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ABSTRACT Background: Diabetes is estimated to complicate 2-5% of all pregnancies. 90% of those are detected during pregnancy i.e. Gestational Diabetes Mellitus (GDM) and the rest are overt or pregestational i.e. either type I or type II. Increased awareness and availability of many screening methods detected Gestational Diabetes Mellitus early and controlled further effects on the mother or fetus and also prevented long term complications to both mother and fetus. In this study we have evaluated the efficacy of RBS and GCT as screening test for GDM.

Aims of the Study

To evaluate the predictive value of Glucose Challenge Test and Random Blood Sugar as a screening test in pregnant women between 24-28 weeks in Low risk group and in High risk group at the first antenatal visit, if negative at 24-28 weeks and, if negative at 32-36 weeks of GA.

To evaluate the preference of two screening tests RBS and OGCT for $\ensuremath{\mathsf{GDM}}$

Materials and methods

The prospective clinical study is done at GGH, Kakinada, over a study period of 1½ year. 200 Antenatal women between the gestational ages 24-28 weeks were screened for oral glucose challenge test (OGCT) Vs random blood sugar (RBS). The study group was divided into high risk and low risk group. All the women are subjected to both OGCT and RBS. Antenatal women were divided into low risk and high risk group according to inclusion and exclusion criteria. Women with the values above the cut off values were subjected to glucose tolerance test (GTT) for confirmation of diabetes.

Results:

In high risk group of 75 (100% antenatal women, 27 (36%) were screened positive by OGCT and 6 (22.22%) were diagnosed GDM by OGTT. Of them 3 were diagnosed GDM at 32-36 weeks, 2 were diagnosed GDM at 24-28 weeks and one was diagnosed GDM at First AN visit. In OGCT screen positive patients of high risk group- 4(6.55%) were in the age group between 26-30 years among 61, one (9.09%) was in the age group between 31-35 years among 11 and one (33.33%) was in the age group >35 years among 3 diagnosed as GDM.

In the low risk group of 125(100%) antenatal women, 7(5.6%) women were screened positive with OGCT and 2 (28.57) were diagnosed GDM by OGTT. In OGCT screen positive of low risk group- One (3.70%) was in the age group <20 years among 27 and one (1.02%) was in the age group of 21-25 years among 98 was diagnosed as GDM.

In high risk group of 75 (100%) antenatal women 31 (41.33%) were screened positive by RBS and 3 (9.67%) per diagnosed as GDM by OGTT. Of them 2 were diagnosed GDM at 32-36 weeks, one was diagnosed between 24-28 weeks and none was diagnosed GDM at First AN

visit. One (1.63%) was in the age group of 26-30 years among 61, One (9.09%) was in the age group 31-35 years among 11 and one (33.33%) was in the age group of >35 years among 3.

In the low risk group of 125 (100%) antenatal women, 5(4%) women were screened positive with RBS. In RBS screen positive of low risk group none was in the age group <20 years and 1(1.02%) in the age group between 21-25 years among 98 diagnosed as GDM. Out of 200 AN cases, 85 were primigravida of which one was diagnosed as GDM. 73 were second gravida of which 1 was diagnosed as GDM. 30 were third gravida of which 3 were diagnosed as GDM. 10 were fourth gravida of which 2 were diagnosed as GDM. 2 were fifth gravida of which 1 was diagnosed as GDM.

The sensitivity was 100% in both high risk and low groups screened with OGCT and RBS. Specificity was 69% for high risk OGCT group, 61% for the high risk RBS group, 96% for low risk OGCT group, 96.7% for low risk RBS group.

Conclusion:

- OGCT could diagnose GDM more accurately than RBS, the same is confirmed by OGTT.
- Positive member of women Screened by RBS were more but when OGCT was done in RBS Positive cases, Positive GDM cases are low. This indicates RBS evaluation gave rise to more number of false Positive cases. So when we compare the efficacy of RBS and OGCT, OGCT is better screening test.
- As the parity and age of the antenatal women increases number of Gestational Diabetes Mellitus increases.

Until superior alternatives become available the 50gm glucose challenge test should be preferred screening test for GDM. GCT is a better investigation for the screening of gestational diabetes than random blood glucose.

Universal screening for gestational diabetes mellitus should be mandatory irrespective of presence or absence of risk factors because it is definite disease entity associated with significant maternal and perinatal complications.

Materials and methods

- The prospective clinical study is done at GGH, Kakinada, over a study period of 1½ year.
- 200 Antenatal women between the gestational ages 24-28 weeks were screened for oral glucose challenge test (OGCT) Vs random blood sugar (RBS). The study group was divided into high risk and low risk group. All the women are subjected to both OGCT and RBS.
- Cut off value for OGCT was taken as 140 mg/dl.
- Cut off value for RBS was taken as100 mg/dl.
- Antenatal women were divided into low risk and high risk group according to inclusion and exclusion criteria.
- Women with the values above the cut off values were subjected to glucose tolerance test (GTT) for confirmation of diabetes.

INCLUSION CRITERIA

High Risk Group:

- Age > 25 years
- Family history of diabetes in first degree relative
- Marked obesity-BMI >2 Kg/m² or >120% ideal body weight
- Previous abnormal glucose tolerance test
- Previous large baby > 4Kg
- Persistent glycosuria
- Previous bad obstetric history- unexplained still birth or congenital malformed babies, unexplained perinatal loss, intrauterine death, Preterm delivery.
- Polyhydramnios
- Previous h/o of preeclampsia
- Previous h/o of GDM

Low risk group:

- Age < 25 years
- No known diabetes in first degree relatives
- Weight normal before pregnancy
- Weight normal at birth
- No history of abnormal glucose metabolism
- No previous history of adverse obstetrical outcome usually associated with gestational diabetes [macrosomia, neonatal hypoglycemia]

EXCLUSION CRITERIA

• Women with known preexisting diabetes (overt)

METHOD OF PERFORMING OGCT

- Fasting was not a prerequisite.
- Irrespective of the time of last meal, 50gms of glucose was dissolved in 200ml of water and asked to drink within 5 min.
- Exactly after 1 hour, venepuncture was made and blood obtained for the study
- Plasma glucose was estimated with glucose oxidase or Hexokinase reagent test
- If result >140mg/dl, then women was subjected to three hour OGTT with 100gm of glucose.

METHOD OF PERFORMING RBS

- Fasting was not a prerequisite
- Venepuncture was made and blood obtained for the study
- Plasma glucose was estimated with glucose oxidase or Hexokinase reagent test
- If the result was >100mg/dl, then the women underwent three hour OGTT with 100gm of glucose.

METHOD OF PERFORMING OGTT

• For at least three days prior to the test, women were

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asked to consume their normal unrestricted diet containing a minimum of 150gm of carbohydrate.

- After an overnight fast of 8 hours, a fasting blood sample is drawn; following which she drank a solution of 100gm of glucose dissolved in 300ml of water within 5 minutes.
- First hour, second hour and third hour samples were collected by vene puncture.
- Plasma glucose estimated with glucose oxidase or Hexokinase reagent.
- Values of four plasma glucose obtained are compared with values done for 100gm OGTT of carpenter and coustan criteria.
- If 2 values were greater than the values of carpenter and coustan criteria, woman was labeled as GDM.

Timing of plasma glu- cose collection	Carpenter and coustan criteria
Fasting	95 mg/dl
1 hour	180 mg/dl
2 hour	155 mg/dl
3 hour	140 mg/dl

SCREENING OF BLOOD SUGARS VALUES TESTING METHOD

Sample collection and processor:

- Intravenous blood of 2ml is collected after taking oral glucose 50gm or 100gm according to the test – 50 gm OGCT and 100gm OGTT.
- Intravenous blood of 2ml is collected irrespective of glucose intake i.e. Random Blood Sugar.
- After separation of serum from sample, it is centrifuged at rate of 3000 rpm for 5min.
- After 5min the test sample is kept in auto analyzer (Glucose Oxidase or Glucose Hexokinase for recording of blood sugar values and results are taken accordingly.

Principle:

The reaction sequence employed in the assay of glucose is as follows:

	Glucose Oxidase	
1. Glucose + 1/2 O2 -	H20 → Gluconate + 1	H_2O_2

Peroxidase

Glucose is oxidized by glucose oxidase and produces gluconate and hydrogen peroxide. The hydrogen peroxide is then oxidatively coupled with 4 – aminoantipyrine and phenol. The amount of coloured complex (quinoneimine) is proportional to glucose concentration in sample that can be measured photometrically.

STATISTICAL METHODS

The Diagnostic values were computed for OGCT and RBS with respect to the final diagnosis.

TEST	GDM	NOT GDM	TOTAL
ABNORMAL	а	b	a + b
NORMAL	с	d	c + d

RESEAR	RCH I	PAPER				
TOTAL		a + c	b + d	n		
Sensitivity=	Sensitivity= a / (a + c)					
Specificity=	= d /	(b + d)				
PPV =	= a/	(a + b)				
NPV =	= d /	(c + d)				

Statistical Software: Microsoft word and Excel have been used to generate graphs, tables etc

Observations and results TABLE 1: OGCT SCREENING IN HIGH RISK GROUP

SCREENING			TOTAL
			n=75(100%)
POSITIVE	6 (22.2%)	21 (77.77%)	27 (36%)
NEGATIVE	0	48 (100%)	48 (64%)

The table shows the number of women who underwent OGCT in the high risk group and number of patients who were diagnosed to be GDM.

Total number of high risk cases is 75. Out of them 27 (36%) antenatal women were OGCT screen positive. These 27 cases underwent OGTT, of them 6 (22.22%) were diagnosed to be GDM and remaining 21 (77.77%) were negative - no Gestational Diabetes Mellitus.

Out of 75 antenatal cases, 48 antenatal women (64%) who were OGCT negative did not undergo OGTT and were considered as having no GDM.

TABLE	2:	OGCT	SCREENING	IN	LOW	RISK	GROUP
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SCREENING	GDM (OGTT)	NO GDM	TOTAL n= 125(100%)
POSITIVE	2 (28.57%)	5 (71.42%)	7 (5.6%)
NEGATIVE	0	118 (100%)	118 (94.4%)

Table 2 shows the number of woman who underwent OGCT in the low risk group and number of patients who were diagnosed to be GDM.

Total number of low risk cases is 125. Out of them 7 (5.6%) antenatal women were OGCT positive. These 7 cases underwent OGTT, of them 2 (28.57%) were diagnosed to be GDM and remaining 5 (71.42%) were negative- no GDM.

Out of 125 antenatal cases, 118 women (94.4%) were OGCT negative and did not undergo OGTT and considered them as having no GDM.

TABLE 3: RBS SCREENING IN HIGH RISK GROUP

SCREENING	GDM (OGTT)	NO GDM	TOTAL n=75(100%)
POSITIVE	3 (9.67%)	28 (90.32%)	31 (41.33%)
NEGATIVE	0	44 (100%)	44 (58.67%)

Table 3 shows the number of woman who underwent RBS in the high risk group and number of patients who were

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diagnosed to be GDM.

Total number of high risk cases is 75. Out of them 31 (41.33%) antenatal women were OGCT positive. These 31 cases underwent OGTT, of them 3 (9.67%) were diagnosed to be GDM and remaining 28 (90.32%) were negative - no GDM.

Out of 75 cases, 44 antenatal women (58.67%) who were OGCT negative did not undergo OGTT and considered them as having no GDM.

SCREENING	gdm (ogtt)	NO GDM	TOTAL n= 125(100%)
POSITIVE	1 (20%)	4 (80%)	5 (4%)
NEGATIVE	0	120 (100%)	120 (96%)

TABLE 4: RBS SCREENING IN LOW RISK GROUP

Table 4 shows the number of woman who underwent RBS in the low risk group and number of patients who were diagnosed to be GDM.

Total number of low risk cases is 125. Out of them 5 (4%) were OGCT positive. These 5 cases underwent OGTT, of them 1 (20%) were diagnosed to be GDM and remaining 4 (80%) were negative - no GDM.

Out of 125 antenatal women, 120 women (96%) who were OGCT negative did not undergo OGTT and considered them as having no GDM.

Visit	OGCT	OGTT (No. AN cases)		OGTT %	
	Positive	Positive	Nega- tive	Positive	Nega- tive
At First AN Visit	8 (10.66%)	1	7	12.5%	87.5%
At Second AN Visit (24- 28 weeks)	9 (13.4%)	2	7	22.22%	77.7%
At Third AN Visit (32.36 weeks)	10 (17.24%)	3	7	30%	70%

TABLE 5: ASSOCIATION OF GDM WITH GESTATIONAL AGE IN OGCT OF HIGH RISK GROUP

At first antenatal visit, out of 75 antenatal women 8 i.e. 10.66% women were screened positive by OGCT. These 8 women were subjected to OGTT, of them 1 (12.5%) was diagnosed to be diabetic and 7 (87.5%) had no gestational diabetes mellitus. Remaining 67 women in high risk group who were OGCT negative at first antenatal visit were subjected to OGCT at 24-28 weeks; of them 9 (13.4%) women were screened positive by OGCT. These 9 antenatal women were subjected to OGTT, of them 2 (22.22%) diagnosed to be diabetic and 7 (7.77%) had no gestational diabetes mellitus.

Remaining 58 women in high risk group who were OGCT negative at 24 – 28 weeks were subjected to OGCT at 32 – 36 weeks, of them 10 (17.24%) women were screened positive by OGCT. These 10 antenatal women were subjected to OGTT, of them 3 (30%) diagnosed to be diabetic

and 7 (70%) had no gestational diabetes mellitus.

Visit	RBS	OGTT (No. AN cases)		OGTT%	
	Positive	Positive	Nega- tive	Posi- tive	Nega- tive
At First AN Visit	11 (14.66%)	0	11	0%	100%
At Second AN Visit (24- 28 weeks)	11 (17.18%)	1	10	9.09%	90.9%
At Third AN Visit (32.36 weeks)	9 (16.98%)	2	7	22.2%	77.7%

TABLE 6: ASSOCIATION OF GDM WITH GESTATIONAL AGE IN RBS OF HIGH RISK GROUP

At first antenatal visit, out of 75 antenatal women 11 (14.66%) women were screened positive by RBS. These 11 women were subjected to OGTT, and all of them had no gestational diabetes mellitus. Remaining 64 women in high risk group who were RBS negative at first antenatal visit were subjected to RBS at 24-28 weeks; of them 11(17.18%) women were screened positive by RBS. These 11 antenatal women were subjected to OGTT, of them 1 (9.09%) diagnosed to be diabetic and 10 (90.9%) had no gestational diabetes mellitus.

Remaining 53 women in high risk group who were RBS negative at 24 - 28 weeks were subjected to RBS at 32 -36 weeks; of them 9 (16.98%) women were screened positive by RBS. These 9 antenatal women were subjected to OGTT, of them 2 (22.22%) diagnosed to be diabetic and 7 (77.77%) had no gestational diabetes mellitus.

TABLE 7: COMPARISION OF LOW RISK AND HIGH RISK GROUP

	HIGH RISK		LOW RISK	
	n=75		n=125	
Screening	Total	GDM	Total	GDM
Test	(Screen Positive)	(OGTT)	(Screen Positive)	(OGTT)
	27	6	7	2
OGCT	(36%)	(22.2%)	(5.6%)	(28.57%)
	31	3	5	1
RBS	(41.33 %)	(9.67%)	(4%)	(20%)

Shows the number of women screened and diagnosed GDM in each group.

In the OGCT high risk group, 27 (36%) antenatal women of 75 were OGCT screen positive. These 27 antenatal women underwent OGTT and of them 6 (22.2%) were found to be diabetic.

In the RBS of high risk group, 31 (41.33%) antenatal women of 75 were RBS screen positive. These 31 antenatal women underwent OGTT and 3 (9.67%) were found to be diabetic.

In the OGCT low risk group, 7 (5.6%) antenatal women of 125 were OGCT screen positive. These 7 antenatal women underwent OGTT and 2 (28.57%) were found to be diabetic.

In the RBS low risk group 5 (4%) antenatal women of 125 were RBS screen positive. These 5 antenatal women underwent OGTT and 1 (20%) was found to be diabetic.

NISK/			
SCREENING	TOTAL		
(OGCT)	n=200 (100%)		
POSITIVE	34	26	8
	(17%)	(76.47%)	(23.52%)
	166	166	0
NEGATIVE	(83%)	(100%)	U

TABLE 8: SCREENING WITH OGCT (High Risk + Low D:-1-1

Table 8 shows the number of patients who underwent screening in the OGCT group including both high risk and low risk factors. Total of 34 out of 200 antenatal women (17%) were OGCT screen positive. These 34 antenatal women underwent OGTT of whom 8 (23.52%) were diagnosed to have GDM and 26 (76.47%) did not have GDM. Remaining 166 i.e. 83% women did not undergo OGTT and considered them as having no GDM.

TABLE 9: SCREENING WITH RBS (High Risk + Low Risk)

SCREENING	TOTAL			
(RBS)	n=200			
	36	32	4	
FUSITIVE	(18%)	(88.88%)	(11.11%)	
	164	164	0	
	(82%)	(100%)	U	

Table 9 shows the number of patients who underwent screening in the RBS group including both high risk and low risk factors. Total of 36 out of 200 antenatal women (18%) were RBS screen positive. These 36 antenatal women underwent OGTT of whom 4 (11.11%) were diagnosed to have GDM and 32 (88.88%) did not have GDM. Remaining 164 i.e. 82% women did not undergo OGTT and considered them as having no GDM.

TABLE 10: TOTAL NUMBER OF ANTENATAL WOMEN ACCORDING TO AGE GROUPS

	No. of Antenatal women	
AGE GROUP	(n=200)	
<20	27 (13.5%)	
21-25	98 (44%)	
26-30	61 (30.5%)	
31-35	11 (5.5%)	
>35	3 (1.5%)	

Figure - 1



Most of the women in the study group were between 21-25 (49%) years of age. There were 98 women screened for OGCT and RBS belonging to this group. Women between 26-30 years were 61 (30.5%). 27 (13.5%) women screened for OGCT and RBS who were less than 20 years of age. In age group of 31-35 years there were 11 (5.5%) patients and in the age group of >35 years there were 3(1.5%) antenatal women who were screened for both GCT and RBS.

TABLE 11:COMPARISIONOFOGCTANDRBSSCREENINGFORGDMACCORDINGTOAGEGROUPSANDNUMBEROFANTENATALCASES

AGE (YEARS)	TOTAL NO. OF ANTENA- TAL CASES	OGCT (GDM)	RBS (GDM)
<20	27	1 (3.70%)	0
21-25	98	1 (1.02%)	1 (1.02%)
26-30	61	4 (6.55%)	1 (1.63%)
31-35	11	1 (9.09%)	1 (9.09%)
>35	3	1 (33.33%)	1 (33.33%)

TABLE 12: OGCT SCREENING IN HIGH RISK ACCORD-ING TO AGE GROUPS FOR GDM

HIGH- RISK	26-30 YEARS		31-35 YEARS		>35 YEARS	
TOTAL n = 75	61		11		3	
CDM	Positive	Nega- tive	Positive	Nega- tive	Positive	Nega- tive
(OGTT)	4 (6.55%)	57 (93.44%)	1 (9.09%)	10 (90.9%)	1 (33.33%)	2 (66.66%)

TABLE 13: OGCT SCREENING IN LOW RISK ACCORD-ING TO AGE GROUPS FOR GDM

LOWRISK	<20 YEARS	5	21-25 YEARS	
TOTAL	27		0.0	
n = 75	27		70	
GDM	Positive	Negative	Positive	Negative
	1	26	1	97
(OGTT)	(3.70%)	(96.3%)	(1.02%))	(98.98%)

Figure - 2



In the high risk group, out of 75 women screened by OGCT, 61 (30.5%) women were in the age group of 26-30 years, 4 (6.55%) patients were diagnosed as GDM by

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OGTT. Out of 11 (5.5%) women in the age group 31-35years, 1 (9.09%) patient was diagnosed as GDM by OGTT. Out of 3 (1.5%) women in the age group >35 years, 1 (33.33%) patient was diagnosed as GDM by OGTT. In the low risk group, out of 125 women screened by OGCT, 27 (13.5%) women in age group of less than 20years, 1(3.7%) patient was diagnosed as GDM by OGTT. Out of 98 (49%) women in the age group of 21-25 years, 1 (1.02%) patients was diagnosed as GDM by OGTT.

TAE	BLE	14:	RBS	SCREE	ENING	IN	HIGH	RISK	ACCOR	RDING
то	AG	ΕG	ROU	PS FO	r gdiv	1				

HIGH- RISK	26-30 YEARS		31-35 YEARS		>35 YEARS	
TOTAL n = 75	61		11		3	
GDM	Positive	Nega- tive	Posi- tive	Nega- tive	Positive	Negative
(OGTT)	1 (1.63%)	60 (98.37%)	1 (9.9%)	10 (90.9%)	1 (33.33%)	2 (66.66%)

TABLE 15: RBS SCREENING IN LOW RISK ACCORDING TO AGE GROUPS FOR GDM

LOWRISK	<20 YEARS		21-25 YEARS	
TOTAL n = 75	27		98	
CDM	Positive	Negative	Positive	Negative
(OGTT)	0	27 (100%)	1 (1.02%)	97 (98.98%)





In the high risk group, out of 75 women screened by RBS, 61 (30.5%) women were in age group of 26-30 years, 1 (1.63%) patient was diagnosed to have GDM by OGTT. Out of 11 (5.5%)women in the age group 31 -35 years, 1(9.09%) patient was diagnosed to have GDM by OGTT. Out of 3 (1.5%) women in the age group 35years, 1(33.33%) patient was diagnosed to have GDM by OGTT.

In the low risk group out of 125 women screened by RBS, 27(13.5%) women were in the age group < 25years, out of which no one had GDM by OGTT. Out of 98 (49%) women in age group of 21-25 years, 1 (1.02%) patient was diagnosed to have GDM by OGTT.

TABLE 16: COMPARSION OF DIAGNOSTIC VALUES OF OGCT AND RBS AS PER RISK GROUP.

	High Risk OGCT in %	High Risk RBS in %	Low Risk OGCT in %	Low Risk RBS in %
Sensitivity	100	100	100	100
Specificity	69	61	95.9	96.7
PPV	22	9.6	28	20
NPV	100	100	100	100

Figure - 4



The table shows comparison of diagnostic values of OGCT and RBS on high and low risk group. The sensitivity is 100% in all 4 groups with variations in the specificity. RBS Low risk group had high specificity of 96.7%, with OGCT low risk group had 95.9% and OGCT high risk group 69%, RBS high risk group 61%.

OGCT low risk group had high PPV 28%, RBS 20% and OGCT high risk group had 22%, RBS 9.6% and NPV has 100% in all the four groups.

TABLE 17: PARITY DISTRIBUTION IN THE STUDY GROUP

GRAVIDA	TOTAL NO. OF ANTEN- TAL CASES N = 200	GDM in total number of cases as per Gravida	Percentage GDM per Gravida in total GDM cases
Primi	85 (42.5%)	1 (1.17%)	12.5%
2 nd Gravida	73 (36.5%)	1 (1.36%)	12.5%
3 rd Gravida	30 (15%)	3 (10%)	7.5%
4 th Gravida	10 (5%)	2 (20%)	25%
5 th Gravida	2 (1%)	1 (50%)	12.5%

Primigravidas were more in the study group. 85 (42.5%) women were primigravidas of which 1 (1.17%) woman was diagnosed to have GDM by OGTT. 73 (36.5%) women were 2^{nd} gravida of which 1 (1.36%) woman was diagnosed to have GDM by OGTT. 30 (15%) women were 3^{nd} gravida of which 3 (10%) women were diagnosed to have GDM by OGTT. 10 (5%) women were 4^{th} gravida of which 2 (20%) women were diagnosed as GDM. 2 (1%) women were 5^{th}

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gravida of which 1 (50%) woman was diagnosed to have GDM by OGTT.

Among 8 GDM cases diagnosed 1 case was primi gravida (12.5%), 1 case was 2^{nd} gravida, 3 cases (37.5%) were 3^{rd} gravida, 2 cases (25%) were 4^{th} gravida and 1 case (50%) was 5^{th} gravida.

TABLE 18: COMPARISON OF RISK FACTORS

Risk Factors	No of Antenatal Women
Family H/O Diabetes Mellitus	15 (20%)
Previous H/O Perinatal Mortality	11 (14.66%)
Previous H/O IUD	13 (17.33%)
Previous H/O LGA	4 (5.33%)
Previous H/O Anamolous Foetus	1 (1.33%)
Previous H/O of Preterm	5 (6.66%)
Polyhydramnios	6 (8%)
Previous H/O Pre Eclampsia	10 (13.3%)
Glucosuria	8 (10.66%)
BMI (>27 kg/m²)	16 (21.32%)





Family H/o Diabetes Mellitus	Previous H/O Perinatal mortality
Previous H/O IUD	Previous H/o LGA
Previous H/o Anamolous Foetus	Previous H/o of Preterm
Polyhydraminos	Previous H/o Pre eclampsia
■ Glycosuria	BMI (>27kg/m2

In high risk group there were 15 (20%) with family H/o. of diabetes mellitus, 11 cases (14.66%) with previous H/o. of perinatal mortality, 13 cases (17.33%) with previous H/o. of IUD, 4 cases (5.33%) with previous H/o. of LGA, 1 case (1.33%) with previous H/o. of Anomolous foetus, 5 cases (6.66%) with previous H/o. preterm, 6 cases (8%) with polyhydraminos, 10 cases (13.3%) with previous H/o. of Eclampsia, 8 cases (10.66%) with Glucosuria, 16 cases (21.32%) with BMI >27 kg/m²). More than One risk factor is present in some cases.

TABLE	19:	ASSOCIATION	OF	OUTCOME	IN	GDM	PA-
TIENTS	i						

	OUTCOME		Number
	Live	Dead	
FTND	4	-	4
Instrumental	1	-	1
Preterm Vaginal Delivery	1	-	1
LSCS	1	-	1
Still Born	1	1	1

There were 4 FTND, INSTRUMENTAL deliveries were 1, 2 patients underwent LSCS, 1 STILL BIRTH with GDM.

Figure - 6



Conclusion

India has become diabetic epidemic due to its fast urbanization, life style changes and genetic make up of Indians. Diabetic mellitus is estimated to complicate 2-5% of all pregnancies. 90% of those cases are detected during pregnancy-called as Gestational Diabetes Mellitus. Approximately 7% of all pregnancies are complicated by Gestational Diabetes Mellitus resulting in greater than 2 lakh cases per annum. To forecast Gestational Diabetes Mellitus, we need to have a strong screening test. Such available screening tests are:

- Glycosuria
- Blood glucose estimation
- RBS
- FBS
- PPBS
- Mixed nutrient meal
- Oral glucose challenge test
- Glycosylated hemoglobin (HbA1c) estimation
- Spot test
- Glucose polymer challenge test
- Fructosamine estimation

We need to have fast, simple, reliable, relatively inexpensive tests. Such two tests are RBS and OGCT.

In this study we have evaluated the efficacy of RBS and GCT as screening test for GDM.

OGCT could diagnose GDM more accurately than RBS, the same is confirmed by OGTT.

Positive member of women Screened by RBS were more but when OGCT was done in RBS Positive cases, Positive GDM cases are low. This indicates RBS evaluation gave rise to more number of false Positive cases. So when we compare the efficacy of RBS and OGCT, OGCT is better screening test.

As the parity and age of the antenatal women increases number of Gestational Diabetes Mellitus increases.

Until superior alternatives become available the 50gm glucose challenge test should be preferred screening test for GDM. GCT is a better investigation for the screening of gestational diabetes than random blood glucose.

Universal screening for gestational diabetes mellitus should

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be mandatory irrespective of presence or absence of risk factors because it is definite disease entity associated with significant maternal and perinatal complications.

REFERENCES:

- 1. Williams Text Book of Obstetrics 23rd edition pg.1106,1113,1114
- 2. Blot: Comptes-rendus de la soc de biologie. 1883; 5: 193.
- Graham G: A case of diabetes mellitus complicated by pregnancy treated with insulin. Proc Roy Soc Med 1924; xvii: 102-4.
- O'Sullivan JB, Mahan CM: Criteria for the oral glucose tolerance test in pregnancy. Diabetes 1964; 13: 278-85.
- Expert committee on the diagnosis and classification of diabetes mellitus. Report on the expert committee on the diagnosis and classification of diabetes mellitus. Diabetes Care 1997;20:1183-97
- American college of Obstetricians and Gynaecologist. Gestational Diabetes. acog practical bulletin 2001;(30).
- First International workshop-conference on Gestational Diabetes mellitus: summary and recommendations, diabetes care 1980; 3:499-501.
- American Diabetes Association: summary and recommendations of second international workshop-conference on gestational diabetes mellitus. Diabetes 1985, 34(suppl .2):123-126.
- American college of obstetricians and gynaecologist. Diabetes and Pregnancy. International journal of gynaecology and obstetrics 1985; 48:331-2.
- Metzger BE, The organizing committee: summary and recommendations of third international workshop-conference on Gestational Diabetes Mellitus. Diabetes 1991, 40(suppl.2):197-201.
- Metzger BE, Coustan DR. Summary and recommendations of the Fourth International Workshop-Conference on Gestational Diabetes Mellitus. Diabetes Carel998; 21:161B-7B.
- V Seshiah, V Balaji, Madhuri S Balaji, CB Sanjeevi, A Greent, Gestational Diabetes in India JAPI 2004; 52:707-11.
- Avi Benn -Haroush, Y Yogen and Moshe.Hod. Epidemiology of Gestational Diabetes and its association with type II Diabetes. Diabetes Med 2003; 21:103-113.
- Rampal K G, Osman A, Lubis SH, Khalid B A K.A Prevalence study of Diabetes Mellitus among rural Malays in Kuala, Selangor, Malaysia, Third World Congress .Diabetes in the tropics and developing countries, 1985.
- Kumar A, Takkar D, Sunesh K. Implications of diagnosis of glucose intolerance during pregnancy; Perinatal Mortality and Morbidity J Obst Gyn of India. 1993;43:759-63.
- Bhattacharya C, Awashti RT, Kumar S, Lamba PS. Routine screening for Gestational Diabetes Mellitus with glucose challenge test in antenatal patients. J Obst Gyn of India.2001; 51:75-78.
- Jindal A, Ahmed F, Bharadwaj B, Chaturvedi B.Prevalence, Clinical Profile and Outcome of Gestational Diabetes Mellitus. J Obst Gyn of India.2001; 51:46-49.
- Harris SB, et al: The Epidemiology of Diabetes in pregnant native Canadians: a risk profile. Diabetes Care 1997; 20(9):1422-1425.
- Henry OA, Beischer NA, Sheedy MT, Walstab JE. Gestational Diabetes and follow-up among immigrant Vietnam bom women. Aust N Z j Obstet Gynaecol; 1993, 33(2):109-114.
- Nauham GG,Haffaker BJ .Racial differences in oral glucose screening test results : establishing race specific criteria for abnormality in pregnancy[see comments] Obste and Gynae 1993;81(4):517-522.
- Rizvi JH et al Experience with screening for abnormal glucose tolerance in pregnancy: Maternal and Perinatal outcome. Asia Oceania J Obstet Gynaecol 1992, 18(2):99-105.
- Miselli V, Paglianiu, Bisi S et al. [epidemiology of gestational diabetes in scandiano health district 12 (usl 12)]. Minerva endocrinology 1994; 19(2):63-66.
- Serirat S, Deerochang C, Sunthomthepvarakul T, Jinayon P. Gestational Diabetes Mellitus.J Med Assoc Thailand 1992;75(6):315-319.
- Jang HC, et al: Screening for Gestational Diabetes Mellitus in Korea. Int J Gynaecol Obstet 1995;51(2):115-122.
- Oded Langer, MD, PhD Yariv Yogev, MD ,Orli Most, MD, Elly M.J. Xenakis, MD .Gestational diabetes: The consequences of not treating American Journal of Obstetrics and Gynecology April 2005, 192 (4) :989-997.

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- Michale Maresh. Diabetes In Pregnancy. Progress in Obstetrics and Gynaecology. John Studd; vol 13:191-207.
- Sutherland H.W, Stowen J M, MC Kenzie C. Simplifying the Clinical problem of Glycosuria in Pregnancy: Lancet 1970; I : 1069-1071.
- Lind T, Anderson J.Does Random blood glucose sampling outdate testing for glycosuria in the detection of Diabetes during Pregnancy. BMJ 1984; 289: 1569 -1571.
- Nasrat AA, Johnstne FD, Hasan S A M. Is random plasma glucose an efficient screening test for blood glucose tolerance in pegnancy. Pro. Journal of Obstetrics and Gynecology 1988; 95: 855 -860.
- Mathai. M; Thomas. T J: Kuruvila S; Jairaj P. Natl Med J India. July -Aug 1994; 7(9): 160-2.
- Coustan DR, Carpenter MW, Widness JA. The "breakfast tolerance test": screening for gestational diabetes with a standardized mixed nutrient meal. Am J Obstet Gynecol 1987; 157:1113-7.
- Huisman F H J And Dozy A M .Studies on the heterogenity of hemoglobin: Binding of hemoglobin with oxidized gluthatione. J of Lab Med 1962; 60:302-319.
- Bunn J F,Harrey D N AND Gabbary K H.Further identification of the nature and linkage of carbohydrate in hemoglobin Alc.Biochemical and Biophysical research communications, 1979:67-103.
- O'Shaughnessy Richard, Russ John, Zuspan F P. Am J Obstet Gynecol .1979;135:783.
- 35. Shah B D, cohen A W, May C, Gabbe SG, Am J. Obstet Gyn: 1982;144: 774.
- Baxi L., Barad D., Reece A.E Ferber R: Obst andGyn. Survery: 1984;64: 347.
- 37. Jovanovic L. Peterson M. Am J of Med 1981; 70: 331.
- Coen Ronald W, Parreco Richard, M Cousins Larry, Sandler J.A. Am J Obstet Gynecol .1980;136: 380.
- Sujata A Dalvi, Sangeeta Agarwal , Nishigandha Desai, Shirish S Seth. J Obstet Gynecol of India 1994; 44(2):181-185.
- Ross I S.Glycosylated Hemoglobin in the detection of Gestational Diabetes in an unselected population.in H W Sutherland, J M Stowers (Eds) Carbohydrate metabolism in pregnancy and new born. Churchill Livingstone, 1984:206-208.
- Jovanovic L, Peterson CM. Screening for gestational diabetes. Optimum timing and criteria for retesting. Diabetes 1985; 34(Suppl 2):21-3.
- Coustan DR, Widness JA, Carpenter MW, et al. Should the fifty-gram, one-hour plasma glucose screening test for gestational diabetes be administered in the fasting or fed state? Am J Obstet Gynecol 1986; 154:1031-5.
- Wilkerson H, O'Sullivan JB. A study of glucose tolerance and screening criteria in 752 unselected pregnancies. Diabetes 1963;12:313-8.
- Carpenter MW, Coustan DR. Criteria for screening tests of gestational diabetes. Am J Obstet Gynecol 1982; 144:768-73.
- Sacks DA, Abu-Fadil A, Greenspoon JS, Fotheringham N. Do the current standards for glucose tolerance testing in pregnancy represent a valid conversion of O'Sullivan's original criteria? Am J Obstet Gynecol 1989; 161:638-41.
- ACOG Practice Bulletin. Clinical management guidelines for obstetrician-gynecologists. Number 30, September 2001 (replaces Technical Bulletin Number 200, December 1994). Gestational diabetes. Obstet Gynec. 2001; 98:525-38.
- ADA Position Statement: gestational diab mell. Diabetes Care 2004; 27(Suppl I):S88-90.
- 48. Consultation WHO. Definition, diagnosis, and classification of diabetes mellitus and its complications: report of a WHO consultation Part I: diagnosis and classification of diabetes mellitus. Geneva: World Health Organization; 1999. American Diabetes Association. Gestational diabetes mellitus. Diabetes Care 2001; 24:S77.
- Posner NA, Silverstone FA, Brewer J, Heller M. Simplifying the intravenous glucose tolerance test. J Reprod Med 1982; 27:633-8.
- Michale Dg Gillmer, M Fathe Raslan .Management of Diabetes in pregnancy. Obstetrics for Postgraduates and Practioners by Sisir Chatopathyay, Bijoy Sree Sengupta James G Thornton, Parthasarthy Sengupta. 1999; 69-77.
- Weeks Jw,Major Ca ,De Veciana M ,Morgan Ma:Gestational Diabetes Mellitus:Does the presence of risk factors influence perinatal outcome ?

- diagnosis and outcome.Diabet Med 2000 Jan;17(l):26-32.
 53. Gajjar F,Maitra N KJntrapartum And Perinatal outcomes in women with gestational diabetes and mild gestational hyperglycemia. J Obstet Gynaecol india2005: 55(2):135-137.
- Vinita Das, Smita Kamara, Amita Mishra, Anjoo Agarwal, C G Agarwal. Screening for gestational diabetes and maternal and fetal outcome. J Obstet Gynaecol india 2004; 54(5):449-451.
- Ray R, Heng Bh,Lim C,Ling SI. Gestational Diabetes In Singaporean women: Use of glucose challenge test as a screening test and identification of high risk factors. Ann Acad Med Singapore 1996, 25(4):504-508.
- Naylor Cd,Sermer M,Chen E, Farine D: Selective Screening For Gestational Diabetes Mellitus:Toronto Trihospital Gestational Diabetes project investigators [see comments] .New England Journal Of Medicine. 1997; 337(22):1591-1596.
- Moses R,Colagiuri S: The extent of undiagnosed gestational diabetes mellitus in New South Wales . Med J of Aust. 1997; 167(I):14-16.
- Wilkins-Hang L, Horton Jn, Cruess Df, Frigoletto Fd: Antepartum screening in the office based practice: finding from the collaborative ambulatory research network. Obste and Gynecol .1996; 88(4 pt) 483-489.
- Donald R,Coustan M D .Making the diagnosis of Gestational Diabetes Mellitus. Clinical Obstetrics and Gynaecology 2000; 43(1):99-105.
- Jovanovic L, Metzger B, Knopp RH, et al. [beta]-hydroxybutyrate levels in type 1 diabetic pregnancy compared with normal pregnancy. Diabetes Care 1998; 21:1-5.
- Lind T, Harris VG. Changes in the oral glucose tolerance test during the puerperium. Br J Obstet Gynaecol 1971; 78:460-3.
- 62. Ferris a m, neubauer shb,bendel r b, green g w, ingardia c j, reece e a. perinatal lactation proctocol and outcome in mothers with and without insulin dependent diabetes mellitus .am j clin nutr 1993;57:43-48.
- Gabbe, Steven G. MD; Graves, Cornelia R. MD. Management of Diabetes Mellitus Complicating Pregnancy. The American College of Obstetricians and Gynecologists October 2003; 102(4): 857-868.
- Towner D, Kjos SL, Leung B, et al. Congenital malformations in pregnancies complicated by MDDM. Diabetes Care 1995; 18:1446-1451.
- F. Galerneau, S.E. Inzucchi / Obstet Gynecol Clin N Am (2004); 31:907-933
- Tuomilehto J, Lindstrom J, Eriksson JG, Valle TT, Hamalainen H, Ilanne-Parikka P, et al.Finnish Diabetes Prevention Study Group. Prevention of type 2 diabetes mellitus by changes inlifestyle among subjects with impaired glucose tolerance. N Engl J Med 2001; 344:1343 -50.
- O'Sullivan JB, Charles D, Mahan CM, Dandrow RV. Gestational diabetes and perinatal mortality rate. Am J Obstet Gynecol 1973; 116:901-4.
- Mesteger BE, Phelps, RL, Dooley SL. Mother in pregnancy complicated by Diabetes Mellitus in Porta D, Sharwin RS, Baron A Editors Ellenbergs and Rifkins Diabetes Mellitus. Newyork, McGraw Hill; 2002.P.626.
- V Seshiah, V Balaji. Pregnancy and Diabetes Mellitus. API Medicine Update 2002(12):357-372.