in pregnancy, regular close follow ups and institutional delivery of these pregnant women will improve pregnancy outcome. Preconceptional counselling of women with risk of hyperglycaemia in pregnancy ensures that she enters pregnancy with an optimal state of health. The need of the hour is to make preconceptional counselling a 'done thing' in India too as in western countries. Large multicentric population based study should be carried out to know the disease burden in the study population and to further strengthen the health system to combat the adverse effect of diabetes on maternal and fetal outcome.

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