

**THE ROLE OF HbA1C AS AN EARLY PREDICTOR OF
GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY-
A PROSPECTIVE STUDY**

By

Dr. SOMAVARAPU DIVYA, MBBS



**DISSERTATION SUBMITTED TO SRI DEVARAJ URS ACADEMY OF HIGHER
EDUCATION AND RESEARCH, KOLAR, KARNATAKA IN PARTIAL FULFILLMENT
OF THE REQUIREMENTS FOR THE DEGREE OF**

**MASTER OF SURGERY
IN
OBSTETRICS AND GYNAECOLOGY**

**UNDER THE GUIDANCE OF
DR. SHEELA S R
PROFESSOR
DEPARTMENT OF OBSTETRICS AND GYNAECOLOGY**

**CO-GUIDE
DR. SHASHIDHAR K N
PROFESSOR
DEPARTMENT OF BIOCHEMISTRY**



**SRI DEVARAJ URS MEDICAL COLLEGE
TAMAKA, KOLAR 563101**

2024

SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION AND RESEARCH

DECLARATION BY THE CANDIDATE

I hereby declare that this dissertation entitled “**THE ROLE OF HbA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY**” is a bonafide and genuine research work carried out by me under the direct guidance of **DR.SHEELA S R** Professor, Department of obstetrics and gynaecology, and co-guidance of **DR. SHASHIDHAR K N**, Professor department of biochemistry Sri Devaraj Urs Medical College, Tamaka, Kolar, in partial fulfillment of the requirement for the degree of **MASTER OF SURGERY in OBSTETRICS AND GYNAECOLOGY**

Place:

Kolar

**DR. SOMAVARAPU DIVYA
POSTGRADUATE IN OBSTETRICS
AND GYNAECOLOGY
SRI DEVRAJ MEDICAL COLLEGE,
TAMAKA , KOLAR**

**SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION
AND RESEARCH**

CERTIFICATE BY THE GUIDE

This is to certify that the dissertation entitled “**THE ROLE OF HbA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY**” is a bonafide and genuine research work done by **DR.SOMAVARAPU DIVYA** in partial fulfillment of the requirement for the degree of **MASTER OF SURGERY in OBSTETRICS AND GYNAECOLOGY** as per regulations of **SRI DEVRAJ URS ACADEMY OF HIGHER EDUCATION AND RESEARCH, KOLAR.**

Date:

Place: Kolar

Signature of Guide:

DR. SHEELA S R

PROFESSOR

**DEPARTMENT OF OBSTETRICS AND
GYNAECOLOGY**

**SRI DEVARAJ URS MEDICAL
COLLEGE TAMAKA, 563101,
KOLAR**

**SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION
AND RESEARCH**

CERTIFICATE BY THE CO-GUIDE

This is to certify that the dissertation entitled “**THE ROLE OF HbA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY**” is a bonafide and genuine research work done by **DR.SOMAVARAPU DIVYA** in partial fulfillment of the requirement for the degree of **MASTER OF SURGERY in OBSTETRICS AND GYNAECOLOGY** as per regulations of **SRI DEVRAJ URS ACADEMY OF HIGHER EDUCATION AND RESEARCH, KOLAR.**

Date:

Place: Kolar

Signature of CO-Guide

DR. SHASHIDHAR K N

**PROFESSOR
DEPARTMENT OF BIOCHEMISTRY
SRI DEVARAJ URS MEDICAL
COLLEGE TAMAKA,
KOLAR, 563101**

**SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION
AND RESEARCH**

**ENDORSEMENT BY THE HOD, PRINCIPAL/HEAD OF
THE INSTITUTION**

This is to certify that the dissertation entitled “**THE ROLE OF HbA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY**” is a bonafide and genuine research work done by **DR. SOMAVARAPU DIVYA** under the guidance and co-guidance of **DR. SHEELA S R and DR. SHASHIDHAR K N**, Professor of Department of Obstetrics and Gynecology And Professor of Department of biochemistry respectively, Sri Devaraj Urs Medical College, Tamaka, Kolar.

Signature of HOD

DR MUNIKRISHNA M

PROFESSOR AND HOD

DEPARTMENT OF OBSTETRICS

AND GYNECOLOGY

SRI DEVARAJ URS MEDICAL COLLEGE

TAMAKA KOLAR. 563101.

DATE:

PLACE: KOLAR

Signature of Principal

DR. PRABHAKAR K

PRINCIPAL,

SRI DEVARAJ URS MEDICAL

COLLEGE TAMAKA,

KOLAR. 563101



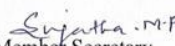

DATE:

PLACE: KOLAR

,

SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION AND RESEARCH, TAMAKA, KOLAR, KARNATAKA

ETHICAL COMMITTEE CERTIFICATE

	<p>SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION & RESEARCH</p> <p>SRI DEVARAJ URS MEDICAL COLLEGE Tamaka, Kolar</p> <p>INSTITUTIONAL ETHICS COMMITTEE</p>	
<p>Members</p> <ol style="list-style-type: none">1. Dr. D.E.Gangadhar Rao, (Chairman) Prof. & HOD of Zoology, Govt. Women's College, Kolar2. Dr. Sujatha.M.P., (Member Secretary), Prof. Dept. of Anesthesia, SDUMC3. Mr. Gopinath Paper Reporter, Samyukth Karnataka4. Mr. G. K. Varada Reddy Advocate, Kolar5. Dr. Hariprasad S, Assoc. Prof Dept. of Orthopedics, SDUMC6. Dr. Abhinandana R Asst. Prof. Dept. of Forensic Medicine, SDUMC7. Dr. Ruth Sneha Chandrakumar Asst. Prof. Dept. of Psychiatry, SDUMC8. Dr. Usha G Shenoy Asst. Prof., Dept. of Allied Health & Basic Sciences SDUAHER9. Dr. Munilakshmi U Asst. Prof. Dept. of Biochemistry, SDUMC10. Dr. D. Srinivasan, Assoc. Prof. Dept. of Surgery, SDUMC11. Dr. Waseem Anjum, Asst. Prof. Dept. of Community Medicine, SDUMC12. Dr. Shilpa M D Asst. Prof. Dept. of Pathology, SDUMC		
<p>No. SDUMC/KLR/IEC/300/2022-23 Date: 20-07-2022</p> <p>PRIOR PERMISSION TO START OF STUDY</p> <p>The Institutional Ethics Committee of Sri Devaraj Urs Medical College, Tamaka, Kolar has examined and unanimously approved the synopsis entitled "The role of HBA1C as an early predictor of gestational diabetes mellitus in early pregnancy - A prospective study" being investigated by Dr. Somavarapu Divya, Dr. Sheela S.R. & Dr. K.N. Shashidhar¹ in the Departments of OBG & Biochemistry¹ at Sri Devaraj Urs Medical College, Tamaka, Kolar. Permission is granted by the Ethics Committee to start the study.</p> <div style="display: flex; justify-content: space-around;"><div><p> Member Secretary Member Secretary Institutional Ethics Committee Sri Devaraj Urs Medical College Tamaka, Kolar.</p></div><div><p> Chairman CHAIRMAN Institutional Ethics Committee Sri Devaraj Urs Medical College Tamaka, Kolar</p></div></div>		

**SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION
AND RESEARCH, TAMAKA, KOLAR, KARNATAKA**

COPY RIGHT

DECLARATION BY THE CANDIDATE

I hereby declare that the Sri Devaraj Urs Academy of Higher Education and Research, Tamaka, Kolar, Karnataka shall have the rights to preserve, use and disseminate this dissertation/thesis in print or electronic format for academic /research purpose.

Date:
Place: Kolar

Signature of the candidate
DR. SOMAVARAPU DIVYA
Postgraduate
Department of Radiodiagnosis
Sri Devaraj Urs Medical
College Tamaka, Kolar

**© Sri Devaraj Urs Academy of Higher Education and Research,
Tamaka, Kolar, Karnataka.**

PLAGIARISM CERTIFICATE



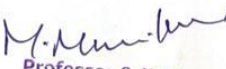
SRI DEVARAJ URS ACADEMY OF HIGHER EDUCATION & RESEARCH
Tamaka, Kolar 563103


Certificate of Plagiarism Check

Title of the Thesis/Dissertation	THE ROLE OF HBA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY
Name of the Student	DR SOMAVARAPU DIVYA
Registration Number	21OG1090
Name of the Supervisor / Guide	DR. SHEELA S.R.
Department	OBSTETRICS AND GYNECOLOGY
Acceptable Maximum Limit (%) of Similarity (PG Dissertation)	10%
Similarity	6%
Software used	Turnitin
Paper ID	2414161766
Submission Date	09/07/2024


Signature of Student


DR. S.R. SHEELA
Professor of OBG
Signature of Guide/Supervisor
Date..... Time.....


Professor & HoD
Obstetrics and Gynaecology
Sri Devaraj Urs Medical College
Tamaka, Kolar.


University Librarian
Senior Librarian
ULLRC, SUAHER
Tamaka, KOLAR-563103


PG Co-ordinator
PG Coordinator
Sri Devaraj Urs Medical College
Tamaka, Kolar-563103

Turnitin Originality Report

Processed on: 09-Jul-2024 14:30:15T

ID: 2414161766

Word Count: 26390

Submitted: 2

THE ROLE OF HbA1c AS AN EARLY PREDICTOR
OF GE... By Dr. Somavarapu Divya

Similarity Index

6%

Similarity by Source

Internet Sources	5%
Publications	5%
Student Papers	1%

include quoted include bibliography excluding matches < 10 words mode: quickview (classic) report print

refresh download

- 1% match (Internet from 31-Oct-2021)
<https://filedownload.com/wp-content/uploads/2020/10/Dewhurst5-Textbook-Of-Obstetrics-.pdf>
- <1% match (Internet from 15-Mar-2024)
<https://listens.online/thesis/thesis-on-gestational-diabetes>
- <1% match (M., Umadevi. "Outcome of Diabetes Mellitus in Pregnancy in a Tertiary Referral Centre", Rajiv Gandhi University of Health Sciences (India), 2023)
M., Umadevi. "Outcome of Diabetes Mellitus in Pregnancy in a Tertiary Referral Centre", Rajiv Gandhi University of Health Sciences (India), 2023
- <1% match ("Abstracts of the IDF Congress in Paris 2003", Diabetologia, 2003)
"Abstracts of the IDF Congress in Paris 2003", Diabetologia, 2003
- <1% match (Internet from 07-Dec-2020)
<https://www.science.gov/topicpages/a/a1c+fasting+plasma>
- <1% match (Internet from 04-Mar-2023)
<https://www.science.gov/topicpages/d/diabetes+mellitus+dm1>
- <1% match ()
H N Rohini, Pushpanathan Punita, Prasanna Kumar Santhekadur, MV Ravishankar, "Gestational Diabetes Mellitus - The Modern Indian Perspective", Indian Journal of Endocrinology and Metabolism
- <1% match (Internet from 18-Oct-2022)
<http://repository-tnmgrmu.ac.in>
- <1% match ()
Shilpa Reddy, T. "A comparative study of platelet profile in gestational diabetes mellitus versus healthy pregnancies: A cross sectional study", 2018
- <1% match (Internet from 20-Nov-2023)
<http://repository-tnmgrmu.ac.in>
- <1% match (Internet from 27-Mar-2019)
<https://link.springer.com/content/pdf/10.1007%2F978-1-60327-250-6.pdf>
- <1% match (Internet from 17-Nov-2017)
<https://link.springer.com/content/pdf/10.1007%2F978-3-319-56440-1.pdf>
- <1% match ()
- <1% match ()
- <1% match ()
Martis, Ruth. "Optimal treatments and experiences for women with gestational diabetes mellitus (GDM): improving health for mothers and babies", 2018
- <1% match (Internet from 09-Nov-2017)
<http://www.nejm.org>
- <1% match (Internet from 12-Jun-2018)
<http://gynecology.sbmua.ac.ir>
- <1% match (Internet from 28-Aug-2019)
<http://gynecology.sbmua.ac.ir>
- <1% match ("International Textbook of Diabetes Mellitus", Wiley, 2015)
"International Textbook of Diabetes Mellitus", Wiley, 2015
- <1% match (Internet from 03-Dec-2023)
<https://vdoc.pub/documents/survival-analysis-a-self-learning-text-3t5onb2e5j3g>
- <1% match (Internet from 24-Mar-2022)
https://www.researchgate.net/publication/8944841_Gestational_diabetes_Problems_associated_with_the_oral_glucose_tolerance_test
- <1% match (Internet from 22-Feb-2023)
https://www.researchgate.net/publication/300372549_Management_of_type_2_diabetes_in_adults_Summary_of_updated_NICE_guides
- <1% match (Internet from 07-Aug-2022)

PROFESSOR
TAMAKA KOLAR

ULCER, STOMACH
TAMAKA KOLAR-563103

https://mdpi-res.com/bookfiles/book/3969/Recent_Advances_in_Gestational_Diabetes_Mellitus.pdf?v=1659459097

<1% match (Helena F Fadl, David Simmons. "Trends in diabetes in pregnancy in Sweden 1998-2012". BMJ Open Diabetes Research & Care, 2016)
[Helena F Fadl, David Simmons. "Trends in diabetes in pregnancy in Sweden 1998-2012". BMJ Open Diabetes Research & Care, 2016](#)

<1% match ("Abstracts of the EASD, Stockholm 2010". Diabetologia, 2010)
["Abstracts of the EASD, Stockholm 2010". Diabetologia, 2010](#)

<1% match (Internet from 21-Feb-2022)
<http://enjournalssu.ac.ir>

<1% match (Internet from 20-May-2019)
<https://www.drkirstencannan.com/blog?category=Women%27s+health>

<1% match ("42nd EASD Annual Meeting of the European Association for the Study of Diabetes", Diabetologia, 2006)
["42nd EASD Annual Meeting of the European Association for the Study of Diabetes". Diabetologia, 2006](#)

<1% match ("Comprehensive Clinical Approach to Diabetes During Pregnancy", Springer Science and Business Media LLC, 2022)
["Comprehensive Clinical Approach to Diabetes During Pregnancy". Springer Science and Business Media LLC, 2022](#)

<1% match (Fang Zhou, Xiao Ran, Fangliang Song, Qinglan Wu, Yuan Jia, Ying Liang, Suichen Chen, Guojun Zhang, Jie Dong, Yukun Wang. "A stepwise prediction and interpretation of gestational diabetes mellitus: Foster the practical application of machine learning in clinical decision". Heliyon, 2024)
[Fang Zhou, Xiao Ran, Fangliang Song, Qinglan Wu, Yuan Jia, Ying Liang, Suichen Chen, Guojun Zhang, Jie Dong, Yukun Wang. "A stepwise prediction and interpretation of gestational diabetes mellitus: Foster the practical application of machine learning in clinical decision". Heliyon, 2024](#)

<1% match (W Y. Fok. "The Influence of Fetal Position on Amniotic Fluid Index and Single Deepest Pocket", Obstetrical & Gynecological Survey, 01/2007)
[W Y. Fok. "The Influence of Fetal Position on Amniotic Fluid Index and Single Deepest Pocket". Obstetrical & Gynecological Survey, 01/2007](#)

<1% match (Internet from 14-Oct-2022)
<https://www.hindawi.com/journals/jdr/2021/5537110/>

<1% match ("A Practical Manual of Diabetes in Pregnancy", Wiley, 2010)
["A Practical Manual of Diabetes in Pregnancy". Wiley, 2010](#)

<1% match ("Abstracts 2007", Diabetologia, 2007)
["Abstracts 2007". Diabetologia, 2007](#)

<1% match (Jianbin Sun, Sanbao Chai, Xin Zhao, Ning Yuan, Jing Du, Yufang Liu, Zhi Li, Xiaomei Zhang. "Predictive Value of First-Trimester Glycosylated Hemoglobin Levels in Gestational Diabetes Mellitus: A Chinese Population Cohort Study". Journal of Diabetes Research, 2021)
[Jianbin Sun, Sanbao Chai, Xin Zhao, Ning Yuan, Jing Du, Yufang Liu, Zhi Li, Xiaomei Zhang. "Predictive Value of First-Trimester Glycosylated Hemoglobin Levels in Gestational Diabetes Mellitus: A Chinese Population Cohort Study". Journal of Diabetes Research, 2021](#)

<1% match (Jianbin Sun, Sanbao Chai, Xin Zhao, Ning Yuan, Jing Du, Yufang Liu, Zhi Li, Xiaomei Zhang. "Predictive Value of First-Trimester Glycosylated Hemoglobin Levels in Gestational Diabetes Mellitus: A Chinese Population Cohort Study". Journal of Diabetes Research, 2021)
[Jianbin Sun, Sanbao Chai, Xin Zhao, Ning Yuan, Jing Du, Yufang Liu, Zhi Li, Xiaomei Zhang. "Predictive Value of First-Trimester Glycosylated Hemoglobin Levels in Gestational Diabetes Mellitus: A Chinese Population Cohort Study". Journal of Diabetes Research, 2021](#)

<1% match (Nwoli, Chidiebere H., "Improving preconception care for women with diabetes: development and feasibility study of a mobile application", 2018)

<1% match ("Nutrition and Diet in Maternal Diabetes", Springer Science and Business Media LLC, 2018)
["Nutrition and Diet in Maternal Diabetes". Springer Science and Business Media LLC, 2018](#)

<1% match (Poornima, C., "The Glucose Challenge Test for Screening Gestational Diabetes in Pregnant Women With No Risk Factors.", Dr. NTR University of Health Sciences (India), 2019)
[Poornima, C., "The Glucose Challenge Test for Screening Gestational Diabetes in Pregnant Women With No Risk Factors.", Dr. NTR University of Health Sciences \(India\), 2019](#)

<1% match (Internet from 26-Jun-2021)
<https://dokumen.pub/varneys-midwifery-6th-ed-9781284127966-1284127966.html>

<1% match (Internet from 27-Jun-2023)
<https://dliem.wms.ac.ir/bitstream/Hannan/996/1/4.pdf>

<1% match (Internet from 25-Dec-2022)
https://pure.rug.nl/ws/files/49804538/Complete_Thesis.pdf

<1% match (Internet from 20-Mar-2023)
<https://www.cfp.ca/content/cfp/49/6/761.full.pdf>

<1% match (A., Amarnath. "Waist-Calf Ratio as a Marker of Carotid Atherosclerosis in Patients of Type 2 Diabetes Mellitus", Rajiv Gandhi University of Health Sciences (India), 2023)
[A., Amarnath. "Waist-Calf Ratio as a Marker of Carotid Atherosclerosis in Patients of Type 2 Diabetes Mellitus". Rajiv Gandhi University of Health Sciences \(India\), 2023](#)

<1% match (Meharban, A., "Evaluation of Hydro-Ethanol Extract of Syzygium Cumini Seeds in High Fat Diet and Streptozotocin Induced Type 2 Diabetes Mellitus in Wistar Albino Rats", Rajiv Gandhi University of Health Sciences (India), 2023)
[Meharban, A., "Evaluation of Hydro-Ethanol Extract of Syzygium Cumini Seeds in High Fat Diet and Streptozotocin Induced Type 2 Diabetes Mellitus in Wistar Albino Rats". Rajiv Gandhi University of Health Sciences \(India\), 2023](#)

<1% match (Internet from 01-Jul-2023)
https://ada.silverchair-rdn.com/ada/content_public/journal/care/issue/46/supplement_1/21/standards-of-care-2023.pdf?Expires=1691198812&Signature=F1xw3-5gQ5IK-B-5s9GQ12s-rt-

6ZqRdsL9Jf, %7GyrdLHMPJd0NymQds.c. WcydmX5yUyYhDQhBamC6Ng

<1% match (Internet from 09-Dec-2022)

<https://diabetes.newlifepoutlook.com/what-is-type-2-diabetes/?all=1>

<1% match (Internet from 11-Apr-2020)

<https://rc-scnd.com/document/158306264/Canadian-Journal-of-Diabetes>

<1% match (Internet from 21-Jan-2023)

<https://www.yumpu.com/en/document/view/63047570/ispad-clinical-practice-consensus-guidelines-2018>

<1% match ("2. Classification and Diagnosis of Diabetes: ", Diabetes Care, 2018)

"2. Classification and Diagnosis of Diabetes: ", Diabetes Care, 2018

<1% match (D., Shwtha. "An Observational Study on the Effect of Eladi Gutika in the Management of Chardi (Vomiting) in Children", Rajiv Gandhi University of Health Sciences (India), 2023)

D., Shwtha. "An Observational Study on the Effect of Eladi Gutika in the Management of Chardi (Vomiting) in Children", Rajiv Gandhi University of Health Sciences (India), 2023

<1% match (Mohammed Bashir, Yassin Fagier, Badredeen Ahmed, Justin C Konje. "An overview of diabetes mellitus in pregnant women with obesity", Best Practice & Research Clinical Obstetrics & Gynaecology, 2024)

Mohammed Bashir, Yassin Fagier, Badredeen Ahmed, Justin C Konje. "An overview of diabetes mellitus in pregnant women with obesity", Best Practice & Research Clinical Obstetrics & Gynaecology, 2024

<1% match (Yi Lai, Hanxiao Chen, Ze Du, Shu Zhou, Wenming Xu, Tao Li. "The diagnostic accuracy of HbA1c in detecting gestational diabetes mellitus among Chinese pregnant individuals", Annals of Translational Medicine, 2020)

Yi Lai, Hanxiao Chen, Ze Du, Shu Zhou, Wenming Xu, Tao Li. "The diagnostic accuracy of HbA1c in detecting gestational diabetes mellitus among Chinese pregnant individuals", Annals of Translational Medicine, 2020

<1% match ("The Diabetes Textbook", Springer Science and Business Media LLC, 2019)

"The Diabetes Textbook", Springer Science and Business Media LLC, 2019

<1% match (student papers from 21-Oct-2019)

Submitted to Florida International University on 2019-10-21

<1% match (M., Ashwini. "Design and Evaluation of Nano-Lipid Carrier (NLC) Loaded Transdermal Patches of Selected Oral Antidiabetic Drug for the Treatment of Gestational Diabetes", Rajiv Gandhi University of Health Sciences (India), 2023)

M., Ashwini. "Design and Evaluation of Nano-Lipid Carrier (NLC) Loaded Transdermal Patches of Selected Oral Antidiabetic Drug for the Treatment of Gestational Diabetes", Rajiv Gandhi University of Health Sciences (India), 2023

<1% match ("Dewhurst's Textbook of Obstetrics & Gynaecology", Wiley, 2018)

"Dewhurst's Textbook of Obstetrics & Gynaecology", Wiley, 2018

<1% match (Dornhorst, Anne, and Catherine Williamson. "Diabetes and Endocrine Disease in Pregnancy", Dewhurst's Textbook of Obstetrics & Gynaecology Edmonds/Dewhurst's Textbook of Obstetrics & Gynaecology, 2012.)

Dornhorst, Anne, and Catherine Williamson. "Diabetes and Endocrine Disease in Pregnancy", Dewhurst's Textbook of Obstetrics & Gynaecology Edmonds/Dewhurst's Textbook of Obstetrics & Gynaecology, 2012.

<1% match (S., Prajwal. "A Comparative Study of Using Glibenclamide Versus Insulin in the Treatment of Gestational Diabetes Mellitus and its Outcome", Rajiv Gandhi University of Health Sciences (India), 2023)

S., Prajwal. "A Comparative Study of Using Glibenclamide Versus Insulin in the Treatment of Gestational Diabetes Mellitus and its Outcome", Rajiv Gandhi University of Health Sciences (India), 2023

<1% match (student papers from 08-Jan-2020)

Submitted to University of Central Lancashire on 2020-01-08

<1% match (Internet from 17-Jul-2023)

<https://downloads.hindawi.com/journals/ecam/2023/3020033.pdf>

<1% match (Internet from 19-May-2019)

<https://onlinelibrary.wiley.com/doi/pdf/10.1002/9781444324808.ch53>

<1% match (Internet from 21-Nov-2022)

<https://resurge-product.shop/does-tricare-cover-diabetic-supplies.html>

<1% match (Internet from 21-May-2014)

<http://www.biomedcentral.com>

<1% match (Internet from 24-Oct-2022)

<https://www.endotext.org/chapter/page/12/>

<1% match (Meena Khandelwal. "GDM: Postpartum management to reduce long-term risks", Current Diabetes Reports, 08/2008)

Meena Khandelwal. "GDM: Postpartum management to reduce long-term risks", Current Diabetes Reports, 08/2008

<1% match (Ribal Kattini, Ruben Hummelen, Len Kelly. "Early Gestational Diabetes Mellitus Screening With Glycated Hemoglobin: A Systematic Review", Journal of Obstetrics and Gynaecology Canada, 2020)

Ribal Kattini, Ruben Hummelen, Len Kelly. "Early Gestational Diabetes Mellitus Screening With Glycated Hemoglobin: A Systematic Review", Journal of Obstetrics and Gynaecology Canada, 2020

<1% match (Internet from 16-Nov-2023)

<https://academic.oup.com/jcem/article/107/3/e1117/6409789>

<1% match (Internet from 01-Oct-2022)

https://diabetesjournals.org/care/article/30/Supplement_2/S251/23932/Summary-and-Recommendations-of-the-Fifth

<1% match ()

PROFESSOR

R. S. K. Hospital & Research Centre

TAMAKA KOLAR

S. S. K. Hospital

ULLR, SQUATER

Tamaka, KOLAR-563102

ACKNOWLEDGEMENT

Firstly, I would like to express my gratitude to my Guide, **Dr. SHEELA S R**, Professor of Obstetrics and Gynaecology, SDUMC, Kolar, and my co-guide **DR. SHASHIDHAR K N**, Professor, Department of Biochemistry for their patience, unwavering support, guidance and contribution. I'd also like to thank her for her constant encouragement and guidance in all aspects of my professional life.

I am sincerely thankful to **Dr. MUNIKRISHNA M**, Professor and Head of Department of Obstetrics and Gynaecology, SDUMC for encouraging me and providing her kind support and valuable suggestions throughout the entire process.

I wholeheartedly thank **Dr. RATHNAMMA P**, Professors in the department of Obstetrics and Gynaecology for their valuable teaching and insights on perseverance and professional ethics, and their moral support and encouragement.

would like to express my heartfelt gratitude to my beloved parents, **DR. SOMAVARAPU KONDALA RAO** and **DR. PRAMEELA DEVI G**, my brother, **DR SAI PRANEETH**, and my husband **DR PHANITEJA P** for always inspiring me and providing me with unwavering support, encouragement, and unconditional love and constantly motivating me throughout the course.

I also want to take this opportunity to sincerely thank my professors **Dr. VIMARSHITHA**, **Dr. AASHRITHA**, **Dr. NANDINI**, **Dr. KAVYA**, **DR SUKHINI**, **DR YAMINI** for their constant support and encouragement, and appreciate their relentless pursuit to teach us.

I thank my colleagues and friends **DR MADHURYA**, **DR AJITHA**, **DR ASHWINI**, **DR LAKSHMI**, **DR MEGHANA**, **DR SAMYUKTANJALI**, **DR RADHIKA** AND **DR SHREYA** for their unflinching support for the past three years.

Last but not the least, I extend my gratitude towards all the patients who agreed to participate in this study without whose precious support, it would not have been possible to conduct this study.

DR. SOMAVARAPU DIVYA

TABLE OF CONTENTS

SLNO.	TITLE	PAGE NO.
1	INTRODUCTION	1
2	AIM AND OBJECTIVES	28
3	REVIEW OF LITERATURE	29
4	MATERIAL AND METHODS	82
5	OBSERVATION AND RESULTS	86
6	DISCUSSION	103
7	SUMMARY	125
8	LIMITATION & RECOMMENDATION	126
9	CONCLUSION	127
10	REFERENCES	128
	ANNEXURE	140
I	CONSENT FORM	
II	PROFORMA	
III	KEY TO MASTER CHART	
IV	MASTER CHART	

LIST OF TABLES

SL .NO	TITLE	PAGE .NO
1	Age distribution of samples tested during 1 ST Trimester	86
2	Distribution of study population with reference to HbA1c values	87
3	Age distribution of abnormal HbA1c	88
4	Age wise distribution of sample with reference to glycemic control	89
5	Incidence of GDM in different age groups	91
6	Parity Distribution Total sample tested-75 1 ST Trimester	93
7	Parity distribution with reference glycemic control	94
8	Incidence of GDM with reference to parity	95
9	Abnormal HbA1c correlates with excessive weight gain Abnormal HbA1c-18	96
10	Glycemic control basing on Booked/Unbooked status	97
11	Incidence of GDM basing on Booked/Unbooked status	98
12	Incidence of GDM basing on social statu	99
13	Glycemic control basing on social status	100
14	HbA1c distribution with oral glucose tolerance test 24-28 weeks	101

LIST OF FIGURES

<u>SL.NO</u>	TITLE	PAGE .NO
1	Risk factors for antagonistic pregnancy result	1
2	CEMACH Report of PNMR in GDM	6
3	Fertilisation, implantation, and the whole pregnancy are all impacted by maternal hyperglycaemia	7
4	Developmental determinants of Metabolic health in GDM and its complications	30
5	HbA1c levels in GDM-FLOW CHART	58
6	PERICONCEPTIONAL HbA1c	60
7	NO OF SD OF HbA1c	60

LIST OF GRAPHS

<u>SL.NO</u>	<u>TITLE</u>	<u>PAGE.NO</u>
1	Age distribution of samples tested	86
2	Distribution of study population with reference to HbA1c values	87
3	Age distribution of abnormal HbA1c	88
4	Age wise distribution of sample with reference to glycemic control	90
5	Incidence of GDM in different age groups.	92
6	Parity Distribution Total sample tested-75 1 ST Trimester	93
7	Parity distribution with reference glycemic control	94
8	Incidence of GDM with reference to parity	95
9	Abnormal HbA1c correlates with excessive weight gain Abnormal HbA1c-	96
10.	Glycemic control basing on Booked/Unbooked status	97
11.	Incidence of GDM basing on Booked/Unbooked status	98
12	Incidence of GDM basing on social status	99
13.	Glycemic control basing on social status	100
14.	HbA1c distribution with oral glucose tolerance test 24-28 weeks	101
15.	ROC Curve for HbA1c in first trimester in predicting GDM	102

LIST OF ABBREVIATIONS

DM	Diabetes Mellitus
GDM	Gestational Diabetes Mellitus
HbA1c	Glycosylated haemoglobin
PE	Preeclampsia
FBS	Fasting blood glucose
PPBS	Post prandial blood glucose
OGTT	Oral glucose tolerance test
FPG	Fasting plasma glucose
ADA	American Diabetic Association
IADPSG	International Association of Diabetes and Pregnant Study Group
WHO	World health organisation
GRBS	Random blood glucose
GCT	Glucose challenge Test
IR	Insulin resistance
NDDP	National Diabetic Data Group
SD	Standard Deviation
PC	Platelet count
DIPSI	Diabetes in Pregnancy Study Group
NDDA	National diabetes data group

ACOG	American College of obstetrician and Gynaecology
RBC	Red blood cells
HAPO	Hyperglycaemia and Restricting Pregnancy Result (HAPO) research.
ICD	International classification of diseases
IGT	Impaired Glucose Tolerance
BMI	Body mass index
CEMACH	Confidential enquiry in to maternal and child health
NICE	National institute for health and care technology
AC	Abdominal Circumference
VRIII	Intravenous insulin injection
UTI	Urinary tract infection
CSII	consistent subcutaneous insulin mixture siphon
MDI	multi-portion insulin,
CGM	constant persistent glucose checking
CAD	coronary artery disease
HKDC1	hexokinase domain containing 1
FDA	Food and Drug administration
DCCT	Diabetes Control and Intricacies Preliminary
FIGO	International federation of Gynecology and obstetrics
ROC	Receiver operating characteristic curve

ABSTRACT:

BACKGROUND: Gestational Diabetes Mellitus (GDM) that starts or is first identified during pregnancy is known as gestational diabetes mellitus (GDM). Universally, gestational diabetes influences 90% of pregnant ladies. One out of ten pregnancies is connected to diabetes. Fetal and maternal results might be extreme in the event that gestational diabetes is either misdiagnosed or not oversaw as expected. A few examinations, nonetheless, have shown that the two moms and offspring of GDM ladies are at increased risk for creating type 2 diabetes later on.

Rates of GDM are a lot higher in India, quite possibly of the most populated country on the planet, with gauges going from 10.3 to 14.3 percent.

AIM AND OBJECTIVES:

To evaluate the HbA1c in all to begin with first trimester pregnant women.

To assess the connection of HbA1c in early pregnancy (8-12 weeks) and expectation of development of diabetes mellitus in second trimester (24-28 weeks) of pregnancy with oral glucose tolerance test (OGTT).

METHODS:

Seventy-five pregnant ladies going to the OB/GYN facility at Sri Devaraj Clinical School in Kolar, D.t., Karnataka's RL Jalappa medical clinic were the subjects of our forthcoming observational exploration. To gauge HbA1C and fasting blood glucose levels in the primary trimester, analysts submitted blood tests from all pregnant ladies who partook in the review. Between weeks 24 and 28, they had a 75 g OGTT, which is a normalized, one-step test.

RESULTS:

Majority of study members are old enough gathering <25yrs comprising 40(53.33%). Among every one of the members 57 (76%) were viewed as with typical HbA1c. Out of 18 ladies with unusual HbA1c values larger part 10(14.28%) have a place with 25-35 age bunch with P esteem viewed as 0.005 which is measurably huge. Comparatively with the movement old enough the frequency of GDM is likewise increasing with X2 esteem being 18.66 and P esteem <0.001 with high importance. Larger part of the ones who were signed up for the review were Primigravida 46 (61.33%). With increase in equality poor glycaemic control is seen with X2 esteem being 9.68 and P esteem 0.002 which is huge. Frequency of GDM is increased with equality with P esteem <0.001 with high importance. It was observed that there is no relationship between's financial status and event of GDM. ROC Bend for HbA1c in first trimester in anticipating GDM viewed as exceptionally critical with P worth of <0.0001. Huge cut off HbA1C esteem in foreseeing diabetes mellitus by OGTT was viewed as 5.7 (with Awareness is 100 percent and Misleading energy rate is 0%)

CONCLUSION:

Estimation of HbA1c in first trimester is a good predictor for occurrence of GDM in second trimester. Life style modification and Medical nutrition therapy can be advised to those women who are with poor glycemic control in first trimester there by preventing the complications of GDM later in pregnancy, intrapartum and postpartum. This may also prevent occurrence of DM in progeny.

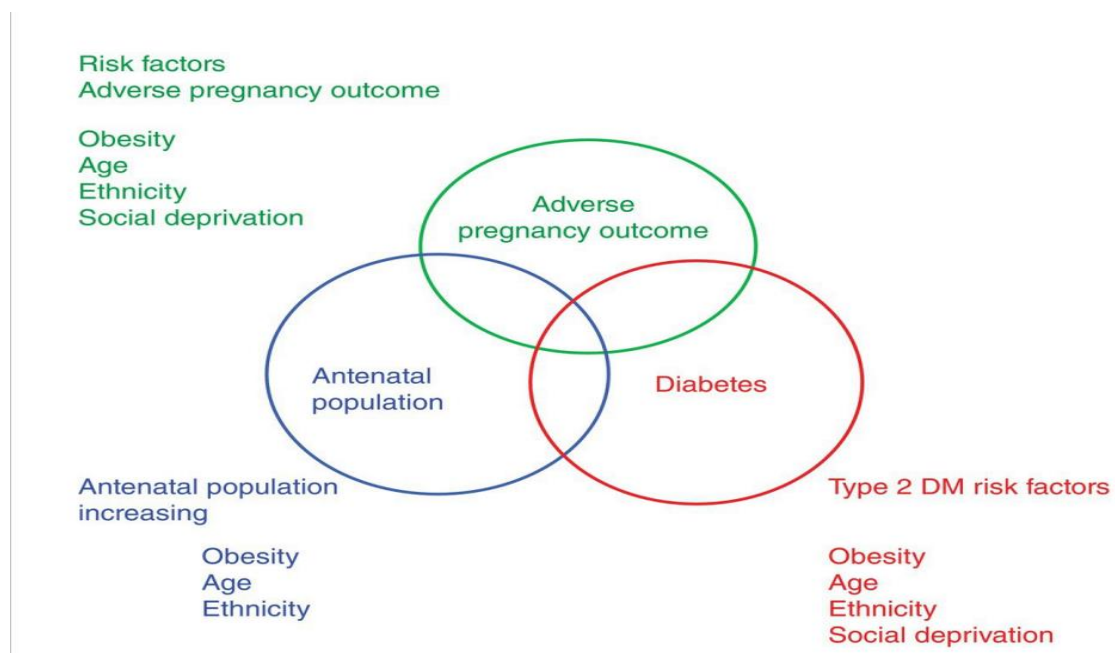
Keywords: FBS,GDM, OGTT

INTRODUCTION

INTRODUCTION

Significant dangers to the mother's and the developing foetus' health, as well as effects on the child's future health, are associated with diabetes throughout pregnancy^{1,2,3}. There is a higher risk of diabetes during pregnancy from pre-gestational diabetes, whether it's type 1 or type 2, compared to GDM, which develops during the pregnancy. This risk grows as the maternal hyperglycaemia level rises.³

Risk factors for antagonistic pregnancy result



From Dewhurst's text book of Obstetrics and Gynaecology¹¹

Figure 1 outlines the connection between prenatal populace socioeconomic and the elements associated with the gamble of diabetes and antagonistic pregnancy results.¹

Ladies who have type 1 diabetes, type 2 diabetes, or gestational diabetes have long haul issues because of the way that they are pregnant with a youngster who likewise has diabetes. Around half of infants conveyed to mothers with pre-gestational diabetes display indications of sped up advancement, with generally 40% of these infants being brought into the world at or over the 90th percentile for their gestational age. Moreover, kids who have a high birthweight and neonatal adiposity are at an increased gamble of developing early-beginning corpulence, metabolic brokenness, and diabetes further down the road, particularly when combined with shoulder dystocia and cesarean segment rates above 60%.³

The St. Vincent Announcement, laid out in 1989, swore to increase the incidence of effective pregnancies among ladies with diabetes contrasted with those without the condition^{5, 6}. Notwithstanding substantial advancement in the fields of obstetrics, neonatal, and diabetes care, this has yet to transpire. Notwithstanding certain advancements, the incidence of perinatal fatalities, stillbirths, and innate anomalies remains two to multiple times higher for ladies with pre-gestational diabetes contrasted with those with type 1 diabetes. Lately, there has been an outstanding flood in the pervasiveness of type 2 diabetes mellitus among ladies of conceptive age who likewise have other pregnancy-harming risk factors: enrolment in a non-white ethnic minority bunch, advanced maternal age, over the top body

weight, high equality, and significant financial disadvantage^{5,8}. An analogy can be drawn between these gamble variables and those associated with GDM. The pervasiveness of gestational diabetes is predominant among ladies who look for obstetric diabetes treatment at perinatal clinics, as the quantity of ladies with pre-gestational sort 2 diabetes right now surpasses that of ladies with type 1 diabetes in European urban communities because of segment shifts^{9,10}. Pregnant ladies who have type 1 diabetes, type 2 diabetes, or gestational diabetes are at increased hazard of developing constant intricacies. 40% of all pregnancies display indicators of sped up development, which influences around half of the infants brought into the world to moms with pre-gestational diabetes. Shoulder dystocia and caesarean segment rates exceeding 60% are associated not just with raised birth weight and neonatal adiposity, yet in addition with untimely beginning of heftiness, metabolic brokenness, and diabetes in the offspring during their later years (4). The kind of diabetes directs the obstetric and diabetic consideration approach.

Many forms of diabetes are seen in obstetric care.¹¹

Type 1 diabetes

The pancreatic β -cell is destroyed by the immune system, leading to an insulin shortage. In only 10% of instances, first-degree relatives are impacted, and symptoms usually appear before the age of 20. Not linked to

excessive fat storage. Almost five percent of all cases of diabetes that do not develop during pregnancy

Type 2 diabetes

Deficiency in relative insulin and insulin sensitivity impairment. Generally, symptoms become apparent subsequent to the 20th birthday, and over 50% of cases involve a first-degree relative. It has a robust association with adiposity. Approximately 90% of diabetes cases that do not manifest during pregnancy are this type.

Monogenetic diabetes

MODY stands for maturity-onset diabetes in young people. This condition develops as a result of a faulty production of insulin by pancreatic β -cells caused by a single gene mutation. Affected from birth and usually identified in a person's twenties or thirties. The condition is inherited autosomally, meaning that almost all cases have an afflicted first-degree relative. Unrelated to excess body fat. This condition is responsible for 1% to 2% of all cases of diabetes that do not occur during pregnancy.

Mitochondrial diabetes

The inability to secrete insulin is the result of a mutation in the DNA of the mitochondria. The risk of stroke, lactic acidosis, neural sensory deafness,

and other health complications are all closely linked. Diabetes, which is passed on from mothers to their children, often manifests around the age of 35. Unrelated to excess body fat. Out of all diabetes cases, this one accounts for less than one percent.

Secondary diabetes

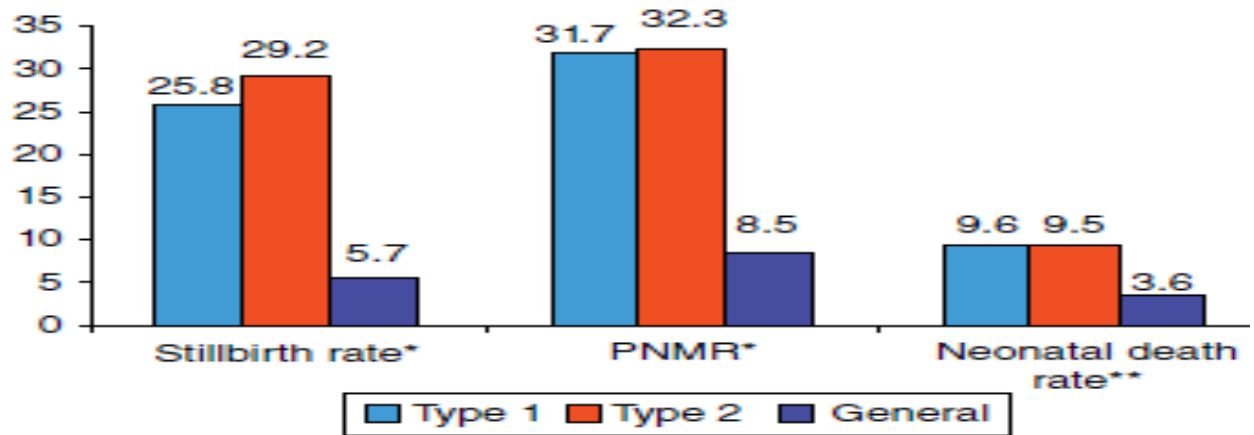
Another medical condition, such as pancreatitis, cystic fibrosis, glucocorticoids, or another medication, might cause diabetes. About 2% of all cases of diabetes not caused by pregnancy fall into this category.¹¹

Recent Guidelines for determination of GDM

Diabetes mellitus type 2 (GDM) occurs when either the mother develops or experiences her first symptoms of glucose intolerance while she is pregnant. In order to assess GDM using a 75-g OGTT, the following metrics are used: Fasting blood glucose readings of 5.6 mg/dl or higher and/or a 120-minute blood glucose level of 7.8 mg/dL as determined by recent in 2015. When using IADPSG, it is necessary to have blood glucose levels of at least 5.1 mg/dL when fasting, 10 mg/dL at an hour, and 8.5 mg/dL at 120 minutes. A higher gamble of gestational diabetes mellitus (GDM) is associated with being overweight (BMI > 30 kg/m²), having a background marked by gestational diabetes during a past pregnancy, having a kid who was 4.5 kg or heavier, having a direct relation with diabetes, or being from an ethnic foundation where diabetes is more normal.

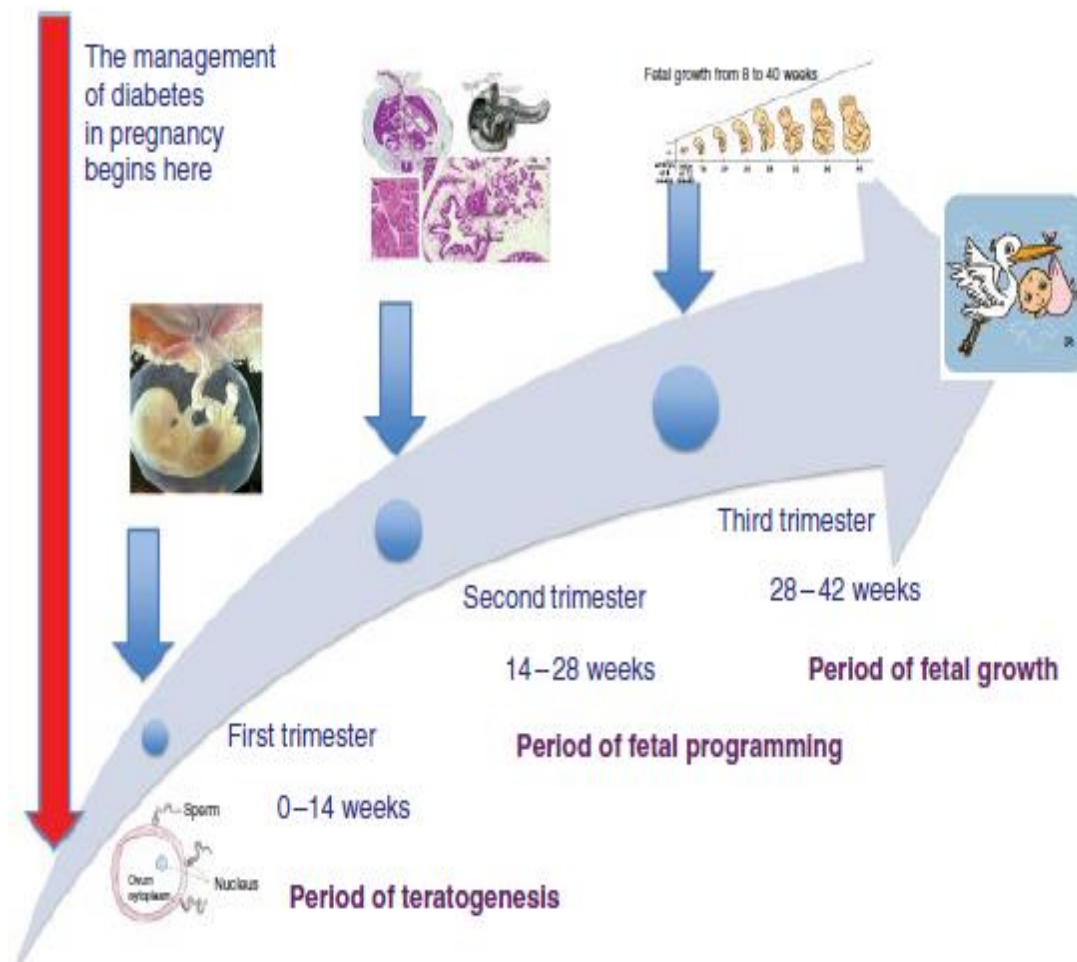
From Dewhurst's text book of obstetrics and gynaecology¹¹

CEMACH Report of PNMR in GDM



From Dewhurst's text book of obstetrics and gynaecology¹¹

Data assembled from the CEMACH Diabetes Review, which is a private investigation into maternal and youngster wellbeing, incorporates paces of perinatal mortality, stillbirths, and neonatal passings as well as paces of prenatal mortality and PNMR for moms with type 1 and type 2 diabetes mellitus (DM) who conceived an offspring or were booked to conceive an offspring in the Unified Realm between Walk 1, 2002, and February 28, 2003. The accompanying not entirely set in stone for each 1000 live births: maternal age changed rates per 1000 outright births ¹¹



From Dewhurst's text book of obstetrics and gynaecology¹¹

Fertilisation, implantation, and the whole pregnancy are all impacted by maternal hyperglycemia.¹¹

During the first twelve weeks of gestation, hyperglycemia begins to have its teratogenic consequences.

If a woman has diabetes and is of reproductive age, she should have pre-conception planning done as part of her diabetes management strategy.

Providing women with pre-conception counselling lowers the risk of congenital malformations and premature deliveries.

Optimal glycemic management, folic acid supplementation (5 mg), and discontinuation or substitution of any teratogenic medicines are all part of preconception care.

Women who are at high risk of pre-eclampsia, thrombophilia, or substantial proteinuria may need to take aspirin or heparin.

Maternal Hyperglycemia and its effects

Fertilisation, implantation, and delivery are all impacted by maternal hyperglycemia. Maternal hyperglycemia may raise the risk of obesity, diabetes, and other health issues in the offspring via altering gene expression during foetal programming, according to mounting data.³. Therefore, maintaining euglycemia during the whole pregnancy is a key component of diabetes care during pregnancy. Several complications, such as increased risk of miscarriage, fetal development acceleration, late stillbirth, delivery trauma, and newborn hypoglycemia, are associated with elevated maternal glycaemia. Having excess body fat may exacerbate this risk.¹² The maternal type of diabetes and its significance Maternal hyperglycemia, not type of diabetes, is the major factor in foetal disturbances, according to the 2002

CEMACH study of 3,800 pregnancies in the UK, which found near fetal catches (perinatal mortality, stillbirth, and neonatal mortality) among ladies with type 1 and type 2 DM.⁶

Women with type 2 DM are less prepared for pregnancy, bound to be socially hindered, and have a higher power of chunkiness, among other social and maternal bet factors that could impact pregnancy. In contrast, ladies with type 1 DM are less inclined to have these issues. Especially in danger are pregnant teenagers and youthful grown-ups with type 2 diabetes, who have an extremely low achievement rate and a fetal deformation pace of over 20%. Glycemic control is fundamental, yet so is the ID and treatment of specific diabetes problems.¹¹ Beginning at the top finish of the typical non-diabetic reach, the relationship between's pre-pregnancy glycaemia levels and innate irregularities starts.¹² Hyperglycemia has a teratogenic effect during blastocyst improvement, embryogenesis, and organogenesis, which occur in the initial twelve weeks of development. Ideal glycemic the executives before early fetal misfortune and inherent deformities. To do this, ideal for ladies have pre-gestational diabetes to painstakingly design their pregnancies, seek help prior to becoming pregnant, and continue to utilize contraception until they acquire incredible control of their glucose levels.¹³

As per the 2015 Pleasant rules, ladies who have a HbA1c level surpassing 86 mmol/mol (10%) are exhorted against imagining. Ideal pregnancy is

accomplished when the HbA1c falls under 48 mmol/mol (6.5%), as there exists a connection between's this limit and extreme fetal irregularities. Since glucose crosses the placental hindrance yet insulin doesn't, increases in maternal glucose cause hyperplasia of insulin-touchy tissues and the creation of insulin in the foetus.¹³

Hyperglycemia in the mother also increases the production of tumor necrosis factor, fetal insulin-like growth factor, and human placental growth hormone. These hormones, like insulin, operate as fetal growth factors, which may lead to complications such organomegaly, macrosomia, and accelerated fetal development. Birth malformations affecting the cardiovascular system common in mothers with hyperglycemia¹.

Heart irregularities including dextrocardia and transposition of the extraordinary supply routes are among the focal sensory system issues that might create, alongside anencephaly, spina bifida, hydrocephaly, and holoprosencephaly. Moreover, irregularities involving the genitourinary and skeletal frameworks might be available. Caudal relapse, an unusual birth imperfection, is more normal in kids brought into the world to moms with diabetes than in children brought into the world to sound moms by a variable of more than 200¹⁴. Issues like hypoglycemia, respiratory trouble, polycythemia, hypocalcemia, and hyperbilirubinemia are bound to happen in children whose moms have hyperglycemia or auxiliary fetal

hyperinsulinemia, as shown by the relationship between's the insulin or C-peptide levels in the blood upon entering the world and these results. Analysts are simply now beginning to check out at the drawn-out impacts of the mother's diabetes on the kid. Children brought into the world to mothers with diabetes have an increased gamble of inborn irregularities and a large group of metabolic diseases¹⁵.

At the point when the transfer of glucose from the mother stops and industrious post pregnancy fetal hyperinsulinemia continues, the most widely recognized of these circumstances is neonatal hypoglycemia. Fetal hyperinsulinemia, in which the mother supplies the developing child with an overabundance of metabolic substrates, is the main driver of these and other transient metabolic diseases in babies. Another normal condition is neonatal hypertrophic cardiomyopathy, which often endures long and doesn't makes no side effects except for may now and then prompt significant disease or passing¹⁶. New review shows that youngsters with a mother with diabetes, whether type 1 or type 2, are at a higher gamble of being overweight and experiencing metabolic irregularities, such glucose intolerance, as far as possible into their teen years. Children are bound to have issues in the event that their moms' diabetes isn't very much treated. Having a mother with type 2 diabetes mellitus increases a kid's gamble of getting the condition. Contrasted with their moms, these youngsters foster diabetes at a prior age.

Pregnant ladies who foster gestational diabetes are bound to have kids who are overweight, have type 2 diabetes, or experience the ill effects of metabolic disorder.¹⁷

It is conceivable that the multi-ethnic Age study will likewise show the connection between's gestational diabetes and a woman's weight quite early on. In this review, kids whose moms often had gestational diabetes mellitus had a faster pace of weight index (BMI) and development rate increase contrasted with unexposed controls, and this increase began between the ages of 10 and 13. Proof proposes that hereditary elements, related with the impacts of intrauterine hyperglycemia, add to the advancement of metabolic disorder and corpulence in the offspring of Caucasian ladies who have GDM. For instance, when contrasted with everyone, the gamble of heftiness is two times as high and the gamble of metabolic disorder is multiple times higher.¹⁸ Increased attention to the dangers associated with uncontrolled diabetes during pregnancy is important considering the way that this condition influences an increasing number of pregnant ladies and their unborn kids.

International Association for the Avoidance and Control of Diabetes (IADPSG) and the Populace Starting point for Wellbeing and Care Importance (Pleasant) in the Assembled Realm led a few gestational diabetes mellitus (GDM) assessments in 2015. The Fair's recommendations

incorporated a more noticeable fasting regard, a diminished 2-hour worth, and the disposal of a 1-hour worth. Research on money related intercession and assessments of monetary mindfulness turned into the basis of the Pleasant rules. The significant prenatal exam is an extraordinary spot to recognize risk factors, and the pleasant measures suggest organizing general screenings based on those discoveries. Some gamble factors include: a weight file (BMI) of at least 30, a background marked by gestational diabetes mellitus (GDM), a first-degree relative with the disease, racial or ethnic minority status, a birth weight of 4.5 kg or more due to macrosomia, etc. A 75-g OGTT ought to be taken by a lady between weeks 24 and 28 in the event that she follows any of these guidelines.¹³ Ladies who have gestational diabetes ought to have an oral glucose resistance test (OGTT) at something like four months into their pregnancy. Assuming the outcomes are standard, it ought to be rehashed in 24-28 weeks. Since it doesn't address maternal age or polycystic ovary condition, the Fair proposed outline of chance elements isn't intended to be comprehensive.¹⁹ Considering this, one ought to reasonably ask why, even with a positive outcome on an OGTT²⁰, evaluating for great bet factors alone could neglect to separate GDM in up to 25% of ladies. From 9.3 to 25.5% of basic HAPO accessory cases were GDM, as indicated by IADPSG rules, which differed by spot of beginning. Peri-conception haemoglobin A1c levels increased from 2 standard

deviations to 8 standard deviations over the normal 2% range, leading to an average foetal deformity incidence of 10% in a thorough evaluation of 1977 diabetic pregnancies from seven US cohort studies conducted between 1986 and 2006.²¹ There was a high link between prenatal glucose control and congenital malformations in 1677 infants delivered to diabetic mothers in England between 1996 and 2008. This finding is consistent with the overall population. Another possible predictor of stillbirth is high HbA1c levels during the periconceptional period.²²

One of the signs of GDM is unusual glucose tolerance, which either starts or manifests itself during pregnancy. Blood glucose levels in GDM don't ascend to the symptomatic limits for diabetes mellitus²³. Ladies with GDM are at a higher gamble of fetal macrosomia, toxemia, and cesarean segments during pregnancy.²⁴ Ladies who have gestational diabetes mellitus likewise have a more prominent chance of developing sort 2 diabetes mellitus later in life.²⁵

Organ improvement in a developing embryo happens all through the main trimester of a pregnant woman's life. During the initial three to about two months of pregnancy, fluctuating glucose resistance might prompt undeveloped improvement issues in some organs.²⁶ In this way, early gestational diabetes mellitus testing is fundamental for preventing fetal distortion. An exact indicator of pregnancy entanglements, the HbA1c level might be distinguished in the main trimester and can uncover toxemia, fetal

macrosomia, and enormous for gestational age.²⁷

Fasting blood glucose (FBG) testing is another early indicator of gestational diabetes. Fasting blood glucose (FBG) requires glucose subsequent to fasting, which makes it more factor and harder to rehash. This means it misses the early phases of GDM while screening. There are various advantages of comparing FBG levels with HbA1c values,²⁸ Each day, there is less pressure and infection, and it's more pragmatic since fasting isn't needed. Hemoglobin A1c (HbA1c) has seen broad use in the determination and management of diabetes mellitus. Be that as it may, as HbA1c readings decrease in the early phases of development, using them to analyze gestational diabetes is problematic.²⁹

During the primary trimester of pregnancy, ladies with a background marked by GDM ought to get locally situated glucose checking (HBGM) as well as guidelines for food and actual activity. Assuming that this neglects to accomplish the HBGM objectives, the following stage is to start taking an oral hypoglycemic medication, which could or probably wo exclude insulin. Oral hypoglycemics metformin and glibenclamide (showcased as glyburide in the US) are both protected and viable during pregnancy³⁰. Metformin represses gluconeogenesis in the liver, increases fringe glucose assimilation, and further develops insulin awareness. Metformin is an extraordinary decision for post pregnancy weight reduction since it is economical, easy to

take, only from time to time causes hypoglycemia, and is generally very much endured. page31. Increasingly more examination is shown that it benefits and doesn't harm pregnant ladies with polycystic ovaries, type 2 diabetes, and gestational diabetes mellitus, despite the fact that it isn't authoritatively authorized for use during pregnancy.³².

It is essential for the glycemic the board suggestions for type 2 diabetes and GDM in the Pleasant guidelines.¹³ There are continuous hypothetical issues about the use of metformin because of the way that it crosses the placenta. While there might be a few transient benefits, more examination is expected to decide the long haul effects.¹⁸ Glibenclamide might be endorsed to pregnant ladies who are restless about beginning insulin, despite the fact that it doesn't yet have business endorsement in the UK.³³ A far reaching ultrasound ought to be directed between the eighteenth and twentieth seven day stretch of growth to find any critical innate irregularities. Remembered for this ought to be an image of the heart's four chambers as well as the pathways for ventricular outpouring.

A raised gamble of toxemia and different types of hypertension is associated with diabetes in pregnant ladies. Risk assessment during pregnancy might be improved with the utilization of Doppler waveform investigations of the uterus at 20 weeks along with routine center circulatory strain and protein tests. Pregnant ladies frequently get ultrasounds at the finish of the

subsequent trimester and afterward at regular intervals (or all the more every now and again if necessary) to follow the child's advancement. Expected indications of development limitation or speed increase might be seen by contrasting the percentile assessment of the fetal stomach circuit at 28 weeks with checks performed at later weeks. Since polyhydramnios is more normal in diabetes pregnancies, keep liquid volume measurements in a specific sequence is fundamental.

Insulin obstruction and insulin needs both consistently ascend during the third trimester. Insulin obstruction safeguards ladies from the advantages of less complex glucose the executives. Toward the finish of the third trimester, insulin needs had begun to decline³³, which is an indication of serious fetal hyperinsulinemia, which decreases maternal blood glucose levels inferable from an increased glucose slope across the placenta or placental disappointment. A family as soon as possible in the event that your insulin needs start to decrease. An individual's insulin need can twofold in the following three days on the off chance that glucocorticoid chemicals are vital for lung improvement.

As a consequence of fetal hyperinsulinemia, a quickly growing fetal stomach boundary (AC) percentile comparative with biparietal width or head periphery is a typical finding in ultrasound studies. The mother's interior organs, including her heart and liver, experience additional strain because of

the developing child's insulin overproduction, and she likewise feels heavier than expected. The expression "macrosomia" is generally used to portray children brought into the world to diabetic mothers; in any case, its definite importance isn't clear. One bunch of measures incorporates outright birthweights that are under 4, 4.5, or 5 kg, while one more arrangement of models incorporates percentile birthweights that are over 90%, 95%, or 97.5%. Outright weight isn't the right measurement to use to characterize birthweight since it differs with gestational age, sex, identity, guardians' level and weight, and mother's weight. The example of fetal improvement has more clinical importance than the outright or percentile birthweight.⁶ Infants brought into the world to overweight mothers with type 2 diabetes or gestational diabetes mellitus type 6 some of the time have unusually high birthweights. Follicle development limitation or asymmetrical development limitation might be recognized all through pregnancies by sequential ultrasonography.³³

The fundamental goal of this study was to decide if first trimester hemoglobin A1c readings may be utilized as a predictor of GDM. Most of metabolic problems, like GDM, occur during pregnancy. A developing number of individuals are being determined to have GDM, and the repeat rates fall anyplace somewhere in the range of 9.32% and 5.5%.¹¹ Increases in maternal age and weight file (BMI), changes in diabetes side effect edges,

and more helpful admittance to prenatal screening tests are factors that could make sense of this improvement.³⁴ The major obstetric entanglements that could create from gestational diabetes influence the prosperity of both the mother and the unborn youngster. Most frequently, gestational diabetes might prompt complexities like toxemia, fetal macrosomia, polyhydromnios, neonatal hypoglycemia, an increased requirement for cesarean segments, and mischief to the mother and youngster.³⁵

The perinatal and post pregnancy results affected by gestational diabetes are associated with the level of the mother's uncontrolled hyperglycemia. To evade this issue, a few associations pushed for prenatal evaluating for glucometabolic jumbles. As a safeguard against unseen previous diabetes, proposals recommend holding back nothing HbA1c in the principal trimester. This is on the grounds that HbA1c is a measure of ordinary glucose over the past three months.³⁶ The perinatal and post pregnancy results influenced by gestational diabetes are corresponded with the level of the mother's uncontrolled hyperglycemia. To avoid this issue, a few associations supported for prenatal evaluating for glucometabolic jumbles. As a safety measure against unseen prior diabetes, proposals recommend holding back nothing HbA1c in the principal trimester. This is on the grounds that HbA1c is a measure of typical glucose over the past three months.³⁷ within the sight of specific gamble factors, the probability of

creating GDM is incredibly improved. Toxemia (OR2.7; 95CI:1.2-5.9) and shoulder dystocia/birth injury (OR24.5; 95% CI: 2.8-214.8) were still altogether associated with fasting glucose (FG) levels somewhere in the range of 5.6 and 7 mmol/L at the primary prenatal visit, even in the wake of representing maternal age and weight file.³⁶

Oral glucose resilience tests are frequently used to assess pregnant people for GDM. Then again, you'll have to fast for eight hours and have two blood tests done, and the test is very conflicting. Sadly, this intended that around 10% of pregnant ladies didn't finish their oral gestational tests.³⁷ When and how might the shipment be made? The dangers of conveying a pregnancy to term ought to be imparted to pregnant ladies with diabetes as early in the pregnancy as is basically conceivable. The best opportunity to talk about the particulars, such the conveyance procedure and when it will be, is after the 36-week development examine, when both the mother and the child can pursue better decisions, despite the fact that ladies ought to in any case be participated here and there with their introduction to the world plans. In the diabetic populace, the opportunity of late shock stillbirth is multiple times more prominent than in the non-diabetic population.^{2,5}. Subsequently, it is prescribed that insulin-subordinate mothers endeavor to give conveyance somewhere in the range of 38 and 39 weeks, as per most specialists. Prompting work in a lady with type 1 diabetes who has never conceived an

offspring before might be a long and fruitless strategy. A cesarean segment is essential after a bombed enlistment endeavor in practically half of pregnancies including ladies with type 1 diabetes⁵.

Then once more, ladies with type 2 diabetes will without a doubt have strong enlistments of badly designed work and to have various pregnancies. Ladies whose GDM is exclusively obliged by diet or metformin could have the decision to appear at a safe gestational age (40 weeks) and expect an unconstrained work in the event that their pregnancies are overall clear. A more conspicuous birth weight is connected to a higher bet of birth hurt. Maternal diabetes mellitus is associated with an increased bet of shoulder dystocia. This is to some degree made sense of by the way that the chest-to-head and shoulder-to-head degrees are higher in the posterity of diabetes moms. Youths brought into the world a few spot in the scope of 4 and 4.5 kg have a 3% opportunity of shoulder dystocia, whereas those brought into the world after 4.5 kg have a 10% to 14% possibility. An arrangement that just utilized the common not entirely set in stone by ultrasonography (4000 or 4500 g) would, regardless, lead to an unsuitable speed of cesarean segments. Choices about the planning of the transport of a giant youngster recognized by ultrasound ought to consider the diabetic mother's obstetric history, her body size, and her tendencies.

Norms for the organization of insulin to pregnant ladies Hypoglycemia is

bound to happen in newborn children whose mothers had high glucose levels during work and conveyance. In this manner, as per the 2015 Decent rules, ladies with diabetes ought to check their fine plasma glucose levels all through work and delivery¹³ to guarantee that their glucose levels stay inside the prescribed scope of 4 to 7 mmol/L. Since most diabetic ladies will be educated regarding their due date, whether by decision or cesarean area, there ought to be clearly characterized shows for the association of insulin needs during work for both the lady and the work ward. Regardless of the way that there is nobody most effective way to oversee insulin during this time, it is normal practice for the mother to begin a variable-rate intravenous insulin infusion (VRIII) with 5% dextrose before she begins to eat during dynamic work and a usable development. It is supported to dependably screen blood glucose levels all through the range of this time. Since insulin requests would be 20-30% lower during pregnancy contrasted with pre-pregnancy levels, ladies who were taking insulin before to the pregnancy ought to reduce their measurement assesses following start. Since most pregnant ladies take long-acting basal insulin around night time major areas of strength for and insulin with their night feast going before pondering having a youngster, it is suggested that ladies who are hospitalized for a booked assurance take their standard insulin conveys at night. If the patient is pregnant or plans to have a cesarean section, short-acting bolus insulin for

feasts while in the facility, and they might be encouraged to change to VRIII. Cautious glucose screens and insulin siphons are turning out to be increasingly standard in the work environment as means great limits.³⁹

Heaps of ladies with GDM don't need intravenous insulin, taking eating regimen pills or metformin. Ladies with insulin-treated GDM who need under 1.0 unit/kg of insulin routinely may frequently be overseen without intravenous insulin, but a VRIII may possibly be started if vital. An elective cesarean area is best acted toward the beginning of the day. It is suggested that the patient take her ordinary bolus insulin with her feast and a critical part of her basal insulin the night prior to the strategy. You might begin a VRIII while on the ward and deny the evaluations as soon as you arrive. The child's insulin necessities return to what they were before pregnancy as soon as the placenta is conveyed. Ladies who have type 1 diabetes during pregnancy might keep taking the insulin they were taking previously, yet they ought to lessen their dose by something like 25% once they can eat and drink ordinarily once more. Without a doubt, even in ladies with very much controlled diabetes before to pregnancy, the all out day to day insulin segment frequently decreases during pregnancy, with the bolus insulin portion seeing the best decrease (40).

Make a point to incorporate the insulin portion for after the child is conveyed in the birth plan. When a lady conceives an offspring, she might return to

taking her oral prescriptions for type 2 diabetes to nurture during pregnancy. In the event that she wasn't utilizing insulin, she can stop. Taking insulin during breastfeeding is totally sans risk since the medication doesn't enter breast milk. Insulin necessities in moms with type 1 diabetes frequently decrease while nursing because of a little increase in metabolic rate. Metformin is likewise viewed as safe for nursing moms to drink as a result of its low fixation in breast milk.¹³ The 2015 Pleasant rules recommend only two oral specialists — glibenclamide and metformin — for the administration of glucose levels during breastfeeding³². During the post pregnancy time, it means quite a bit to give direction on measures to forestalling beginning. The executives of diabetes mellitus during pregnancy...A fasting blood glucose level under 5.3 mmol/L and a 1 hour postprandial glucose level under 7.8 mmol/L should be the comparable glycemic targets for pregnant women with GDM as they are for other diabetic women.¹³

Numerous ladies with GDM depend on insulin to accomplish their weight reduction and action objectives, regardless of whether others might be moved to do as such by way of life changes. In a few nations, individuals actually question the real role of experts in oral hypoglycemia. Many actually believe that sulphonylureas, regardless of metformin, are less effective and less adaptable than insulin solutions, despite the fact that the

latest exploration on this subject proposes that they are protected when utilized following 15 weeks of hatching. On the other hand, oral hypoglycemic meds are a lifeline in areas where insulin is scant or nonexistent. Subsequent to giving conveyance, numerous ladies with GDM who don't have undiscovered diabetes recover ordinary glucose resilience. Anyway, during the accompanying twenty years, the larger part of everybody will develop sort 2 diabetes⁴¹. Conveying the diabetes quality during pregnancy is critical areas of fortitude for an of creating sort 2 diabetes later on. A month and a half straightforwardly following conceiving an offspring, and again dependably beginning there forward, all pregnant ladies ought to have their fasting glucose or hemoglobin A1c tried¹³.

One of the aims of lifestyle treatments that try to restrict weight gain and encourage physical activity is to lower the chance of acquiring diabetes during the following four to five years¹⁵. Therefore, it is essential that postpartum women with GDM get basic lifestyle advice and learn about the requirement of diabetes testing annually.

Life style intervention and management of GDM

Compared to the general population, women with diabetes have almost a fourfold increased chance of having a baby that is born unexpectedly and at a late stage.

Between weeks 38 and 39 of gestation, women using insulin should give birth.

It is possible for women to safely reach the 40th week of their pregnancies when they are managed with metformin or if they rely only on nutrition.

For women who have pre-gestational diabetes, the first step after giving birth is to lower their insulin dosage to 25-30% of what they were taking throughout pregnancy.

Women with gestational diabetes mellitus (GDM) may usually cease taking their diabetic medicines after giving birth. Some women with type 2 diabetes mellitus (DM) can even stop using insulin.

Nursing mothers may safely use insulin, Glibenclamide, or metformin.

Haemoglobin A1c (fasting blood glucose level) testing is necessary for women with gestational diabetes mellitus (GDM) six weeks after giving birth and annually after that.

To primarily identify type 2 diabetes mellitus, glycosylated hemoglobin (HbA1c) is used. The HbA1c test shows the average glucose concentration over the last two or three months, in comparison to FBS and OGTT. Due to its reduced intra-person changeability and the fact that fasting, the test is more suitable for all patients. Despite the fact that HbA1c seems to be a straightforward, valid, and accommodating lab test in GDM diagnosis, the feasibility of using first-trimester HbA1c well evaluated, and none of the

existing guidelines for GDM address this issue.^{27,32,37}

We want to assess the demonstration tool for first-trimester HbA1c in early pregnancy in this way. Screening for gestational diabetes and other important components of evaluating diabetic pregnancies, beginning at 6–12 weeks of gestation and continuing through 26–28 weeks, have been emphasized in this dissertation. Predictor of GDM by evaluating the threshold values for diagnosing and prescribing first-trimester GDM.

AIMS & OBJECTIVES

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at the right end of the horizontal line. The horizontal line starts under the 'A' of 'AIMS' and extends to the right edge of the page. The vertical line starts from the top of the page, goes down, crosses the horizontal line, and continues to the bottom of the page.

AIM AND OBJECTIVES:

- To evaluate the HbA1c in all to begin with first trimester pregnant women.
- To assess the connection of HbA1c in early pregnancy (8-12 weeks) and expectation of development of diabetes mellitus in second trimester (24-28 weeks) of pregnancy with oral glucose tolerance test (OGTT).

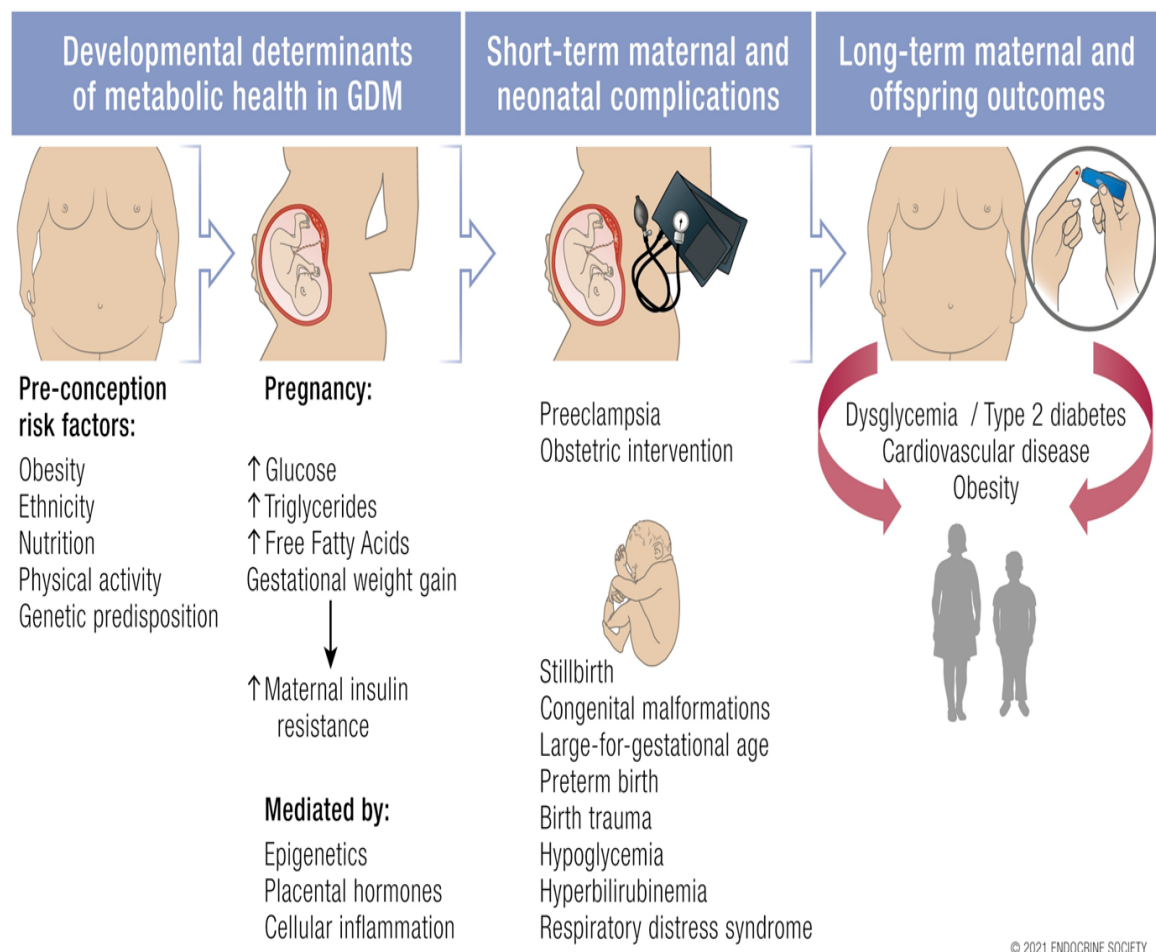
REVIEW OF LITERATURE

A decorative graphic consisting of a thick horizontal line and a thick vertical line intersecting at a right angle. The horizontal line is positioned below the word 'LITERATURE' and extends across the width of the page. The vertical line is positioned to the right of the horizontal line and extends upwards, crossing the horizontal line.

REVIEW OF LITERATURE

Diabetes or glucose narrow mindedness that starts during pregnancy or is first identified during pregnancy is known as type 2 diabetes mellitus (GDM). This gathering incorporates individuals whose diabetes was undetected for quite a while. The increasing commonness of heftiness is generally liable for the ascent in the level of ladies determined to have pre-gestational sort 2 diabetes mellitus (DM) and gestational diabetes (GD). Type 1 diabetes mellitus is likewise on the ascent, however more leisurely, with an expected 3-4% yearly event in Europe. An ever increasing number of pregnancies will happen as the normal age at which type 2 diabetes starts to show declines. Physiological changes in the mother's glucose digestion frequently cause gestational diabetes mellitus to foster in the last 50% of the subsequent trimester (10). It is basic to distinguish this high-risk bunch with the goal that clinical treatment can mirror the subject of who ought to be screened and how⁵³. In situations where the going with conditions are fulfilled when not pregnant: a plasma glucose level of 7.0 mmol/L or higher while fasting; a plasma glucose level of 11.1 mmol/L or higher following 2 hours during a 75-g oral glucose versatility test (OGTT); an unpredictable plasma glucose level of 11.1 mmol/L or higher when hyperglycemic secondary effects are free; and a hemoglobin A1c level of 6.5% or higher, which is comparable to 48 mmol/mol or earlier.

Clinical appraisal of the patient's experience plan and acknowledged risk factors may habitually assist with perceiving type 1 and type 2 diabetes mellitus. Tests for the specific pancreatic islet autoantibodies Stray, IA-2, and ZnT8 could attest the finish of type 1 diabetes in around 80% of situations where this is impossible. White people could address as much as half of the instances of type 1 diabetes mellitus among adults who get the condition after the age of 30, according to mounting confirmation.⁸



- From Dewhursts text book¹¹

Developmental determinants of Metabolic health in GDM and its complications

A very dangerous complication of GDM is preeclampsia. Preeclampsia afflicts about 10% of GDM sufferers. In addition to pregnant weight gain, preeclampsia was more prevalent in younger, obese, and nulliparous women with GDM. When proteinuria is present and diabetes is known to induce nephropathy, this becomes an even greater concern. Microalbuminuria during the first trimester of pregnancy increases the chance of superadded preeclampsia by 35-60%. Preterm birth, chorioamnionitis, polyhydramnios, and UTIs are all issues that might arise during pregnancy when a woman has diabetes. In situations of premature labour, tocolytics, steroids, and beta2 agonists may amplify hyperglycemia and increase the likelihood of ketoacidosis. Miscarriages are more common in women whose diabetes is not well managed before becoming pregnant. In order to treat nephropathy with angiotensin II receptor blockers or angiotensin-converting enzyme inhibitors—which raises the risk of preterm delivery and other complications—it is vital to measure renal function previous to and during pregnancy. Diabetic nephropathy risk or advancement may be mitigated or halted with better glucose management and hypertension medication. Because of the risk of fetal proximal renal tubular dysgenesis and oligohydramnios, these medications should not be used while pregnant.

Hence, further pharmacological therapy with drugs like labetalol or alpha methyldopa has to start right away if a pregnancy is diagnosed.

An important maternal health concern is GDM. Severe perinatal problems may occur if screenings, diagnoses, and treatments are delayed. It is linked with the following irregularities: encephalocele, meningomyelocele, and anencephaly are all birth abnormalities affecting the central nervous system. cardiovascular problems (e.g., hypoplastic left heart, ventricular and atrial septal abnormalities, transposition of great vessels, and others) Physical: spinal abnormalities, caudal regression syndrome Urinary tract: cysts, leaky kidneys, renal agenesis Internal: anorectal deformity, duodenal atresia Pregnant women with glycosylated hemoglobin levels in the first trimester may have a better idea of how likely it is that their unborn child will have a congenital abnormality. moms with hemoglobin A1c levels below 7% are at the same risk for birth defects as moms without diabetes, according to studies. The likelihood of abnormalities increases to 22% at 1.10%, from 5% in 7-8.5%.⁸

The effect of hyperglycaemia in the mother on complications for the mother and foetus	
The impact of hyperglycaemia	
1st trimester	
Implantation	Trachectoderm differentiation is inhibited.
Embryogenesis	Elevated levels of oxidative stress impair the

	expression of vital genes that are indispensable for embryogenesis.
Organogenesis	Congenital defects are exacerbated as the diacylglycerol-protein kinase C cascade is activated.
Miscarriage	Elevated premature programmed cell demise of primordial progenitor cells
2nd trimester	
Endocrine pancreas	Inhibits foetal B-cells
Fetal growth	Induces foetal hyperinsulinemia, which manifests as an ultrasound-detected acceleration in growth by 26 weeks.
3rd trimester	
Fetal growth	A significant foetal substrate and determinant of rapid foetal development
Adipose disposition	Inducing hyperinsulinemia, which facilitates fat elimination, including that of fat within the abdomen.
Lung maturation	Promotes hyperinsulinemia, which inhibits surfactant proteins and thereby delays lung maturation.
Stillbirth	Is correlated with placental maturation abnormalities that elevate the likelihood of foetal hypoxia.
Delivery	
Birth trauma	Accelerating foetal growth increases the likelihood of developing shoulder dystocia, which increases the susceptibility to birth trauma and asphyxiation.
Neonate	
Hypoglycaemia	Induces foetal hyperinsulinemia, which increases the risk of hypoglycemia in the neonate.
Hypoglycaemia	Modifies the expression of calbindin mRNA in the placenta, which influences the calcium status at birth.
Polycythaemia	Similar to foetal hypoxia, it induces foetal hyperinsulinemia, which improves antepartum haemopoiesis.
Cardiomyopathy	Induces foetal hyperinsulinemia, which increases the

	risk of developing hypertrophic cardiomyopathy.
Adolescence/adulthood	
Obesity	The development of the metabolic syndrome is predisposed to intrauterine exposure, irrespective of any genetic predisposition.
Type 2 diabetes	Type 2 diabetes risk is increased by intrauterine exposure, irrespective of genetic susceptibility.

From Dewhursts text book of obstetrics and gynaecology¹¹

An examination of the correlation between antenatal glycosylated hemoglobin (HbA1c) and congenital abnormalities³ It is essential to test for and treat particular diabetic complications in addition to glycemic control. Birth defects are more likely to occur in women whose blood sugar levels are high before pregnancy, and this association begins at the higher end of the normal non-diabetic range¹². During the first twelve weeks of gestation, hyperglycemia has teratogenic consequences that become apparent during blastocyst development, embryogenesis, and organogenesis. Early foetal loss and congenital abnormalities may be significantly reduced if pregnant women maintain proper glycemic control. It is best for women with pre-gestational diabetes to obtain advice before becoming pregnant, plan their pregnancies carefully, and use contraception continuously until their blood sugar levels are under control¹³ The prevalence of congenital abnormalities and premature delivery is inversely correlated with preconception counselling (42). These clinics create a safe space for pregnant women to discuss their options for prenatal care, stress the need for close monitoring of

both the mother and the unborn child, and reassure them that their blood sugar levels can be managed to reduce the risks of a diabetic pregnancy. One possible approach is to prescribe 5 mg of high-dose folic acid supplements and increase glycemic control. Another is to quit utilizing or find more secure options in contrast to prescriptions that might cause pregnancy issues.⁴³ Pregnant ladies who have over the top proteinuria (>4 g each 24 hours), are at high gamble of toxemia or thrombophilia, or who have diabetes could have their condition assessed to see whether aspirin or heparin is fundamental.¹³

Treatment and determination of gestational diabetes mellitus further foster pregnancy results, according to two significant randomized controlled investigations. One analysis was finished in Australia, and the other in Canada. In these two examinations, which used different models to analyze GDM, pregnant women were randomly assigned to get either standard pre-birth care or dynamic treatment, which included dietary changes, glucose monitoring, and insulin, if needed⁷. Following change for maternal age, race or ethnic gathering, and balance, the general bet of outrageous perinatal challenges was 1% in the 490 pregnancies assigned to intervention care contrasted with 4% in the 510 pregnancies assigned to routine thought in an Australian preliminary of 1000 women determined to have blocked glucose tolerance according to World Wellbeing Organization (WHO) measures. The

speed of induction of work was 10% higher in the intervention bundle contrasted with the normal thought pack, whereas the speed of cesarean sections remained unchanged⁷.

Dynamic treatment diminished a few elements, as per a Canadian survey incorporating 958 ladies with gentle gestational diabetes. These incorporated the “mean birthweight (3302 g versus 3408 g), neonatal fat mass (427 g versus 464 g), repeat of huge for-gestational-age infant youngsters (7.1% versus 14.5%), birthweight more prominent than 4000 g (5.9% versus 14.3%), shoulder dystocia (1.5% versus 4.0%), and the” requirement for cesarean conveyance (26.9% versus 33.8% in the benchmark group). Treatment with dynamic diminished the rate of blood poisoning and gestational hypertension in women.³⁸ In excess of 23,000 non-diabetic ladies' pregnancies were considered from 2000 to 2006 by analysts in nine unique countries. Between the 24th and 32nd seven day stretch of incubation, all pregnant ladies were given a 75-g OGTT that lasted for two hours. Macrosomia (adjusted birthweight >90th centile), essential cesarean conveyance, are four significant perinatal results that ought to be better perceived assuming moms have glucose levels underneath the diabetes demonstrative edge. Fasting glucose levels, levels tried one hour after a 75-g glucose burden, and levels recorded two hours after the heap were undeniably associated with the four essential results. These discoveries

educated the improvement regarding refreshed GDM indicative models by the Global Association of Diabetes and Pregnancy Study Gatherings (IADPSG), which have now gotten endorsement from the American Diabetes Association (ADA) and the World Wellbeing Association (WHO). Yet, in 2015, the English Public Organization for Wellbeing and Care Greatness (Decent) recommended new standards for GDM. Among them, there was a more limited 2-hour esteem, a bigger fasting esteem, and the 1-hour esteem was taken out. These are unmistakable from the measures gone ahead by the IADPSG. While the Pleasant models don't cover ladies with fasting glucose levels somewhere in the range of 5.1 and 5.6 mol/L, the IADPSG measures incorporate. In spite of a 2-hour esteem beneath 7.8 mmol/L, 38% of these pregnancies have children who are huge for their gestational age, and these moms have an increased gamble of becoming obese⁵⁵. A disagreeable conversation perseveres over the treatment's viability for ladies whose glucose bigotry falls underneath the Pleasant rules.⁵⁵.

dealing with a diabetes pregnancy and what it means for both the mother and her unborn kid Babies with diabetes mothers are at increased risk for various metabolic issues and inborn abnormalities⁵⁶. Neonatal hypoglycaemia is the most well-known kind, and it happens when the mother's glucose transport stops and the embryo keeps on having tireless post pregnancy

hyperinsulinaemia. Fetal hyperinsulinaemia, in which the mother supplies the creating child with an abundance of metabolic substrates, is the main driver of these and other transient metabolic diseases in babies. Another normal condition is neonatal hypertrophic cardiomyopathy, which might cause passing or serious disorder in uncommon cases however frequently just lasts a brief time and has no side effects at each of the 57. Late studies¹⁸ shown that offspring of moms with type 1 or type 2 diabetes had a higher commonness of corpulence and metabolic issues in puberty, like glucose bigotry. Complexities are bound to arise in babies whose moms' diabetes is ineffectively controlled. Having a mother with type 2 diabetes mellitus increases a kid's gamble of getting the condition. These children get diabetes at a prior age than their mothers did¹¹. Children whose moms had diabetes during pregnancy had an increased gamble of becoming overweight, creating type 2 diabetes, and encountering metabolic disorder in advanced age. There might be a relationship between's gestational diabetes and youth weight, as per the multi-ethnic Age research. Weight list (BMI) and development speed increased all the more quickly in kids whose moms for the most part had gestational diabetes mellitus (GDM) contrasted with unexposed controls, and this speed increase started between the ages of 10 and 13. There is solid proof that the impacts of intrauterine hyperglycemia and hereditary inclination put the offspring of Caucasian ladies with gestational diabetes

mellitus at a higher gamble of creating metabolic condition and heftiness. When contrasted with everybody, the probability of stoutness is multiplied and metabolic disorder is quadrupled. Creature studies have shown that embryos whose moms had diabetes during pregnancy are bound to have disabled glucose resilience, diabetes, and overabundance muscle versus fat. It is challenging to unravel the impacts of intrauterine hyperglycemia from acquired impacts in people, and proof recommends an epigenetic system of diabetes move from mother to kid because of perinatal programming of the hatchling. This features the need of better glycaemic the executives all through each pregnancy including a diabetic mother. The gamble of late-term surprising stillbirth is multiple times higher in the diabetic populace contrasted with everyone.

The frequency of fetal confusions, like perinatal passing, stillbirth, and neonatal mortality, were comparable in ladies with type 1 and type 2 diabetes mellitus in the 2002 CEMACH review of 3800 pregnancies in Britain, Ridges, and Northern Ireland. This features the way that kind of diabetes isn't the fundamental element deciding fetal inconveniences, but instead maternal hyperglycemia.⁶ Contrasted with other pregnant ladies, those with type 2 diabetes are more inclined to be overweight, socially impeded, and ill-equipped. Possibly influencing the pregnancy are these and other cultural and maternal gamble factors. On the other hand, these issues are more

uncommon in ladies who have type 1 diabetes. Pregnancy confusions and anomalies in the posterity are more normal among teenagers and youthful grown-ups who experience the ill effects of type 2 diabetes. Type 1 diabetic ladies are at increased risk for retinopathy and extreme episodes of hypoglycemia.

Both glycemic control and the discovery and the board of explicit diabetic confusions are of most extreme significance. The association between pre-pregnancy glycaemia levels and inherent contortions starts at the most elevated place of the ordinary, non-diabetic range¹². During the initial twelve weeks of development, hyperglycemia has a teratogenic impact on blastocyst development, embryogenesis, and organogenesis. To extraordinarily decrease the probability of unexpected labor and other birth irregularities, ladies ought to hold back nothing control preceding getting pregnant. This might be accomplished assuming ladies with pre-gestational diabetes make the accompanying strides: they ought to look for advising prior to getting pregnant, cautiously plan their pregnancies, and keep utilizing contraception until their glucose levels are under extraordinary control.¹³ years old. To more readily oversee glucose levels, it is prescribed to take folic corrosive enhancements at a high dose of 5 mg. Meds like angiotensin-changing over protein inhibitors ought to be either halted or supplanted with more secure other options. 43, 44. It very well might be

important to assess the aspirin or heparin necessities of pregnant ladies who are at high gamble of toxemia, thrombophilia, huge proteinuria (>4 g each 24 hours), or diabetic complications.¹³ Ladies from ethnic minorities, the most financially burdened, and those with type 2 diabetes are excessively underrepresented in the utilization of these administrations. As per the 2015 Pleasant suggestions, ladies shouldn't become pregnant on the off chance that their HbA1c is higher than 86 mmol/mol (10%). Preferably, they ought to attempt to keep their HbA1c under 48 mmol/mol (6.5%), as this level is associated with critical birth defects.⁶ As indicated by the ongoing rules, ladies with type 1 diabetes ought to abstain from eating anything with a glucose level lower than 4-7 mmol/L before feasts and between 5-7 mmol/L subsequent to awakening from fasting. Ladies with type 1 diabetes who need to accomplish this level of glycemic control need either a subcutaneous insulin blend siphon (CSII) or multi-segment insulin (MDI), which requires infusion four to multiple times everyday. Regularly, patients on the MDI routine are given a straightforward basal insulin with a long-acting impact either at the same time toward the beginning of the day or separated into two portions over the course of the day. Injectable bolus insulin, which acts rapidly, is taken with every feast. A few devices that could be useful to ladies keep up with better control incorporate glucose screens, insulin bolus number crunchers that propose compelling insulin measurements based on

carb admission, and novel insulin siphon creations. Enhancements in shut circle insulin conveyance frameworks that coordinate PC estimations for persistent nonstop glucose observing (CGM) with CSII can possibly normalize glycaemic control and lessen the gamble of hypoglycemia later on.⁵⁹

Type 1 and type 2 diabetic issues manifest diversely in females. Diabetic microvascular disease is more normal in ladies with type 1 DM than to type 2 DM. The more extended a lady has type 1 diabetes and the higher her gamble of creating diabetic entanglements, the more probable it is that this will be the case. The circumstance might move assuming the pace of new occasions of type 2 diabetes continues onward down. Treatment of diabetic retinopathy and nephropathy during pregnancy has the possibility to slow their turn of events. Subsequent to controlling for other gamble factors, the gamble of coronary course calcification and coronary vein disease (computer aided design) is more noteworthy in ladies who have type 1 diabetes for over 20 years and are as of now 40 years of age. Coronary supply route disease (computer aided design) is bound to happen in more seasoned ladies who have type 2 diabetes or gestational diabetes mellitus. There is major areas of strength for a between maternal dismalness and a background marked by coronary vein disease (computer aided design) or a conclusion of computer aided design during pregnancy. Since insulin can't pass the placental

boundary, a height in maternal glucose levels triggers insulin creation by the embryo and the hyperplasia of insulin-delicate tissues⁶². Maternal hyperglycemia additionally upgrades the creation of fetal insulin-like development factor, cancer putrefaction element, and human placental development chemical. Similarly that insulin is a fetal development factor, these chemicals might cause organomegaly, macrosomia, and sped up fetal turn of events. Resting alone, renal hindrance, autonomic neuropathy, gastroparesis, and a long history of type 1 diabetes are additionally perceived risk factors. Taking fast-acting insulin during pregnancy was associated with less serious hypoglycemia episodes during the night⁶. Individuals ought to be instructed on the most proficient method to give glucagon, and individuals ought to get directing on when to eat and nibble steadily and when to exercise to bring down their gamble of hypoglycemia. All ladies on insulin ought to continuously have a card on them with their solution data and how to address hypoglycemia. Try not to drive in the event that you are encountering a hypoglycemic episode, whether or not you feel any side effects or not. Glucose levels in pregnant ladies who don't have diabetes ascend after feasts however fall in the wake of fasting when the subsequent trimester begins. Because of increased degrees of placental chemicals and coursing maternal unsaturated fats, the mother's insulin opposition begins to develop during the center of the subsequent trimester.

Hereditary and epigenetic elements may both play a part in the beginning of GDM, as per concentrates on led regarding the matter by Rohini HN, Punita P, and others. While hereditary elements are frequently connected to things like quality transformations or single-nucleotide polymorphisms (SNPs), epigenetic factors incorporate things like quality methylation, histone change, and microRNAs that tight spot to courier RNA (mRNA).¹¹

In South Indian ladies, GDM was associated with the CDKAL1 SNPs rs7754840 and rs7756992, whereas in GDM and T2D, hereditary variations in the HMG20A (rs7178572) and HNF4A (rs4812829) qualities were demonstrated to be related.³⁹ Studies directed in Rome found an association between IRS-1 quality polymorphisms and type 2 diabetes.¹² Six unmistakable quality polymorphisms were associated with GDM in a meta-examination. A portion of the qualities were MTNR1B, TCF7L2, IRS-1, IGF2BP2, and TNF-alpha. Besides the fact that GCK associated with is GDM, yet KCNJ11 and CDKAL1 are as well.¹²

Following a case-control research, researchers in India found that gestational diabetes mellitus (GDM) patients had higher measures of microRNA7 in their placental tissue, maternal blood, and line blood. An increase in miRNA7 was recommended by the decreased articulation of IRS1, IRS2, and RAF1 in maternal blood. One examination out of India observed that GDM was additionally connected to mitochondrial tRNA quality changes or

inclusion cancellations. There is some proof that varieties in the hexokinase space containing 1 (HKDC1) quality might add to GDM. Our work gives the principal proof connecting this quality to GDM.

Popova PV, Smile Eva EN, and others concentrated on their associations with GDM risk to lay out the ease of utilizing typical genetic gamble types to recognize cases of GDM in Russian ladies. Two MTNR1B variations in Russian ladies were displayed to have areas of strength for a with GDM, out of eleven SNPs associated with type 2 diabetes and GDM in different populaces. In any case, these progressions didn't work on the exactness of distinguishing GDM patients. I wouldn't uphold the continuous utilization of GRS in GDM expectation because of the restrictive cost of hereditary testing and its restricted handiness in distinguishing GDM cases. Entire genome sequencing or GWAS might simplify it to disentangle the hereditary basis of GDM in Russia.

Fadl HE, Simmons D. Drifts et al. seen that the prevalence of robustness is straightforwardly related with the increase in the amount of cases of diabetes during pregnancy. Changes in GDM, type 1 and type 2 diabetes, and other pregnancy issues were centered around in a statewide group based center around in Sweden from 1998 to 2012. The data used to break down this all inclusive community based associate came from Sweden's public clinical birth register. The decided lose the faith took the mother's age, character, and

BMI into thought straightforwardly following arranging the timeframes into 3-year stretches. There was an increase in each sort of diabetes all through the 15 years contemplated. The pervasiveness of type 1 diabetes was for the most part significant among ladies of Nordic fall. The increase in GDM and, to a lesser extent, type 2 diabetes, may be made sense of by factors, for example, start, weight record (BMI), and mother age. The speed of new cases of gestational diabetes mellitus all through that period was genuinely predictable in the Nordic countries. These outcomes show that over the 15-year timespan, the transcendence of a broad assortment of diabetes during pregnancy increased in Sweden. In a general sense, the increase in gestational diabetes and type 2 diabetes might be credited to the mother's pre-pregnancy weight file (BMI). To stop the disturbing ascent of diabetes during pregnancy and the ominous results it has on both the mother and the young person's prosperity, we should execute individuals based intercessions to cut down the probability of becoming pregnant with a raised weight record.

Assessing for potential catches of GDM: a couple of things shouldn't in any way unequivocally settled at the head pre-birth plan. (This is a section from the Techniques for the Fourth overall Studio on GDM, which occurred in August 1998). All pregnant ladies ought to be pursued for sickliness at their most basic pre-birth framework. A couple of ethnic social occasions might be

at a lower risk than others relying on their weight report. 1 Hyperglycemia, polycystic ovary tangle, cardiovascular disease, hypertension, unprecedented lipid levels, a first-degree relative with diabetes mellitus, a birth weight of 4000 gm or more, an establishment put away by GDM or prediabetes, a stillbirth or an unexplained irregularity following entering the world, glucose in pee, the utilization of plans that raise glucose levels (such steroids, beta-mimetics, or odd antipsychotics), momentous lipid levels, or cholesterol levels.

Low risk: When all of the following are true, it is not necessary to frequently measure blood glucose levels.

Identified as belonging to a population where type 2 diabetes is uncommon.

We do not have any first-degree relatives who have diabetes.

Young adulthood.

Pregnancy weight is typical.

Delivery weight is within the typical range. I have no prior records of glucose metabolism issues.

Exceptional obstetrical success rate not documented.

Average risk: Between weeks 24 and 28, use one of the two-step strategies to check blood glucose levels. Individuals who show up at the cutoff regard in the GCT one-step method will do a suggestive 100 gms OGTT resulting to completing a 50 g oral glucose challenge test. Test 100 milligrams. OGTT

was administered to each individual.

High risk: If you notice any of these aftereffects, it's important to test your blood glucose levels right once using the methods referred to beforehand. Very overweight, with a substantial genetic inclination to type 2 diabetes and a foundation set apart by GDM. Glycogen abundance with crippled glucose processing. Rehash blood glucose testing should be performed at 24-28 weeks incubation or whenever incidental effects or indications indicate hyperglycemia if GDM isn't as of now analyzed.

19 Williams' Obstetrics, 26th Release — Investigation coordinated by Feig DS et al. distinguishes different bet factors for GDM, including heftiness, a foundation set apart by GDM in past pregnancies, a first-degree relative with diabetes, and ancestry from a high-risk racial or ethnic gathering. A couple of clinical organizations are pushing for risk-based screening to begin in the main trimester of pregnancy rather than the customary 24-multi week range.

25 To this end many biochemical indicators have been the point of convergence of investigation into potential screening gadgets. The examinations show that women with diabetes often have intricacies during pregnancy. Scientists in Ontario, Canada, examined information from 1,109,605 mothers who considered an offspring between the years 1996 and 2010. There were 1,050,943 women who didn't have diabetes and 45,384 who had gestational diabetes (GDM). Likewise, 13,278 women had

pregestational diabetes (pre-GDM). The speeds of main perinatal outcomes were analyzed among gatherings and years using Poisson backslide. Likewise, the age-changed speeds of diabetes in pregnancy were determined annually. The incidence of inborn anomalies decreased by 23% anyway the speeds of perinatal passings were by and large unaffected. There was an increased bet of perinatal mortality for pre-GDM women, and the normality of inborn anomalies was higher in women with GDM contrasted with those without diabetes. The repeat of both gestational and pre-gestational diabetes has duplicated in the last fourteen years, and the by and large cost of this condition has been on the ascent. Regardless of the way that diabetes women have seen a decline in the ordinariness of perinatal mortality, the bet of inherent anomalies and perinatal passing is still much higher than it is for nondiabetic women. We truly want to get different things done to diminish these antagonistic effects: which the American Diabetes Association worked with in 2014.

The normal range of hemoglobin A1c in sound pregnant women is 4.5-5.7%, and it rises pointedly in pregnant women who test positive for protein in their urine. Glycated hemoglobin is reduced as HbA1c. Some glucose links to hemoglobin (the oxygen-carrying protein in red platelets) as it circles in the blood isolated. Hemoglobin A1c is the shortening for this combo. Blood glucose levels are comparative with the quantity of hemoglobin A1c made.

Blood glucose levels are raised, leading to more prominent HbA1c levels, when diabetes isn't especially managed. Since RBCs have a half-presence of three to four months, HbA1c values don't differ often. This means that HbA1c levels in the blood are indicative of the normal blood glucose levels over the last a couple of months. More glycated hemoglobin is made when blood glucose levels are high. Transporting oxygen from the circulatory framework to every cell in the body is hemoglobin's fundamental role. The glucose in the stream coats hemoglobin, causing it to become glycated. A more prominent A1c level is the result of the hemoglobin protein attracting more glucose in the blood, which in turn causes the external layer of the hemoglobin protein to reflect higher glucose levels.⁹

“The "leaned toward screening and suggestive 2-step" way to deal with treating gestational diabetes mellitus, as proposed by Diabetes Canada in 2018, has our full help. From week 24 to week 28, all pregnant ladies ought to have a standard non-fasting 50-g glucose challenge screening test (GCT) with plasma glucose (PG) measured one hour after the occasion (III-B). Further testing isn't required on the off chance that the outcome is under 7.8 mmol/L.1.2. The patient is told to do a 75-g oral glucose opposition test with fasting PG (FPG), 1-hour PG, and 2-hour PG like clockwork assuming the GCT is inside the scope of 7.8 to 11.0. When the accompanying circumstances are met or surpassed: I. FPG ≥ 5.3 mmol/L ii. 1-h PG ≥ 10.6

mmol/L iii.2-h PG \geq 9.0 mmol/L, then gestational diabetes mellitus is getting looked at. At the point when the GCT result is 11.1 mmol/L or above, gestational diabetes mellitus is considered. According to Diabetes Canada's 2018 suggestions, the "elective 1-step definite" approach is adequate. As a feature of this arrangement, pregnant ladies between the ages of 24 and 28 weeks ought to go through normalized oral glucose opposition testing utilizing fasting plasma glucose (FPG), 1-hour plasma glucose (PG), and 2-hour PG. The accompanying circumstances should be fulfilled for this to be valid: gestational diabetes mellitus is assessed while the accompanying levels are met: I. FPG \geq 5.1 mmol/L; ii.1-h PG \geq 10.0 mmol/L; and iii.2-h PG \geq 8.5 mmol/L. We recognize that the "enjoyed" and "elective" procedures have different illustrative constraints, which could prompt misconceptions in specific settings. Following the current Diabetes Canada suggestions is, as per the warning board, vital. Consequently, it is firmly supported that each care office pick one of the two strategies and execute system to ensure the predictable and dependable bookkeeping of preliminary results.¹⁸ Ladies who are at high gamble for gestational diabetes mellitus because of a blend of hazard elements ought to have screening or testing during the primary trimester of their pregnancy. Assuming the outcomes are typical during the primary trimester, they ought to go through recurrent testing between weeks 24 and 28. In the event that the gamble of later-beginning gestational

diabetes was clinically thought or disregarded for any reason, a screening or interesting test ought to be thought of.”

Certain drawbacks of the ADA/IADPSG recommendations and cutoffs have been discussed by Seshi et al.¹⁰: With the exception of Bangkok and Hong Kong, the HAPO research mostly included Caucasians. They postulate that ethnic Asian pregnant women may have elevated blood glucose levels due to increased insulin resistance. Additionally, the majority of pregnant women do not fast before their prenatal appointments. Therefore, when requested to return for an OGTT, the dropout rate is very high, particularly in underdeveloped nations where prenatal checkups are few. Due to financial constraints and a shortage of appropriately trained personnel, glycolylated haemoglobin cannot be produced in settings with limited resources. The Diabetes in Pregnancy Study Group India (DIPSI) proposed a "single step" diagnosis process for all patients, often known as universal screening, to address these issues in developing nations. Regardless of the pregnant woman's fasting state or the date of her last meal, she is given an oral 75 g glucose load following the preliminary assessment in the prenatal clinic. Diabetic mellitus is defined as blood glucose levels 140 mg/dl or above after 2 hours. The idea behind this test is that an ordinarily glucose-tolerant woman would have the choice to keep her glucose levels stable in any occasion, when introduced to a high glucose load. The hyperglycemia trip is

substantially more expressed in patients with GDM. Both the Indian government's Ministry of Wellbeing and the World Wellbeing Organization have given their blessing to this one-step process. The DIPSI technique has the following advantages: Fasting isn't needed for this framework, which may be finished on the initial visit to the trained professional and is best when rehashed in the second and third trimesters of pregnancy. Does not significantly influence the woman's regular schedule. Is both a screening as well as an indicative cycle.

“Threshold Values for Diagnosis of Gestational Diabetes with 75-g OGTT			
Plasma Glucose	Glucose Concentration Threshold ^a		Above Threshold (%)
	Mmol/L	Mg/dl	Cumulative
Fasting	5.1	92	8.3
1 -hr OGTT	10.0	180	14.0
2 -hr OGTT	8.5	153	16.1 ^b
<p>To diagnose gestational diabetes, one or more of these 75-g OGTT levels must be met or surpassed.</p> <p>Also, out of the first cohort, 1.7% were not blindfolded due to fasting plasma glucose levels greater than 5.8 mmol/L (105 mg/dL) or 2-hour oral glucose tolerance test results greater than 11.1 mmol/L (200 mg/dL), for a total of 17.8%.</p>			

Oral glucose tolerance test means OGTT.”
--

From the Williams text book of obstetrics

The Overall Relationship of Diabetes and Pregnancy Review Get-togethers (IADPSG) and the American Diabetes Affiliation propose the 75 g, 2-hour oral glucose opposition test as an expressive instrument. The limits of this OGTT are according to the accompanying:

Free base metal concentration: more than - 92 mg/dL something like 180 mg/dL following 1 hour following 2 hours, more than 153 mg/dL Diabetes mellitus finding with OGTT (WHO, FIGO, IADPSG)

The American Diabetes Affiliation prescribes using plasma glucose estimations to examine diabetes. A1C, 2-hour plasma glucose (2-h PG) following a 75-g oral glucose deterrent test (OGTT), and fasting plasma glucose (FPG) are in like manner fundamental for these standard working frameworks (21). It is extensively believed that A1C, 2-hour PG during 75-g OGTT, and FPG are inside the ordinary reach concerning logical screening. True to form for the two individuals and social events, the disclosure speeds of different screening tests could vary. In like manner, centers around that have shown the reasonableness of medications for principal abhorrence of type 2 diabetes have precluded people with limited hindered fasting glucose (IFG) or weakened glucose adaptability (IGT) related paying little regard to

raised fasting glucose. The Public Glycemic Status Program or the Diabetes Control and Complexities Primer (DCCT) reference measure should be advised for driving A1C tests (ngsp.org). While giving treatment, In both CLIA-oversaw and CLIA-conceded settings, an A1C test that has been NGSP confirmed and endorsed by the U.S. Food and Medicine Association (FDA) may be used to screen glycemic control in diabetics. All A1C tests may simply be coordinated by labs or various workplaces that have gotten FDA underwriting for clinical benefits use, have gone through examination by CLIA, and meet CLIA's quality standards. Staff people are supposed to fulfill explicit requirements and to take part in an embraced capacity testing program something like multiple times every year (29-32). A1C partakes in various high grounds over FPG and OGTT, including being more profitable (no fasting is required), having more preanalytical strength, and having less regular interferences due to push, dietary changes, or disease. The use of A1C testing partakes in the two advantages and downsides. According to one point of view, it is more excessive and less accessible in non-modern nations. On the other hand, there isn't for the most part an optimal association among's A1C and typical glucose in unambiguous individuals. According to measurements from the Public Prosperity and Sustenance Assessment Outline (NHANES) (33), only 30% of the diabetes patients found out and out using A1C, FPG, or 2-h PG are investigated using the A1C test. The

definite edge for A1C is 6.5%, or 48 mmol/mol. In spite of these disclaimers, A1C was incorporated the 2009 definite norms by the Overall Master Board to augment screening rates (39). A1C should not be used to break down diabetes without besides considering various conditions that could influence hemoglobin glycation close to glycemia, similar to hemodialysis or pregnancy, since it is an indirect indication of typical blood glucose levels.

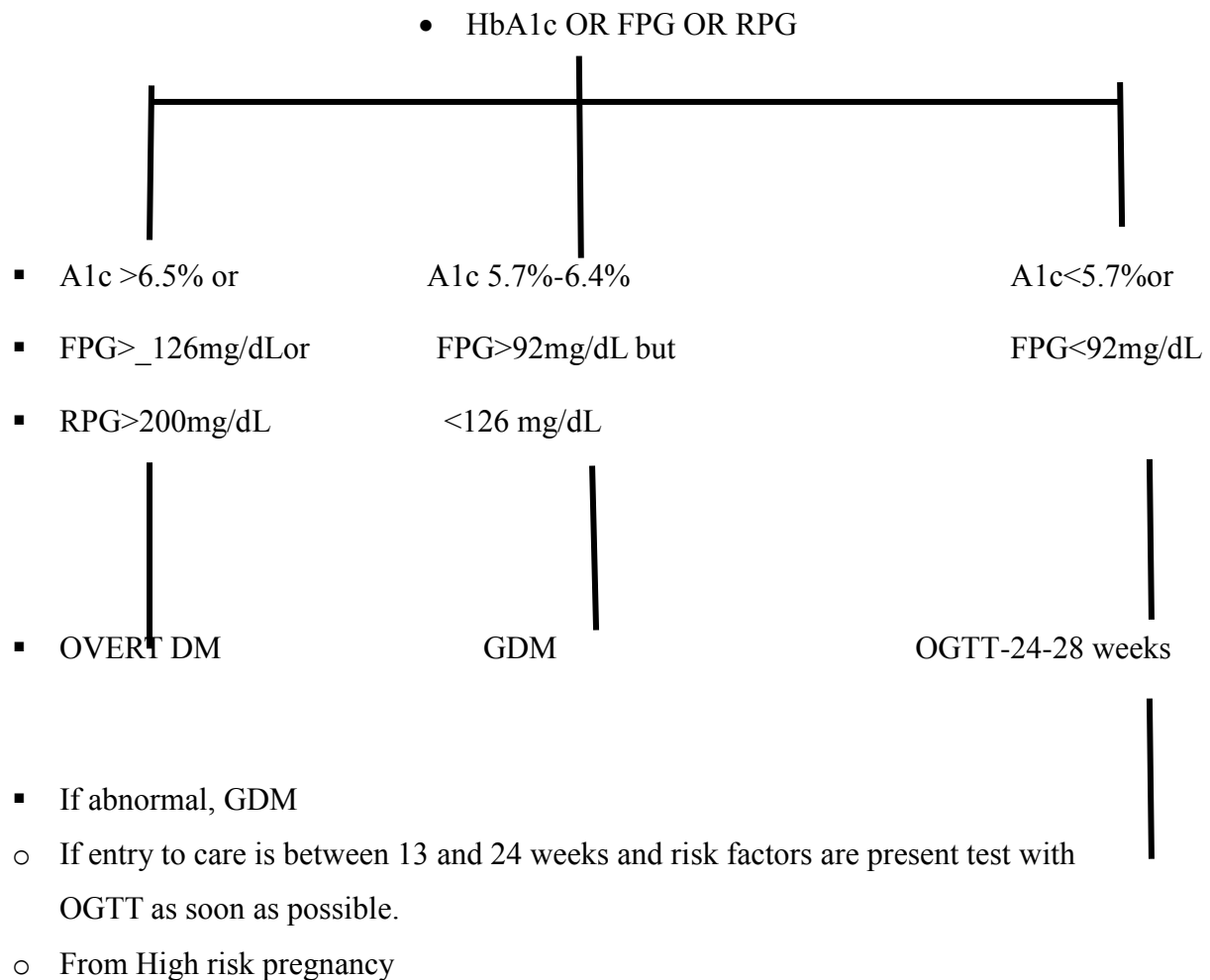
“OGTT criteria for diagnosis of Diabetes Mellitus (WHO, FIGO, IADPSG)			
75 G – OGTT	Normal	Gestational Diabetes Mellitus (GDM)	Diabetes Mellitus In Pregnancy (DIP)
FBS	< 92 mg/dl (5.1 mmol/l)	92 -125 mg/dl (5.1 – 6.9 mmol/l)	>=126mg/d (7.0 mmol/l) ¹
1 hr Post	< 180 mg/dl (10 mmol/l)	>= 10 mmol/l	
2 hr Post	< 153 mg/dl (8.5 mmol/l)	153 to 199 (8.5 - 11.0 mmol/l) mg/dl	>= 200 mg/dl (11.0 mmol/l)”

In 2011, the ADA From Dewhurst’s text book of obstetrics and gynaecology¹ and the IADPSG collaborated to propose the following methodology for

diabetes in pregnancy: 1

Fetal risks generally include Issues with fetal oxygenation, including macrosomia, advancement hindrance, and deformations; Post pregnancy substance imbalances; Fetal oxygenation challenges, including respiratory wretchedness problem, cardiovascular breakdown, and unforeseen fetal downfall; and Long stretch results. A wide range of gestational diabetes address the bet of strange fetal development. The most predominant anomaly is macrosomia, which is defined as a fetal weight that is more than the 90th percentile for that gestational age or a normal fetal heap of 4000 grams or higher. Maternal hyperglycemia makes an overabundance of glucose transferred across the placenta, which in turn induces hyperinsulinemia in the developing kid, a condition known as macrosomia (Pedersen speculation). Fetal pancreatic islet cells begin to encourage the ability to release insulin in light of hyperglycemia around the 20-week normal for incubation, at which point this impact becomes obvious. Moreover, hyperinsulinemia may cause fetal hepatomegaly, splenomegaly, and cardiomegaly. Right when the abdominal edge is higher than the 95th centile, the positive farsighted incentive for detecting macrosomia is above 90%. According to ACOG, the bet of macrosomia could arrive at 20% if gestational diabetes isn't perceived or treated.

HbA1c levels in GDM

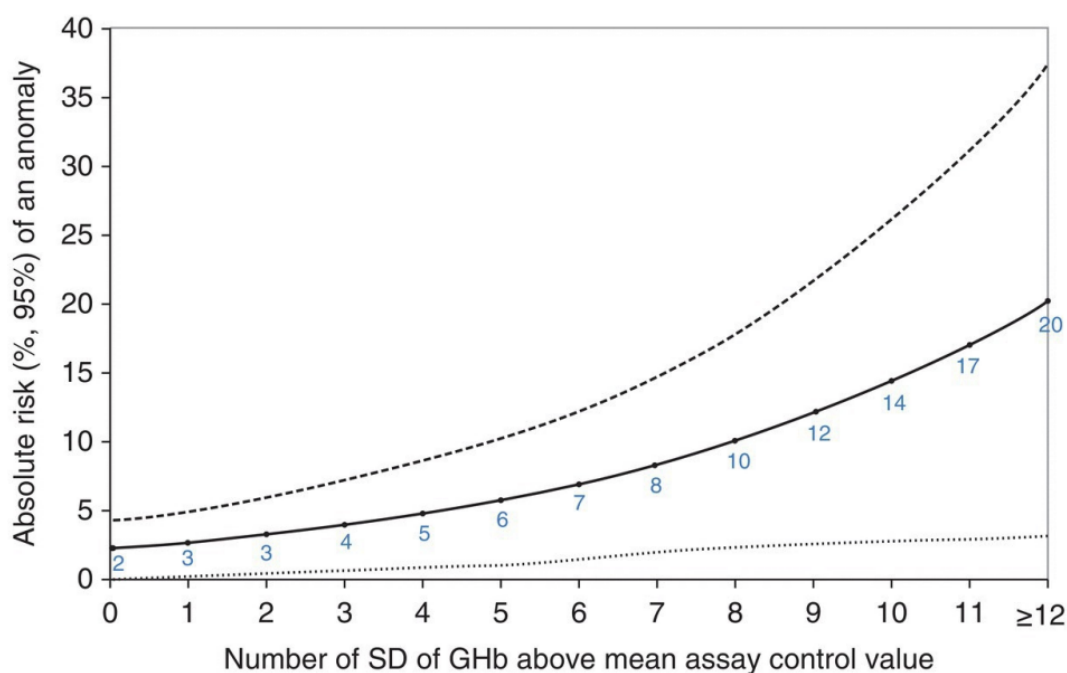
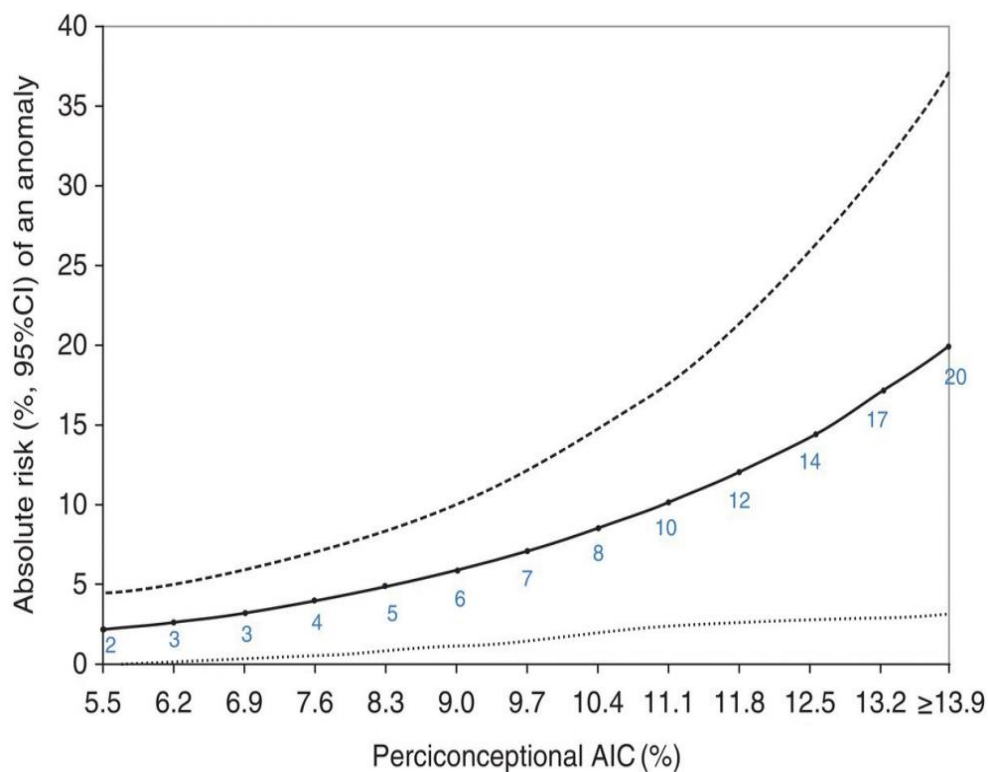


Fernando arias text book of obstetrics and gynecology

“Fifth International Workshop-Conference: Metabolic Assessments Recommended After Pregnancy with Gestational Diabetes		
Time	Test	Purpose
Postdelivery (1 – 3 d)	Fasting or random plasma glucose	Detect persistent, overt diabetes
Early postpartum (6-12 wk)	75-g, 2 hr OGTT	Postpartum classification of glucose metabolism
1-yr postpartum	75-g, 2-hr OGTT	Assess glucose metabolism
Annually	Fasting plasma glucose	Assess glucose metabolism
Triennially	75-g, 2-hr OGTT	Assess glucose metabolism
Prepregnancy	75-g, 2-hr OGTT	Classify glucose metabolism
Classification of the American Diabetes Association (2013)		
Normal Values	Impaired Fasting Glucose or Impaired Glucose Tolerance	Diabetes Mellitus
Fasting < 100 mg/dl	100-125 mg/dl	>=126 mg/dl
2 hr < 140 mg/dl	2 hr >=140-199 mg/dl	2 hr >= 200 mg/dl
Haemoglobin A1c	5.7-6.4%	>=6.5%

< 5.7%		
OGTT = oral glucose tolerance test.		
Data from American Diabetes Association, 2013, 2017a; Metzger, 2007.”		

- From text book of Williams obstetrics 24 th edition



The 20th week of pregnancy marks the beginning of a rise in the amount of insulin that is required, and this increase continues for the whole of the pregnancy. Because of this, it is possible that women who have diabetes that is under control will not be needed to raise the total amount of insulin that they take until the twenty-first week of their pregnancy. It is possible that the amount of insulin that is required during the second trimester of pregnancy will rise by a factor of three in comparison to the first trimester. Women are able to self-adjust their insulin levels in response to HBGM if they have the knowledge and the confidence to do so. This is something that they are capable of accomplishing. In order to prevent hypoglycemia from occurring, you should cut the quantity of glucose that you ingest throughout the night by fifty percent. Insulin resistance The majority of patients have insulin resistance, which makes oral hypoglycemic medications inefficient in managing blood sugar levels. This is because insulin resistance is a common occurrence. It is possible that a woman who has type 2 diabetes may need 300 units of insulin on a daily basis after the completion of her pregnancy. The possibility exists that metformin offers the additional advantage of lowering the total amount of insulin that is necessary to be administered. This is due to the fact that it has the capability to have an impact on insulin-

sensitive tissues as well as the liver's capacity to produce glucose. In order to reduce the possibility of hypoglycemia developing, individuals should be informed on the best times to snack and eat, as well as appropriate exercise routines. Additionally, family members should be trained on how to administer glucagon. Always carrying a card that has information on their medicine and instructions on how to react in the case of an emergency involving hypoglycemia is something that women who use insulin should do with them at all times. If a person is undergoing a hypoglycemic episode, it is strongly recommended that they refrain from driving. This is a piece of particular advice for drivers. It makes no difference whether they are feeling any signs of disease or not; this is always the best course of action. How can people improve their ability to tolerate glucose while they are in the second stage of their life when they are mothers? During the beginning of the second trimester of pregnancy, the levels of glucose in the blood decrease while the mother is fasting, while the levels of glucose in the blood after eating increase. This occurs in pregnancies that are not impacted by diabetes. Because of increased levels of placental hormones and circulating maternal fatty acids, insulin resistance starts to develop during the middle of the second trimester of pregnancy. This is because of the combination of these two factors. The 20th week of pregnancy marks the beginning of a rise in the amount of insulin that is required, and this increase continues for the whole

of the pregnancy. Because of this, it is possible that women who have diabetes that is under control will not be needed to raise the total amount of insulin that they take until the twenty-first week of their pregnancy. It is possible that the amount of insulin that a woman has to take throughout her pregnancy will increase by a factor of three from the beginning of the pregnancy until the end of the pregnancy. Ten Women are able to self-adjust their insulin levels in response to HBGM if they have the knowledge and the guts to do so.

It is possible that women with type 2 diabetes who experience the beginning of insulin resistance during the second trimester of their pregnancy will be needed to take oral hypoglycemic dosages that are more than 300 units per day. Since insulin-agents often fail to provide appropriate glycemic control and metformin modulates hepatic glucose synthesis, it is likely that insulin dosages might be lowered by using metformin. This is on the grounds that metformin impacts the creation of glucose in the liver. The improvement of tissues that are delicate to insulin is one of the distinctive qualities of a pregnancy that is viewed as better. An assessment into the many sorts of inborn irregularities that might be available A widely inclusive ultrasound that is completed between the eighteenth and twentieth seven day stretch of pregnancy can reveal any serious innate irregularities that might be available. With the utilization of this specific contraption, you should have

the option to look at the ventricle surge courses notwithstanding the four offices of the heart. Diabetes during pregnancy is related with a fundamentally expanded possibility of creating toxemia and different sorts of hypertension. Pregnant ladies who have diabetes ought to be observed for both the counteraction and checking of unexpected issues all through pregnancy. The uterine Doppler waveform examination at 20 weeks might be of help in identifying which ladies are most in danger of fostering the problem. This is notwithstanding the pee protein tests and the normal circulatory strain readings that are acted in the facility. After the finishing of the second trimester of pregnancy, it is typical to do ultrasounds once like clockwork (or on a more regular basis on the off chance that it is passed judgment on vital) to screen the improvement of the creating baby. This is done in order to monitor the progress of the baby's growth. By comparing the percentile analysis of the baby's abdominal circumference at 28 weeks to scans performed later in the pregnancy, it is possible to assess if the baby is developing at a normal speed or an unusually rapid pace. This may be done by comparing the two sets of data. Due to the fact that the presence of diabetes is linked to an increased risk of polyhydramnios, it is of the utmost importance to record measurements of fluid volume in a certain order during the course of a diabetic pregnancy. The situation as it now stands The most effective therapy for blood sugar As the third trimester develops, both insulin

resistance and insulin needs continue to gradually grow during the course of the pregnancy. Insulin resistance makes it simpler to control blood sugar levels and provides women with protection from critically low glucose levels in their blood. Insulin resistance is a kind of diabetes. The appearance of reduced glucose levels in the mother's blood during the third trimester of pregnancy may be a sign of severe hyperinsulinemia in the foetus. It is plausible that this disease might be brought on by a larger glucose gradient across the placenta or placental dysfunction³⁷. If insulin needs were to decrease, it is feasible that it would be reasonable to contemplate becoming pregnant sooner. Assuming that glucocorticoid hormones are necessary for lung expansion, it is possible that a person's daily insulin need may rise by a factor of two over the course of the subsequent three days. Consider the following scenario as an illustration: a patient receives injections of beclometasone once every twelve hours. These injections are given into the veins of the patient. This woman has to be hospitalised and given an intravenous insulin infusion of seventy-two hours, which is often referred to as an insulin sliding scale. The infusion will be administered over the course of twenty-four hours. She should also be instructed to take a higher dosage of insulin as an alternative strategy. Identifying hyperinsulinemia-induced fast foetal growth may be accomplished by an assessment of the developing foetus. This can be accomplished by comparing the percentile of the

abdominal circumference to the biparietal diameter or the head circumference. In order to do this, multiple ultrasound exams are required. The mother's internal organs, like as her heart and liver, are exposed to increased strain as a consequence of the overproduction of insulin that occurs during pregnancy in the growing infant. A further symptom is that she is experiencing a heavier sensation than she typically would. When it comes to the actual definition of the word "macrosomia," which is used to describe to babies who are delivered to mothers who are dealing with diabetes, there is a lack of agreement among health professionals. The absolute birthweight may be less than four kilogrammes, four and a half kilogrammes, or five kilogrammes, according to some standards; nevertheless, the percentile birthweight may be more than ninety percent, ninety-five percent, or ninety-seven point five percent of the total birthweights. In the process of trying to quantify birthweight, absolute weight is not the right measurement to use since it varies based on variables such as the gestational age of the mother, the gender of the mother, the ethnicity of the parents, the height and weight of the parents, and the weight of the mother. In the context of clinical practice, the pattern of foetal development is more crucial than either the absolute or percentile birthweight. This is because the pattern of foetal development is more essential. The birthweights of a significant number of newborns who are delivered to moms who have type 2 diabetes or

gestational diabetes mellitus type 17 are much higher than average. The majority of this may be attributed to the effects of maternal obesity, which encompasses both being overweight and being obese. By using serial ultrasonography, it is feasible to discover foetal growth limitation, asymmetrical growth restriction, or a little increase in AC centile development in compared to head circumference. All of these conditions may be identified without any difficulty. There is a possibility that women who have type 1 diabetes and also have renal impairment, vascular disease, or hypertension would display this pattern of development, which is an indication of uteroplacental insufficiency. There is a possibility that women who have hypertension will also exhibit this pattern of development. In the event that it is found, it is possible that it will be required to make preparations for its delivery at a later time. Which time and where will the delivery take place, and what time is it? When it comes to diabetes education, it is essential that initiatives begin as early as possible in order to educate pregnant women with knowledge about the risks that are involved with carrying a pregnancy to term. During the time that they are pregnant, it is essential for women to have a say in the choices that are made about their birth plans. Details such as the method of delivery and the time should not be considered until after the 36-week growth scan, when both the mother and the baby will have more information at their disposal. Examples of such

details include the technique of delivery and the timing. A woman who has diabetes is predicted to have a probability of having a late surprise stillbirth that is roughly four times higher than a woman who does not have any form of diabetes. This is in contrast to a woman who does not have any form of diabetes. It is generally agreed upon among professionals in the medical field that nursing mothers who are dependent on insulin should make every effort to deliver their kids between the 38th and 39th week of their pregnancy. Inducing labour for a woman who is giving birth for the first time and has type 1 diabetes may be a procedure that is both time-consuming and fruitless. It is safe to anticipate that caesarean sections account for around fifty percent of all deliveries in women who have type 1 diabetes level six. This is due to the fact that a significant proportion of caesarean sections are performed following unsuccessful efforts at induction of labour. On the other hand, women who have type 2 diabetes have a propensity to have a bigger number of pregnancies and a higher success rate with procedures that induce early birth. This is because type 2 diabetes is a more advanced form of diabetes. It is conceivable for women whose gestational diabetes is totally managed by diet or metformin to reach a safe gestational age (40 weeks) and expect that labour will begin without the need for medical intervention. One way to do this is by ensuring that the diabetes is completely controlled. This is based on the assumption that their pregnancies are free of any

complications. There is a correlation between having a larger birth weight and an increased likelihood of experiencing birth defects and complications. In compared to moms who do not have diabetes mellitus, those who do have the disease have a greater likelihood of suffering shoulder dystocia. However, this is not the case for all mothers. This specific population is at a danger that is five times higher than it was before. In light of the fact that this ratio is high, it is possible that this may be better understood in light of the fact that pregnancies in which the mother has diabetes are marked by an increased chest-to-head and shoulder-to-head ratio. Approximately three percent of newborns born between four and four and a half kilogrammes are affected by shoulder dystocia, and ten to fourteen percent of infants born over four and a half kilogrammes are affected by this condition. If, on the other hand, the plan depended just on the anticipated birthweight as determined by ultrasonography (four thousand or four thousand and five hundred grammes), then the number of caesarean sections would be far higher than what is considered acceptable. The diabetic lady who had a scan that indicated a big foetus should have an obstetric decision made for her, taking into consideration her choices, the medical history of her family, and her own dimensions.²⁶ Diabetes care throughout pregnancy Maintaining glycaemic targets, which include a fasting blood glucose level that is below 5.3 mmol/L and a 1 hour postprandial glucose level that is below 7.8

mmol/L, is crucial for the treatment of gestational diabetes throughout the duration of pregnancy, just as it is for other diabetic women. 13. Some women find that making modifications to their diet and the amount of physical activity they participate in gives adequate respite from the symptoms of type 2 diabetes, despite the fact that insulin is often necessary for the treatment of the condition. When it comes to oral hypoglycemic drugs, the exact aim of these prescriptions is still a point of contention in a number of countries all over the globe. Even though many people continue to assume that sulphonyl urea drugs are less effective and flexible than insulin, it would seem that the most current sulphonyl urea medications are safe to use beyond 15 weeks of gestation. This is despite the fact that insulin is still the most often used treatment. The use of oral hypoglycemic medicines, on the other hand, offers a potentially helpful option in regions where insulin is in short supply. Women who have gestational diabetes but have not been diagnosed with diabetes often have a return to normal glucose tolerance following birth. Throughout the span of the following twenty years, type 2 diabetes will without a doubt impact most of the population.^{43.1} The presence of the diabetes quality during pregnancy is perhaps of the main component that could add to the improvement of type 2 diabetes in later life's stages. As a component of the post pregnancy diabetes test, it is suggested that all ladies who have been determined to have

gestational diabetes mellitus have their hemoglobin A1c or fasting glucose tried a month and a half after the conveyance of their kid, and afterward again yearly after that. a 44 A change to one's lifestyle that empowers actual activity and assists with staying away from It is conceivable that your possibility having diabetes might diminish in the event that you consume a less than stellar eating routine during the accompanying four to five years. Ladies who have recently conceived an offspring and have been determined to have gestational diabetes mellitus (GDM) ought to be prompted that they are obliged to get a diabetes test once over time, as well as getting fundamental way of life direction. It is suggested that you conceive an offspring between weeks 38 and 39 on the off chance that you are a woman who needs insulin to get past the course of work and conveyance. Ladies who can monitor their glucose levels all through their pregnancy by embracing a fair eating regimen or taking metformin have a risk of having a sound pregnancy up to week 40. This is plausible. In the prompt fallout of conceiving an offspring, a pregnant lady who has pre-gestational diabetes ought to lessen the amount of insulin she takes to a level that is somewhere in the range of 25 and 30 percent of the aggregate sum she took during her pregnancy. A few ladies who have type 2 diabetes might have the option to stop taking insulin treatment after the introduction of their youngster. This is a beneficial result for these ladies. Moreover, it is practical for a critical

number of ladies who are determined to have gestational diabetes to cease the utilization of every single diabetic prescription. With regards to the use of metformin, insulin, or glibenclamide, there is no requirement for any nursing mother to be stressed over the circumstance. An assessment of the hemoglobin A1c, which is likewise frequently alluded to as the fasting blood glucose level, is something that diabetic ladies ought to have done.

This is in reference to Mac MC, Fleming KM, Bailey JA, as well as their partners and teammates. An examination was done in Britain, Ribs, and Northern Ireland to decide the recurrence of perinatal mortality and inherent mutations among babies who were brought into the world to moms who had either type 1 or type 2 diabetes. After the mother had conceived an offspring, the review was completed a month and a half later, and afterward it was rehashed year after that. The prevalence of diabetes during pregnancy is on the rise, and as a result, there is a pressing need to lay a greater emphasis on the dangers that uncontrolled diabetes poses to the mother, the developing infant, and the subsequent generation as a whole. In this discussion, a number of significant features of the treatment of diabetic pregnancies have been discussed. These aspects include screening for gestational diabetes as well as other component that is vital. The use of a multidisciplinary team that is headed by a consultant and the execution of national therapeutic recommendations that are based on evidence are both crucial in order to

maximise the efficacy of glycemic therapy and limit the possibility of complications that are connected with maternal diabetes.

In terms of GDM, which disproportionately affects pregnant and nursing women, India is quickly becoming known as the diabetic capital of the globe.⁴⁷ According to the World Health Organization's projections, the number of people with gestational diabetes would increase from 135 million in 1995 to 300 million in 2025, a total increase of more than 120 percent.

Rohini HN et al.⁴⁶ reports that researchers in many Indian states looked at GDM.

The following table summarises data from many GDM investigations carried out in India.

“List of various studies on Gestational diabetes Mellitus in India

Place	Year of study	Sample size	Test done	Prevalence	Author
Rural Assam	July 2019-September 2019.	1212	Oral GTT	16.7%	Subrata Chanda <i>et al.</i>
Kolkata	Aug 2016-July 2018	416	IADPSG DIPSI	37.3% 31.3%	Lipika Das Mukhopadhyay <i>et al.</i>
Lucknow, Uttar Pradesh	Aug 2016-Sept 2017	162	8 h of overnight fasting and 75 g anhydrous glucose	22.64%	Arpit Gupta
Telangana	January	32 428	elevated	5.4%	Goutham

	2015- December 2016		random blood glucose		Swaminathan
Kerala	January 2015 and December 2016	32 428	elevated random blood glucose	4.5%	Goutham Swaminathan
West Bengal	January 2015 and December 2016	32 428	elevated random blood glucose	2.3%	Goutham Swaminathan
Uttarkhand	January 2015 and December 2016	32 428	elevated random blood glucose	1.2%	Goutham Swaminathan
Bihar (east),	January 2015 and December 2016	32 428	elevated random blood glucose	1.2%	Goutham Swaminathan
Madhya Pradesh (central)	January 2015 and December 2016	32 428	elevated random blood glucose	1.2%	Goutham Swaminathan
Uttar Pradesh (central)	January 2015 and December 2016	32 428	elevated random blood glucose	1.2%	Goutham Swaminathan
Meghalaya	January 2015 and December 2016	32 428	elevated random blood glucose	0.23%	Goutham Swaminathan
Delhi	December 2015– October 2016	100	75 g oral glucose test	14%	Samreen Siddiqu

Bhilai	(December 2015–October 2016)	65	1 hour 75 gm glucose	10.7%	Samreen Siddiqui
Muzaffarpur	(December 2015–October 2016)	65	Fasting blood glucose	3.07%	Samreen Siddiqui
Pune, Maharashtra	September 2012 to June 2014	989	2 h post 75 g glucose	9.5%	Anjali A. Bhatt
Lucknow	June 2012 to July 2013	332	2 hrs 75 g OGTT	41.9%	V Gopalakrishna
Punjab	August 2009–December 2012.	5100	WHO 2013 criteria WHO 1999 criteria	35% 9%	Geeti P Arora <i>et al</i>
Kashmir	April 2011 - March 2012.	306	75 g OGTT	7.8%	Malik Waseem Raja <i>et al.</i>
Rohtak	June 2009 to January 2011	607	75 g 2 h OGTT	7.1%	Rajesh Rajput
Chennai	February–December 2001	1251	2 h 75 g glucose ≥ 140	17.7%	V Seshiah <i>et al.</i>
Bangalore	1991	302	3 hrs OGTT	6.3%	Jaya Narendra <i>et al.</i> ”

They arrived at the resolution in their examination that Indian dietary patterns might have been affected by fascinating impacts, and that our living

style and way of life appropriations have likewise been affected by capricious movements.

AResearchers Valadan M, Bahramnezhad Z, et al.⁴⁷ looked investigated how HbA1c levels in the principal trimester might assist with distinguishing gestational diabetes in its beginning phases. From 2018 to 2020, specialists at a tertiary college emergency clinic followed 700 pregnant ladies who visited the perinatology facility. During the main trimester of pregnancy, all ladies were tried for hemoglobin A1c and fasting blood glucose (FBG). Then, a 100-g oral glucose opposition test (OGTT), the best quality level, was used for a GDM screening test inside 24-28 weeks of pregnancy. We used the standards set out by the American Diabetes Connection (ADA) to investigate GDM. Using the beneficiary working brand name (ROC) twist, we chose the responsiveness, disposition, positive farsighted worth (PPV), and negative perceptive worth (NPV) of HbA1c and FBG. Their assessment suggests that first-trimester HbA1c is neither fragile or unequivocal enough to substitute OGTT in diagnosing GDM.

As indicated by Berger, Gagnon, and partners⁴⁸, rule no. 393 addresses diabetes during pregnancy. The reason for this rule is to give the most exceptional data on the analysis and the executives of gestational diabetes. Both present moment and long haul impacts were considered for the moms, including toxemia, diabetes, and other cardiovascular issues; for the children,

these included intrinsic deformities, stillbirth, macrosomia, birth injury, hypoglycemia, and adverse consequences. We used the principles laid forward in the Canadian Team on Preventive Medical care Report to decide how great the proof was. That was all there was to it for the short comments. Diabetes issues during pregnancy are generally brought about by hyperglycemia and the laid out metabolic milieu. In the event that a pregnant lady has diabetes or some other ongoing ailment, she ought to take drug to control her glucose and different side effects. It is more gainful for both the mother and the child or infant when a multidisciplinary group cooperates to direct the mother's glucose levels and screen the embryo appropriately in pre-birth and post pregnancy diabetes mellitus pregnancies (II-2). Ladies who had pre-gestational diabetes mellitus were bound to have a stillbirth before 40 weeks of incubation, as per review concentrates on contrasting them with the overall obstetrical populace. Results from late enormous scope accomplice and reproduction concentrates on pregnant ladies with gestational diabetes mellitus show that the gamble of stillbirth increments between weeks 36 and 39 of incubation.⁴⁸

Treatment for diabetes mellitus should aim at achieving and maintaining euglycemia (II-2B) in women, whether they are pregnant or not. It is recommended to begin a baseline screening at 28 weeks for individuals with pre- or gestational diabetes mellitus. To find out how maternal glycemic

management affects amniotic fluid volume and foetal development rate (II-2B), serial assessments of foetal growth should be done every three to four weeks after that. Women who have type 1 or type 2 gestational diabetes mellitus, or the two kinds, ought to begin pre-birth checking week after week at 36 weeks. Ladies endeavoring to control their gestational diabetes mellitus with dietary alterations ought to start observing their fetal status week by week at 36 weeks into their pregnancies. Soon, there are a couple of perceived ways of assessing the fetal wellbeing. A few instances of these incorporate the non-stress test and the biophysical profile, or they might be utilized related to each other (1III-A).⁴⁹ We suggest starting fetal wellbeing observing prior or potentially on a more regular basis in the event that the mother has any co-morbidities including heftiness, ill-advised glycemic the board, being large for gestational age (>90%), a background marked by stillbirth, hypertension, or being little for gestational age (<10%). In the event that fetal development limitation (II-2A) is being thought of, a Doppler investigation of the umbilical and fetal center cerebral veins could be useful.⁵⁰ Acknowledgment of pregnant ladies with gestational diabetes mellitus or pre-gestational diabetes mellitus ought to happen somewhere in the range of 38 and 40 weeks of growth (II2-B), contingent upon their glycemic control and other co-dismalness contemplations.⁵¹

Ignoring how there is constant discussion, it has been suggested that moms

with moderate hyperglycemia, as opposed to diabetes mellitus, may have more critical confusions during pregnancy (HAPO Study Wonderful Examination Get-together, 29). The move in peril happened at no especially glaring spots. Optional results comparatively shown essential affiliations, yet less significantly. The assessment found that higher string blood serum C-peptide levels and birth weight were reliably connected with moms whose glucose levels were constantly under the logical edge for diabetes. Issues during pregnancy will without a doubt happen in moms with glucose radicalism levels lower than type 2 diabetics, as per the Hyperglycemia and Restricting Pregnancy Result (HAPO) research.

In expansion to the blood tests got for the arbitrary plasma glucose level estimation, the oral glucose resilience tests' fasting and 2-hour examples were likewise surveyed by the middle's labs. Patients whose irregular plasma glucose levels were 160 mg/dL or higher (8.9 mmol/L or higher), whose 2-hour plasma glucose levels were symptomatic of diabetes (>200 mg/dL, 11.1 mmol/L), or whose levels were under 45 mg/dL, 2.5 mmol/L or higher were not dazed due to moral and security concerns. The main individuals who realized the ladies' or alternately carers' glucose levels in the HAPO research were the lab laborers. Subsequently, we restricted support to ladies whose information had been dazed and who had not had any further glucose testing methods after the HAPO trial. In request to gauge plasma glucose and serum

C-peptide levels, blood tests were gathered from the string after conveyance. Immunoassays were performed on the examples at the focal research facility to decide levels of serum C peptide and plasma glucose. To survey fetal β -cell capability, we used the serum C-peptide level in rope blood as opposed to insulin level for three reasons: first, hemolysis is known to improve insulin debasement, which doesn't affect C-peptide level; second, the discharge of insulin and C-peptide is equivalent; and third, we meant to forestall predisposition. Hemolysis is identified in around 15% of rope blood tests when serum or plasma are isolated.

The standard practice at each centre was used to establish prenatal care, birth time, and neonatal care. No facility regularly conducted caesarean sections at a predetermined gestational age or caused preterm deliveries at will. Information on the pregnancy, birth, postpartum, and infant periods was culled from medical records.

The four significant results were clinical infant hypoglycemia, essential cesarean conveyance, fetal hyperinsulinemia (a degree of serum C-peptide in rope blood that is over the 90th percentile), and birth weight over the 90th percentile for gestational age. Unexpected outcomes included hyperbilirubinemia, toxemia, shoulder dystocia, serious baby care needs, and ensuing episodes of shoulder dystocia. Extra information was recovered at the focuses for the situation that a potential extreme unfavorable occasion,

(for example, demise, shoulder dystocia, birth injury, or critical disfigurement) was recognized. Without knowing the mother's glycemic status, individuals from the result audit board of trustees assessed the information to decide if the frequency had occurred. For significant peculiarities, we utilized the ICD-10 groupings, and for pre-birth demise, we used the Australian and New Zealand Predecessor Characterization of pre-birth passing measures. The HAPO information and security observing panel audited data on unfavorable occasions and passings in the wake of being advised about the aftereffects of the oral glucose-resistance test and the arbitrary plasma glucose levels.

OBJECTIVES

To assess the HbA1c in all first trimester pregnant women

To evaluate the relation of HbA1c in early pregnancy (8-12 weeks) and prediction of Gestation diabetes mellitus in second trimester (24-28 wks.) with oral glucose tolerance test OGTT.

MATERIALS &

METHODS

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at a right angle. The vertical line is positioned to the right of the horizontal line, and they intersect near the right edge of the page.

MATERIALS AND METHODS

SOURCE OF DATA:

Pregnant women in first trimester by the 12th week of their gestational age attended OPD in RLJH and research Centre, Tamaka who fulfill inclusion and exclusion criteria.

Study design- a prospective longitudinal study

Study period- From September, 2022 to December, 2023

INCLUSION CRITERIA :

In this research, we included all pregnant women who were at least 18 years old and who were either outpatients or intrapartum patients who would visit our hospital throughout their first trimester, which ends at the 12th week of gestation.

“EXCLUSION CRITERIA:

Pregnant women with Type I or II diabetes mellitus,

Positive family history.

Previous birth of an overweight baby (>4kg)

Obesity

Unexplained perinatal loss.

Polyhydramnios

Recurrent vaginal candidiasis in present pregnancy

Further the patients who withdrew at any phase during the study.”

Sample size n=75

Sample size is estimated based on HbA1c as an indicator for GESTATIONAL DIABETES MELLITUS according to study conducted by A prospective analysis performed in a tertiary care hospital, Considering positive predictive value and absolute error of 10 percent.

The reference article for calculating sample size is BMJ OPEN DIABETES research and care 2020:

Latife Bozkurt et al study was the basis for calculation of sample size. (reference) ^[9]

With 95% confidence levels and a margin of error 7%, population proportion 90%, the sample size was calculated to be 71, rounded up to 75

pregnant ladies who didn't have diabetes mellitus or pre-diabetic circumstances. Test size was determined using online example size adding machine accessible at: [https://www.calculator.net/test size-calculator.html](https://www.calculator.net/test-size-calculator.html) Using equation:

$$n = \frac{z^2 \times p(1-p)}{\epsilon^2}$$

z is the score

ε is the margin error

p is the population proportion

METHODOLOGY

Following the patient's written informed permission and meeting the study's inclusion criteria.

All patients had comprehensive medical histories documented at the beginning of the study.

It was required that all participants undergo testing for first trimester hemoglobin A1c and fasting blood glucose levels. When individuals enrolled between 8 and 12 weeks gestation, blood samples were taken in accordance with a predetermined technique.

HbA1c levels were monitored during the first trimester in 75 pregnant women who visited our RL Jalappa hospital.

Blood samples were obtained from all patients between 8-12 weeks of gestation, and then they underwent a standardised one-step 75 g OGTT between 24-28 weeks of pregnancy prior to screening.

No matter what time of day it is or whether you've eaten recently, the oral glucose challenge test requires you to swallow 50 grammes of anhydrous glucose. It was then tested an hour later, the glucose level in the veins. The sensitivity of the test is defined by the diagnostic cutoff. The test has a 90% sensitivity level, hence 130 mg/dl is considered the top limit.

An overnight fast of 8 to 14 hours was followed by the OGTT. They maintained an unlimited diet and level of physical activity for the preceding

three days. The patient was instructed not to smoke and to remain sitting for the whole exam. The patient was administered a solution that included 100 g of anhydrous glucose powder. Three samples of blood were taken at one-hour intervals for a total of three hours. An abnormality should be present in at least two of the following values. Diabetes Data Group of the United States standards: Fasting 105, 1hr -180, 2hrs -155, 3hrs-140mg/dl. This oral glucose tolerance test (OGTT) makes use of 75 g of anhydrous glucose. There are 82.5 grammes of glucose available for purchase, with 75 grammes being the anhydrous form. This glucose is a glucose monohydrate.

STATISTICAL METHODS

Using counts and percentages, we summarised the categorical variables and compared them using Pearson's χ^2 test.

Mean \pm SD, median, and IQR were used to summarise continuous data, accordingly.

The necessary statistical performance metrics, such as sensitivity, specificity, predictive values, and 95% CIs, are also included.

A significance threshold of 0.05 has been chosen for both sides.

- **Will patients or any other people or animals need to undergo any kind of examination or intervention for this research to be approved?**
 - YES. Blood specimens were collected in all participants. No financial burden imposed on participants

RESULTS

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at the right end of the horizontal line. The vertical line extends both above and below the horizontal line.

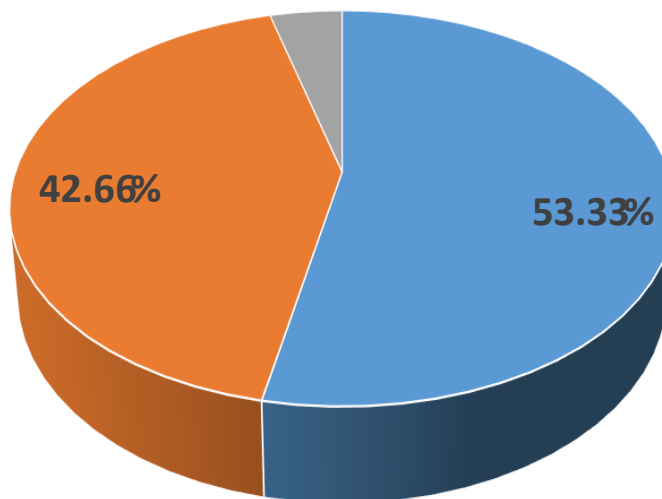
RESULTS

- **Table 1. Age distribution of samples tested Total number of samples tested – 75 1ST Trimester**

Age	number	percentage
< 25	40	53.33 %
25-35	32	42.66 %
> 35	3	4.2 %

- **Graph 1: Age distribution of samples tested**

- ■ < 25 ■ 25-35 ■ > 35

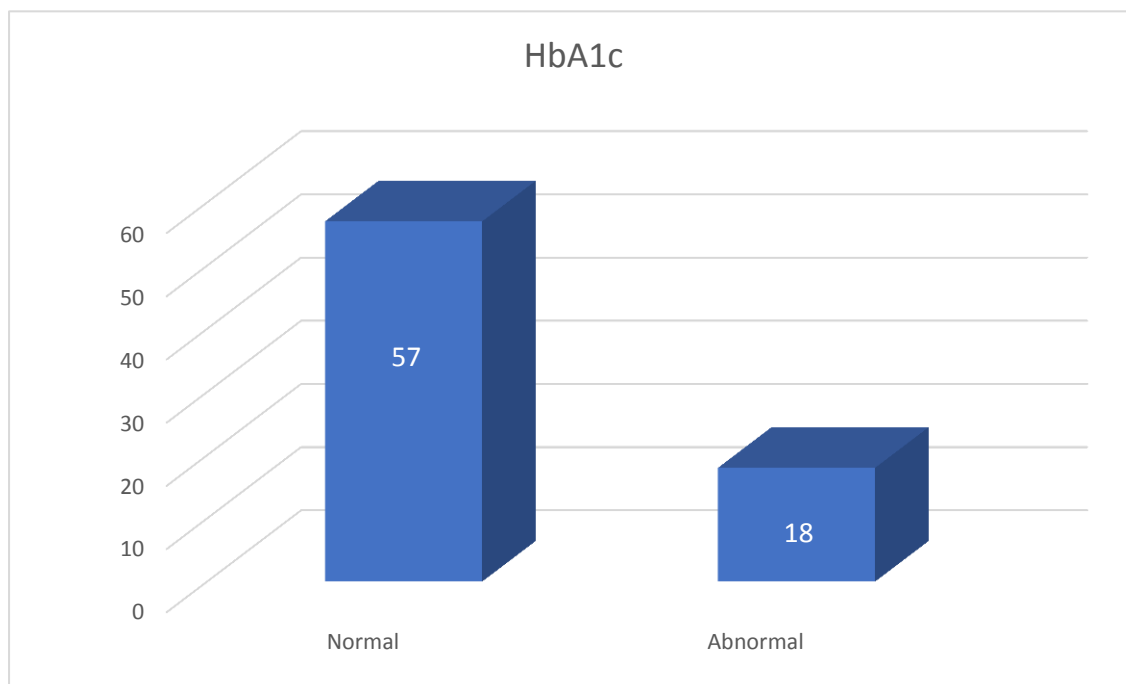


- Total sample tested were 75. Among which <25 years were being 53.33%, 25-35 years being 42.66%. and more than 35 years being 4.2%. More than 35 years elderly group were very less compared to that of <25 years and 25-35 years.

Table 2: Distribution of study population with reference to HbA1c values

HbA1c	number	percentage
Normal	57	76%
Abnormal	18	24%

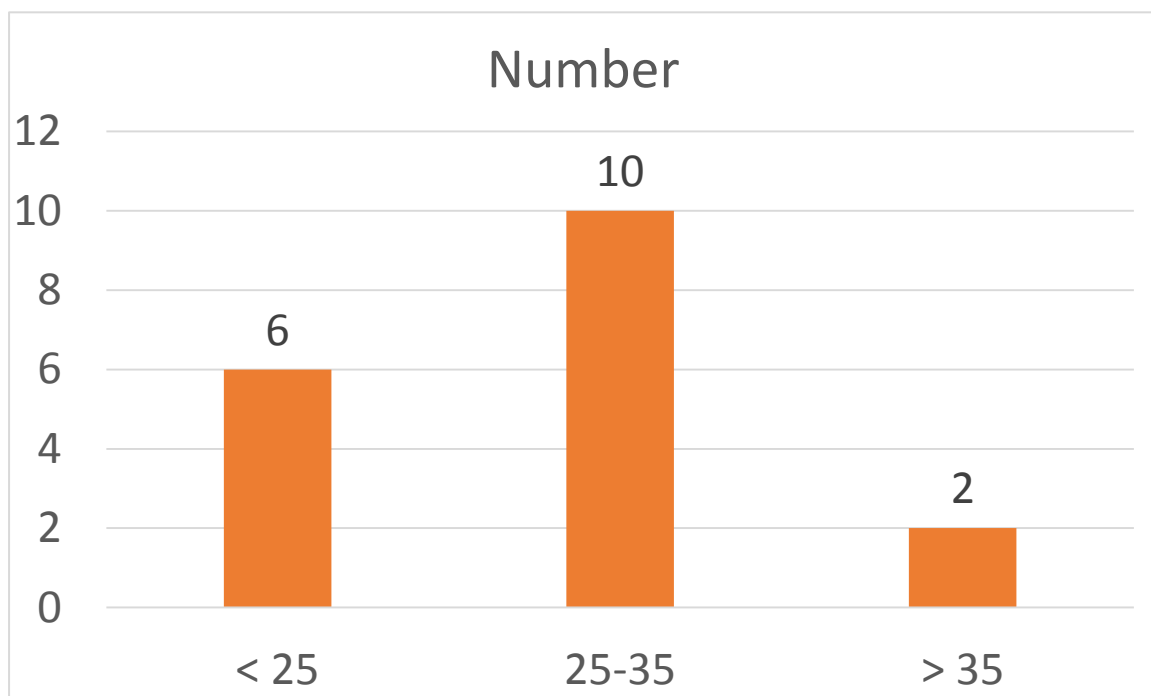
Graph 2: Distribution of patients with reference to HbA1c values



Out of 75 study participant 57 (76%) are with normal HbA1c and 18 (24%) are with abnormal HbA1c.

Tabel 3: Age distribution of abnormal HbA1c

Age	Number	Percentage
< 25	6	8.57%
25-35	10	14.28%
> 35	2	2.8%



Graph 3: Age distribution of abnormal HbA1c

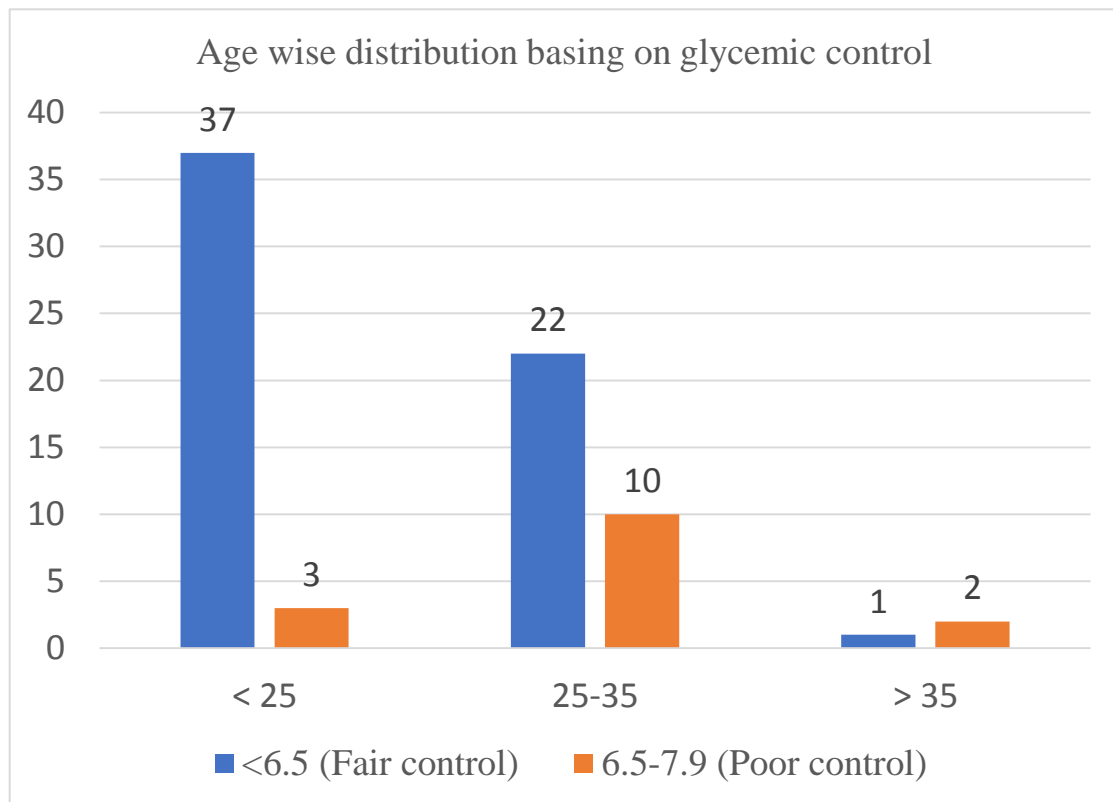
Table 4: Age wise distribution of sample with reference to glycemic control.

Age group	HbA1C group		
	<6.5 Fair control n (%)	6.5-7.9 Poor control n (%)	Total n (%)
< 25	37(92.5)	3(7.5)	40(100.0)
25-35	22(68.8)	10(31.3)	32(100.0)
> 35	1(33.3)	2(66.7)	3(100.0)
Total	60(80.0)	15(20.0)	75(100.0)

In the present study out of 75 pregnant women who were enrolled in first trimester 18 women were found to the with abnormal HbA1c that is above 5.7% among which 6 (8.57%) members were below 25 years age 10 (14.28%) were between 25-35 years and 2 (2.8%) were more than 35 years out of 32 (42.66%) members of 25–35-year age group 10 were found to be with abnormal HbA1c which is statistically significant.

X² Value- 10.52, P value 0.005*-significant

Graph 4: Age wise distribution of sample with reference to glycemic control.

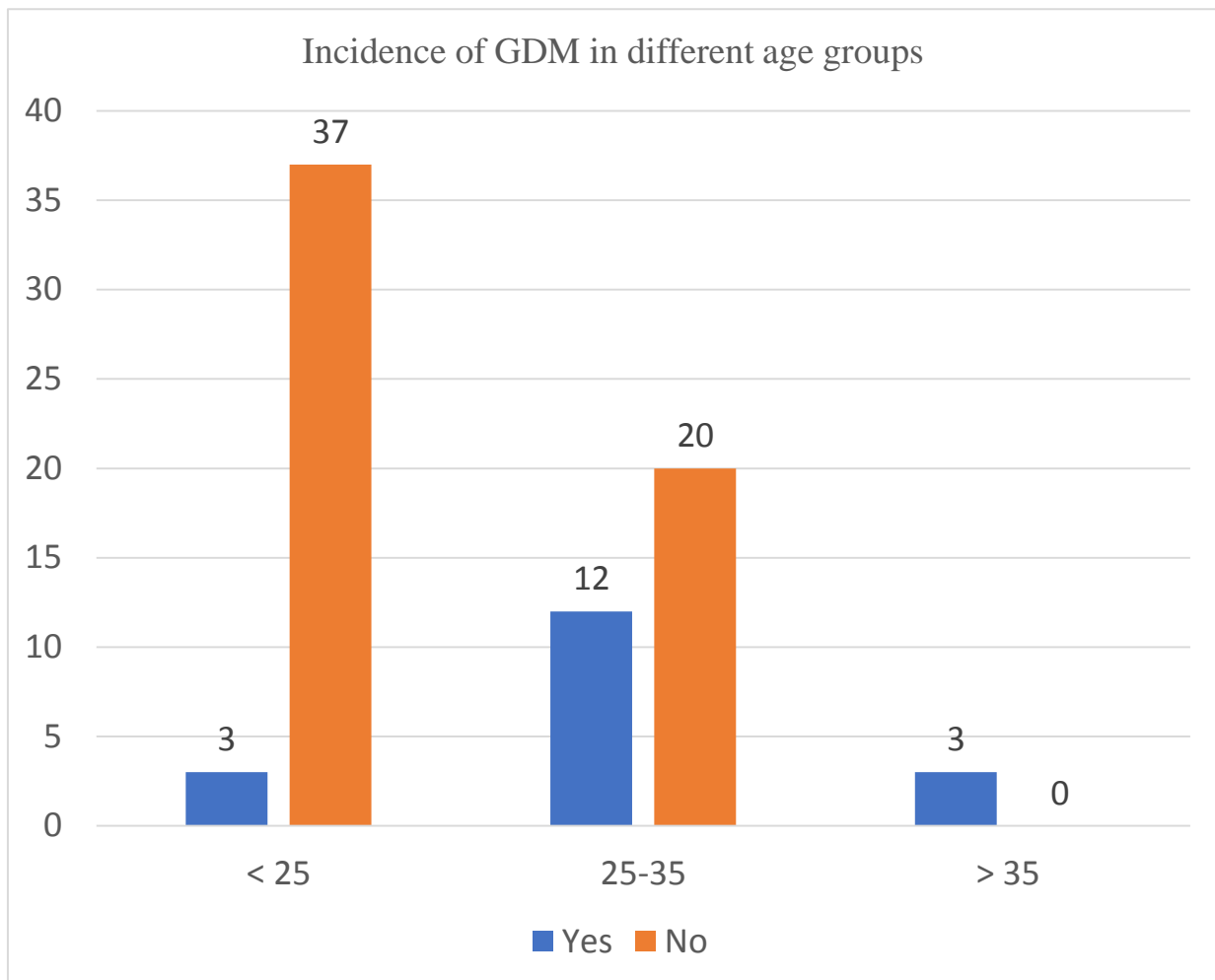


Tabel 5: Incidence of GDM in different age groups.

Age group	GDM		
	Yes n(%)	No n(%)	Total n(%)
< 25	3(7.5)	37(92.5)	40(100.0)
25-35	12(37.5)	20(62.5)	32(100.0)
> 35	3(100.0)	0(0.0)	3(100.0)
Total	18(24.0)	57(76.0)	75(100.0)

Among the 22 pregnant women (68.8%) in the 25-35 age range, poor glycemic control becomes more apparent as one moves older, as seen in table 4 of the age-wise distribution of the sample with respect to glycemic control. Among the members, 10 (31.3%) had poor glycemic control, meaning their HbA1c levels were between 6.5 and 7.5%. Among the members older than 35 years, 3 (4.2%) were also affected. Of the participants, 2 (66.7%) had inadequate glycemic control and 1 (33.3%) had fair glycemic control, which is less than 6.5%. A statistically significant relationship exists between age and glycemic control (chi-square = 10.5; P = 0.005).

X2 Value- 18.66, P value <0.001* Highly significant



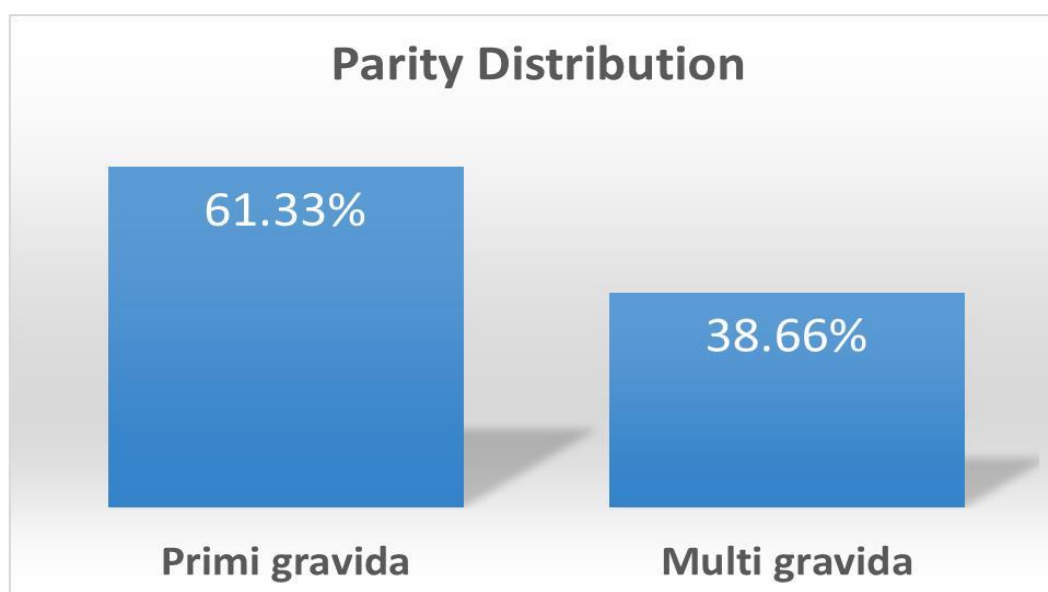
Graph 5: Incidence of GDM in different age groups.

Out of 75 pregnant women participated in the study 18 members developed GDM in second trimester. Among 18, 3 members belong to <25 years age group, 12 belong to 25-35 years age group and 3 belong to more than 35 years age group. Chi – square value being 18.66 and p value being <0.001 which is highly significant. This in turn shows the impact of age over development of Gestational Diabetes Mellitus.

Table 6: Parity Distribution Total sample tested-75 1ST Trimester

Parity	Number	Percentage
Primigravida	46	61.33%
Multigravida	29	38.66%

Table 6 shows the parity distribution of study population. Out of 75 study participants, 46 (61.33%) pregnant women belong to primigravida, 29 (38.66%) belong to multigravida. Majority belongs to primigravida.



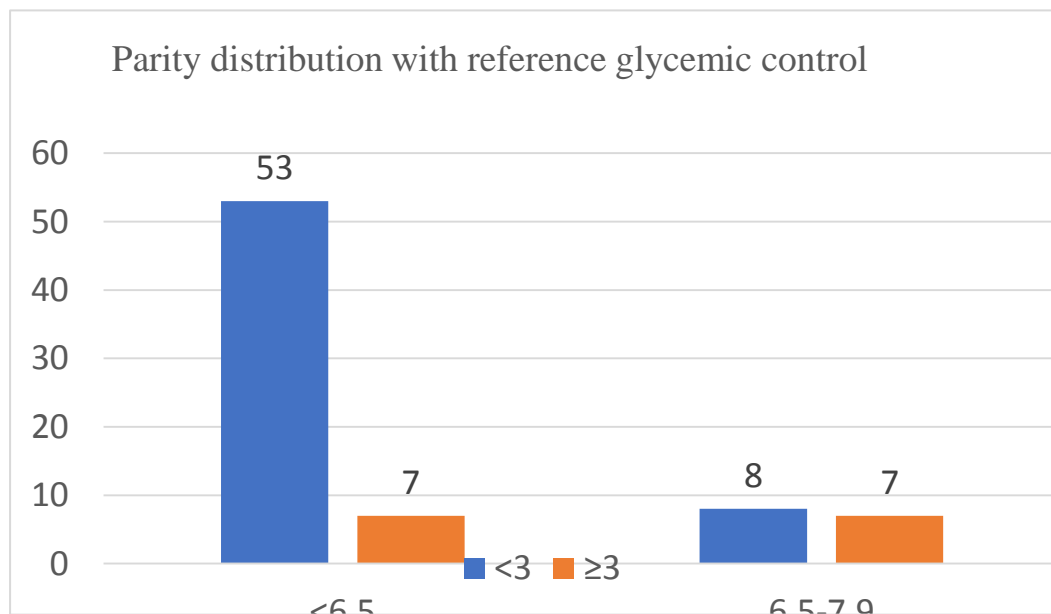
Graph 6: Parity Distribution Total sample tested-75 1ST TRIMESTER

Table 7: Parity distribution with reference glycemic control

Parity	HbA1C group		
	<6.5 n(%)	6.5-7.9 n(%)	Total n(%)
<3	53(86.9)	8(13.1)	61(100.0)
≥3	7(50.0)	7(50.0)	14(100.0)
Total	60(80.0)	15(20.0)	75(100.0)

The above table shows the parity distribution with reference to glycemic control. Among 75 study participants 60 were with fair glycemic control and 15 were with poor glycemic control. Out of 15 poor glycemic control participants 8 (13.1%) belong to parity of <3 and 7 (50%) belong to parity of ≥3 parity group. This distribution is with chi-square value 9.68 and p value of 0.002 which is highly significant. This clearly shows that women with low parity are with poor glycemic control.

X² value-9.68, P value 0.002*, Significant

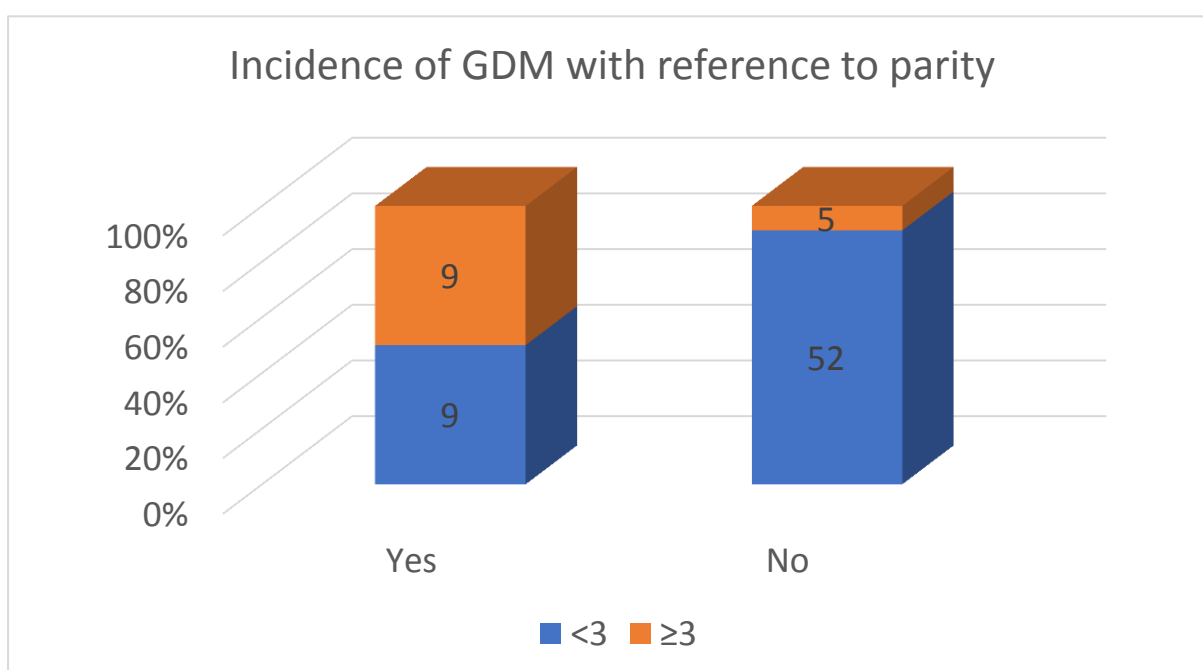


Graph 7: Parity distribution with reference glycemic control

Table 8: Incidence of GDM with reference to parity

Parity	GDM			Diabetes
	Yes N (%)	No N (%)	Total N (%)	
<3	9(14.8)	52(85.2)	61(100.0)	<0.001*
≥3	9(64.3)	5(35.7)	14(100.0)	
Total	18(24.0)	57(76.0)	75(100.0)	

X² value-15.31, P value <0.001*-Highly significant

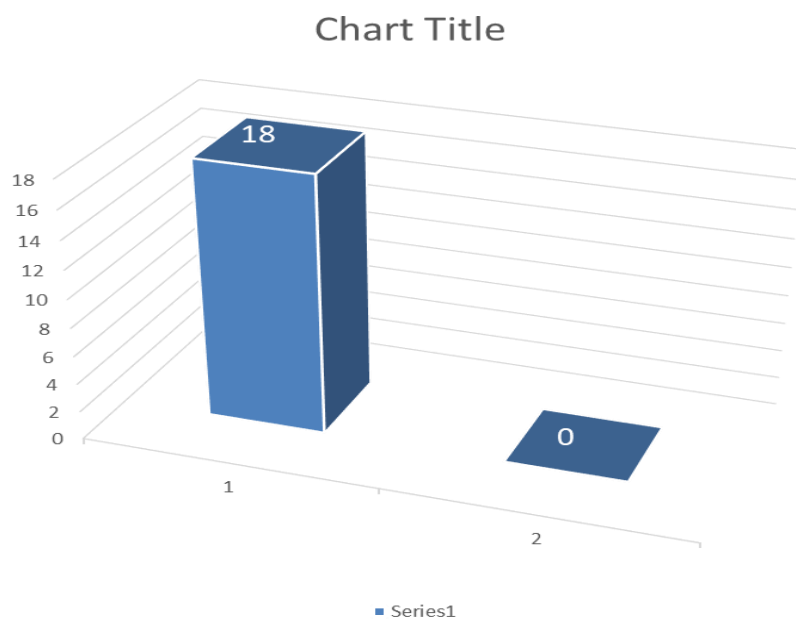


Graph 8: Incidence of GDM with reference to parity

The above table and graph clearly depict incidence of GDM with reference to parity. Out of 61 study participants belonging to <3 parity group 9 (14.8%) developed Gestational Diabetes Mellitus. Out of 14 study participants belonging to ≥3 parity groups 9 developed Gestational Diabetes Mellitus. So out of 75 study participant 18 (24%) developed Gestational Diabetes Mellitus. This is with chi-square value of 15.31 and p value of <0.001 which is highly significant. So, parity is having a significant impact over development of GDM.

Table 9: Abnormal HbA1c correlates with excessive weight gain
Abnormal HbA1c-18

Present	Number	Percentage
Present	18	100%
Absent	0	0%



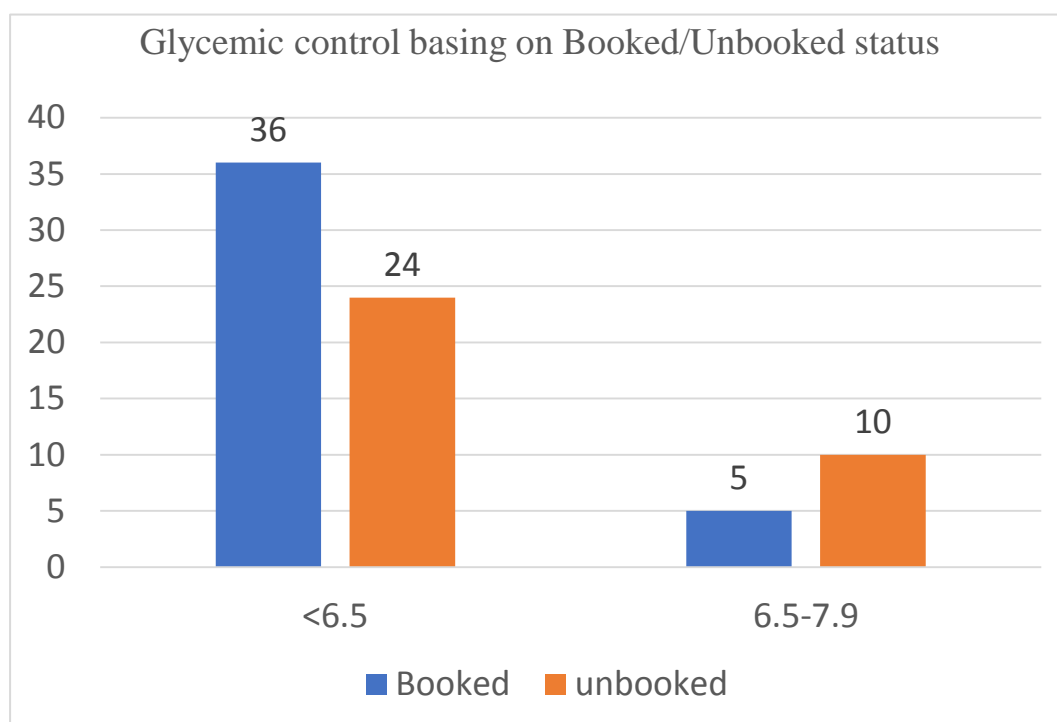
Graph 9: Abnormal HbA1c correlates with excessive weight gain
Abnormal HbA1c-18

Out of 75 study participants 18 were with abnormal HbA1c. All the 18 (100%) women have shown excess weight gain during their 2nd trimester.

Table 10: Glycemic control basing on Booked/Unbooked status

Booked/unbooked	HbA1C group		
	<6.5 n(%)	6.5-7.9 n(%)	Total n(%)
Booked	36(87.8)	5(12.2)	41(100.0)
unbooked	24(70.6)	10(29.4)	34(100.0)
Total	60(80.0)	15(20.0)	75(100.0)

X² value-3.44, P value- 0.064 Not significant



Graph 10: Glycemic control basing on Booked/Unbooked status

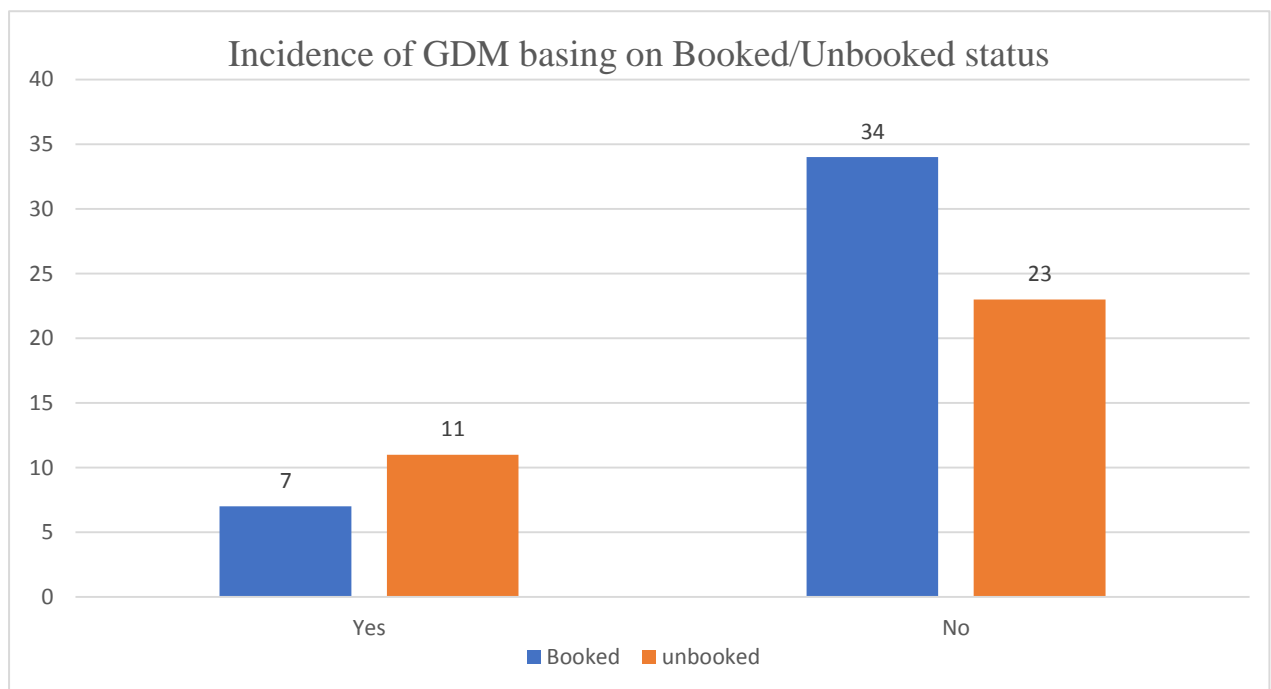
Above table and graph shows the booked and unbooked status of 75 study participants with reference to HbA1c values. Out of 75, 41 were booked cases and 34 were unbooked cases. Among 41, 36 (87.8%) were with fair

glycemic control and 5 (12.2%) were with poor glycemic control. Out of 34 unbooked cases 24 (70.6%) were with fair glycemic control and 10 (29.4%) were with poor glycemic control. This result is with chi-square value of 3.44 and p value of 0.064 which is not significant. This shows there is no correlation between booked status and glycemic control.

Table 11: Incidence of GDM basing on Booked/Unbooked status

X² value-2.37, P value-0.123 Not significant.

Booked/unbooked	GDM		
	Yesn(%)	Non(%)	Total n(%)
Booked	7(17.1)	34(82.9)	41(100.0)
unbooked	11(32.4)	23(67.6)	34(100.0)
Total	18(24.0)	57(76.0)	75(100.0)



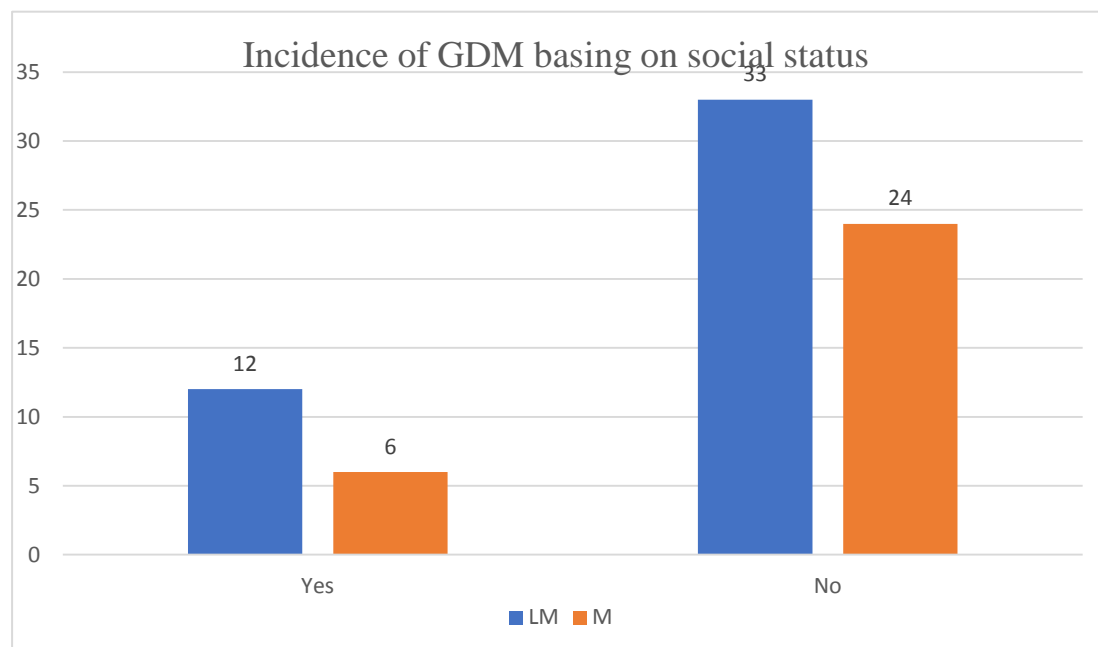
Graph 11: Incidence of GDM basing on Booked/Unbooked status

Above table and graph shows the incidents of GDM based on booked and unbooked status. Out of 41 booked cases 7 (17.1%) were developed GDM in 2nd trimester. Out of 34 unbooked cases 11 developed GDM in 2nd trimester. This is with chi-square value of 2.37 and p value of 0.123 which is not significant. This depicts that there is no impact of booked status over development of GDM.

Table 12: Incidence of GDM basing on social status

X² value-0.43, P value-0.508 Not significant

Social status	GDM		
	Yes n(%)	No n(%)	Total n(%)
LM	12(26.7)	33(73.3)	45(100.0)
M	6(20.0)	24(80.0)	30(100.0)
Total	18(24.0)	57(76.0)	75(100.0)



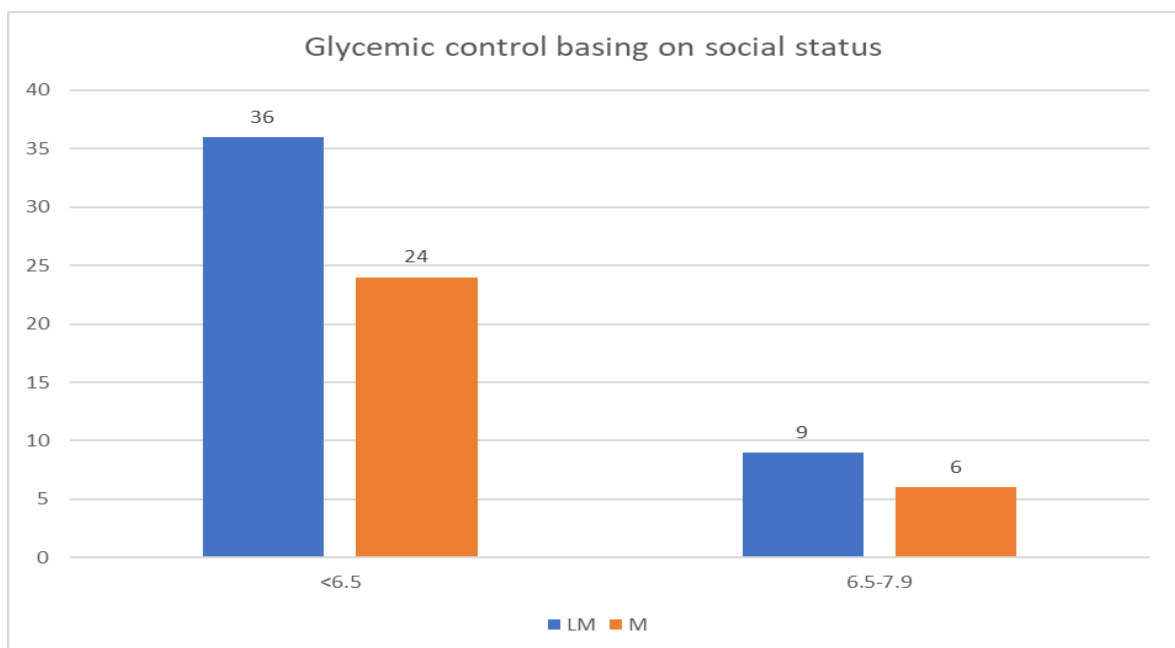
Graph 12: Incidence of GDM basing on social status

Among 75 study participants 45 belong to lower middle class income status and 30 belong to middle class income status. Out of 45 lower middle class 12 (26.7%) developed GDM and out of 30 middle class group 6 (20%) developed the GDM. This with χ^2 value of 0.43 and p value of 0.508. which is not significant. This shows no correlation between income status and GDM.

Table 13: Glycemic control basing on social status

Social status	HbA1C group		
	<6.5 n(%)	6.5-7.9 n(%)	Total n(%)
LM	36(80.0)	9(20.0)	45(100.0)
M	24(80.0)	6(20.0)	30(100.0)
Total	60(80.0)	15(20.0)	75(100.0)

χ^2 value – 0.00 p value – 1.00 Not significant



Graph 13: Glycemic control basing on social status

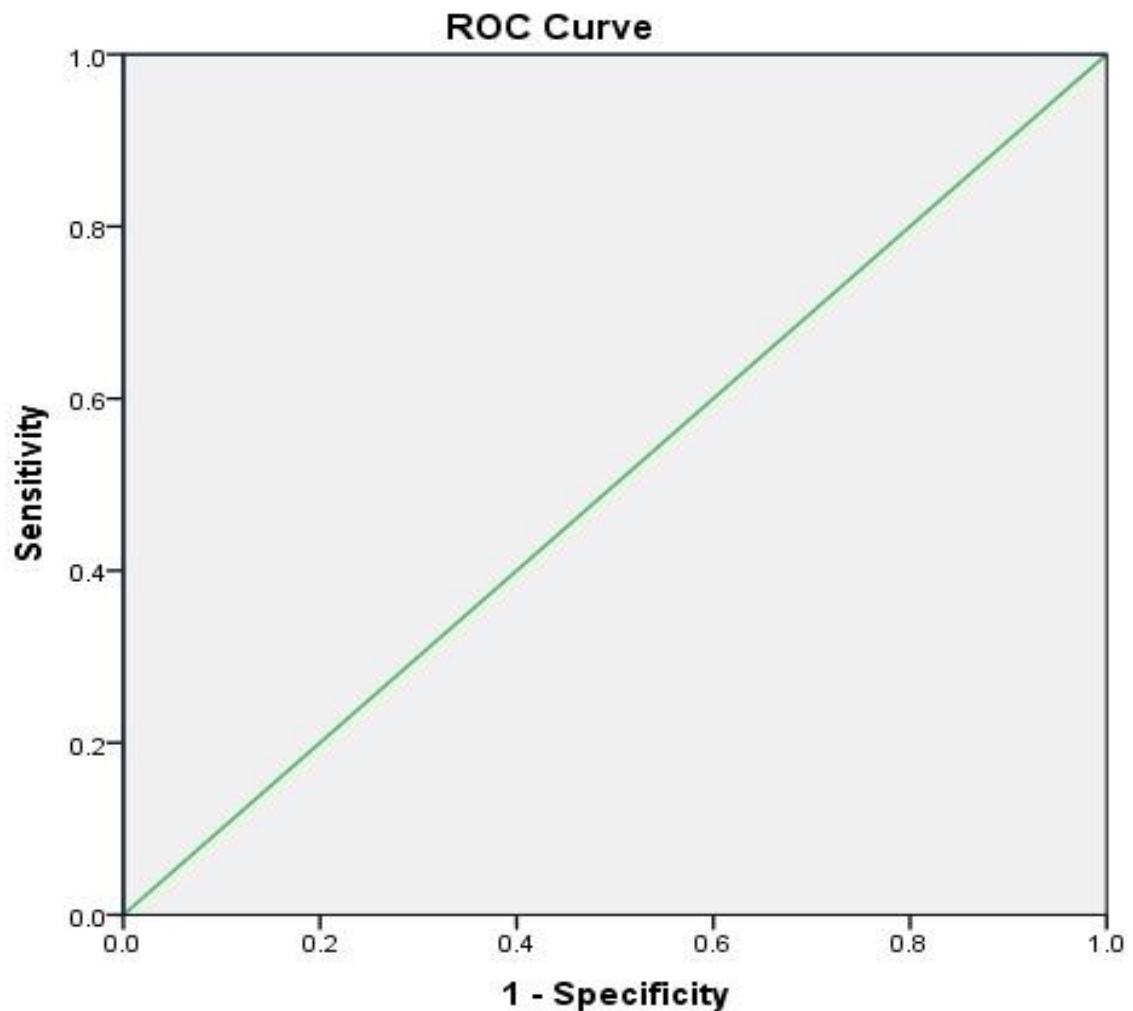
Out of 45 lower middle class social status 36 (80%) were with fair glycemic control and 9 (20%) were with poor glycemic control. Out of 30 middle class social status 24 (80%) with fair glycemic control and 6 (20%) poor glycemic control. This is with chi-square value of 0.00 and p value of 1.00 with is not significant. This shows that social status as no affect over the glycemic control.

Name of the Diagnostic test		OGTT		
		Positive	Negative	Total
HbA1c	Positive	18	0	18
	Negative	0	57	57
	Total	18	57	75

Table 14: HbA1c distribution with oral glucose tolerance test 24-28 weeks

Among 75 study participants 18 were with abnormal HbA1c in 1st trimester and all have shown OGTT test positive and developed gestational diabetes mellitus in 2nd trimester.

Graph 15: ROC Curve for HbA1c in first trimester in predicting GDM



Area under the curve = 1

P value = <0.0001*(Highly significant)

Significant cut off HbA1C value in predicting diabetes mellitus by OGTT was found to be 5.7 (with Sensitivity is 100% and False positivity rate is 0%)

DISCUSSION

A decorative graphic consisting of a horizontal line and a vertical line intersecting at the right end of the horizontal line. Both lines have a thin, light gray shadow offset to the right and bottom, creating a 3D effect.

DISCUSSION

My research proved my hypothesis that early prediction of Gestational Diabetes mellitus can be done by using HbA1c in first trimester of pregnancy, based on my results.

SUMMARY OF RESULTS.

Out of all the samples that were analysed, 53.33 percent were under the age of 25, 42.66 percent were between the ages of 25 and 35, and 4.2 percent were 45 and over.

Out of 75 participants, 18 (or 24% of the total) had HBA1C levels that were considered abnormal.

Among those aged 25–35, abnormal Hba1c levels were more common.

People in the 25–35 age bracket were more likely to have inadequate glycemic control, as measured by HbA1c values ranging from 6.5-7.9.

The prevalence of GDM was highest among those aged 25–35.

Our research included 75 patients; 61.33 percent were first-time mothers, and 38.56% were mothers of multiples.

Eight out of fifteen patients with poor glycemic control had a parity of less than three, whereas seven had a parity more than three. This indicates that women with low parity also have poor glycemic control.

All of the participants in our research who had abnormal haemoglobin A1c levels gained a lot of weight in the second trimester.

The incidence of GDM was not correlated with the booked status of pregnancy.

We found no evidence that women's socioeconomic status was associated with the development of GDM.

SCREENING AND DIAGNOSIS OF GESTATIONAL DIABETES MELLITUS

Among metabolic issues that could make during pregnancy, gestational diabetes mellitus is by far the most normal.

How Typical is GDM? Is rising right now a result of following causes.

Maternal age and weight record both create.

Raising the level of life

A steadily increasing number of people are choosing to check for diabetes.

With an extent of 9.3-25.5%, the overall pervasiveness of GDM is 17.8%.

Somewhere in the range of 24 and 38 weeks of hatching, a glucose challenge test or glucose obstruction test is habitually used to break down gestational diabetes melitus. Exactly when a pregnant woman has hyperglycemia, her youngster turns out to be faster in light of the fact that glucose crosses the placenta. The hatchling could have persevered through harmed if the opportunity to self-screen blood glucose levels and give medications had not

sneaked past in view of a concede in lab testing. 44It is critical to test for metabolic abnormalities in early pregnancy in light of the rising event of strange glucose levels among pregnant women.

We want more information to make any recommendations on screening systems or shorts at the present time.

A two-step process is supposed to confirm an assurance of gestational diabetes mellitus. Two of these tests are open: a 75-gram oral glucose challenge test that simply requires one sitting, and a 50-gram oral glucose screening test that, for individuals who test positive on the screening, is followed by a demonstrative 100-gram oral glucose obstruction test.

Early screening, assurance, and treatment of gestational diabetes mellitus (GDM) should not be recommended due to a shortfall of trustworthy evidence on the risks and advantages of doing as such.

It is feasible to test for and treat contorted glucose absorption frameworks in pregnancy during the essential trimester.

Indications of insulin resistance, adipocytes, blood glucose, and bothering are early predictors of GDM; in any case, the last two are not every now and again used in clinical practice. 52

Glycemic pointers like hemoglobin A1c, fasting glucose, and postprandial glucose are among them.

Insulin deterrent pointers integrate sex synthetic limiting globulin and combustible markers consolidate c-responsive protein and development rottenness factor-alpha. Insulin levels while fasting.

Markers made from adipocytes: leptin and adiponectin.

Placental advancement factor, placental exosomes, and markers made from the placenta, (for instance, follistatin-like 3).

Glyconectin, ferritin, alanine amino transferase, dissolvable virtuoso renin receptor, and others.

Markers most often used consolidate blood glucose and glycosylated hemoglobin.

The American Diabetes Association consolidated the hemoglobin A1c test to its overview of expressive principles for everyone in 2010. that year (19). The suggestive end was set at 6.5%. Unfavourable pregnancy results were similarly associated with unusual HBA1C levels, according to the HAPO research.

Using HbA1c, one can see the regular blood glucose levels over the course of 90 days.

Advantages of Hba1c

Its pre-analytical stability is higher.

The diversity among individuals is minimal.

It undergoes less daily changes.

There is less fluctuation than with glucose testing, and it is easy to use and doesn't need fasting.

Things like mood, stress, and food have little effect on haemoglobin A1c.

I will go over the factors that impact hba1c.

Blood disorder

Decreased rate of intestinal transit

Expectant mothers' physiological hydremia

The exposure of haemoglobin to glucose and the increased red cell turnover are both affected by changes in the RBC life span. A decrease in red blood cell (RBC) life span will impact total hematocrit (HbA1c) production because glycosylation of haemoglobin occurs at a lower rate. Glycosylated haemoglobin varies mostly due to differences in red blood cell (RBC) life span, which are unrelated to changes in blood glucose levels. Underestimation of blood glucose levels as shown by HbA1C detection value occurs in type 2 DM patients due to their reduced life duration.⁵³

Changes to the diet

Abnormal haemoglobins.

Glycosylation of haemoglobin without the need of enzymes.

Weak kidneys throughout time.

Haemoglobin A1c problems include

It drops a lot when you're pregnant. While pregnant, the upper normal limit of haemoglobin drops from 6.3% to 5.7% in the first trimester and to 5.6% in the third.

Disadvantages of oral glucose tolerance test are

Careful planning for the patient.

This is a pain

A night without food is necessary.

Decreased repeatability

Three blood draws are necessary.

Symptoms of nausea and vomiting, such as delayed stomach emptying, are more common in pregnant women.

For the reasons given, not all women will undergo oral glucose tolerance testing.

Many major obstetric complications are more likely to occur in a pregnant woman with gestational diabetes. Some of these complications include problems with the baby's growth, shoulder dystocia, birth injuries, premature delivery, and a rise in the use of caesarean sections [1]. Oral glucose tolerance tests may be administered in either a "two-step" or a "one-step" fashion; the former involves a 50 g (non-fasting) screening test and the latter involves a 100 g OGTT for those who come up positive²⁵. On the other hand, some pregnant women report that the oral gestational test (OGTT) is

uncomfortable, tiresome, and takes too much time. The patient must fast for at least three hours, sit for at least three hours, and have their blood drawn at least three times. The delayed emptying of the stomach may also cause nausea and vomiting in the gravida. These disadvantages account for around 10% of the pregnant women who do not finish the glucose challenge test¹.

Compared to fasting plasma glucose (FPG) and oral glucose tolerance testing (OGTT), glycosylated haemoglobin (HbA1c) has a few benefits, such as being more consistent prior to analysis, easier to use (no fasting required), and less affected by strain and disease concerning ordinary variance. Factors that affect the red blood cell half-life or non-enzymatic glycation of hemoglobin³³ mostly affect haemoglobin A1c, despite its strength as an indication of glycemia. The rapid red cell turnover or iron deficiency during pregnancy makes HbA1c testing for gestational diabetes mellitus (GDM) more challenging. Regardless, the IFCC has standardised reference intervals for HbA1c for each trimester. More and more research suggests that monitoring HbA1c during the first trimester might help identify pregnant women at increased risk of obstetric complications and the recognition of hyperglycemia. For the purpose of assessing for previously undiscovered diabetes (clear diabetes) using similar analytical procedures employed for the general population, HbA1c findings have been used at the necessary pre-birth meeting.³ A woman and her unborn child are more likely to have

complications if her haemoglobin A1c level is 5.9% or higher (41 mmol/mol) during the first trimester of pregnancy, regardless of whether gestational diabetes mellitus (GDM) is detected in the second trimester.^{4,5}

COMPARISON OF RESULTS WITH OTHER STUDIES.

The 25–35 age group had the highest prevalence of abnormal HbA1c readings in our research, at 14.28%.

In study conducted by Pavithra, Sreelatha et al⁵⁴ at ESIC MC and PGIMSR, BANGALORE, majority were in the age group of 26-30, similar to our study, where range of abnormal HbA1c were more in age group of 25-35 years. In the above study patients diagnosed with GDM were taken and their HbA1C levels were measured and maternal and perinatal outcomes assessed.

In study conducted by Madleen Lemaitre et al⁵⁵. To study the association between HbA1c levels and adverse pregnancy outcomes in patients in pregnant women with type1 diabetes mellitus, mean age of abnormal HbA1c and patients with GDM was 30.1 +/- 4 years., which is similar to our study.

The mean age of GDM in years was 32.1 +/- 5.2 years, similar to our study, in the study conducted by sofıya amylydi et al⁵⁶ with p value of 0.02.

In the study conducted by Marie Parfait uzimana et al.⁵⁷ HbA1c levels greater than 5.5 was found more in the age group of >35 years ,with 44.2%.

Name of the study	Mean Age
Pavithra et al ⁵⁴	28 +/_ 2Years
Madleen Lemaitre et al ⁵⁵	30.1 +/_ 4 years
sofiya amyliidi et al ⁵⁶	32.1 +/_ 5.2 years
Marie parfait uzimana et al ⁵⁷	>35 years
Mehraz valadan et al. ⁴⁷	32.64+/_5.49

Abnormal HbA1c levels were more found in multigravida with percentage being 13.33%.

In study conducted by Pavithra, Sreelatha et al⁵⁵ at ESIC MC and PGIMSR,BANGALORE,majority were in multigravida (53.93%.

In the study conducted by sofiya amyliidi et al⁵⁶, prevalence of abnormal hba1c was more in multiparous women (72%).

In the study conducted by marie parfait et al.⁵⁷., abnormal HbA1c was found more in women with multiparous women. (55.8%)

Abnormal HbA1c

Name of the study	Nulliparous	Multiparous
Pavithra et al ⁵⁴	----	Multigravida (53.93%).
sofiya amyliidi et al ⁵⁶	----	Multiparous women (72%)
Marie parfait et al. ⁵⁷	----	Multiparous women (55.8%)

All women with abnormal HbA1c shown excessive weight gain corresponding to 100%.

Gestational weight gain decreases Insulin resistance or worsening of preexisting insulin resistance might develop in certain women due to their subcutaneous storage capacity.

In the study conducted by Marie Parfaitetal⁵⁷, abnormal hba1c was found in women with normal pre pregnancy BMI.(53.6%).

Bozkurt et al. discovered that higher Hba1c levels were linked to higher pre- and post-pregnancy body mass indexes. - A Researchers Latife Bozkurt, Christian S. Göbl, and colleagues⁵⁸ aimed to compare the clinical and pathophysiological features of women whose GDM began early or late, up to the 21st week of the pregnancy. Insulin resistance, which is in part explained by a greater degree of fat, affects the early presentation of GDM. But GDM also showed signs of β -cell malfunction. Defective compensating mechanisms already emerge in early pregnancy, suggesting a late onset. In conclusion, using the IADPSG criteria, we analysed dynamic indices derived from OGTT measurements and found that women with early and later GDM showed significant differences in insulin sensitivity and B-cell function parameters, which were partially explained by the degree of overweight or obesity. Insulin resistance was greater in women with early-onset GDM compared to controls, although it was similar in those with later-onset. This

might suggest that women who have early signs of gestational diabetes may have a longer period before the disease officially begins if they had insulin resistance to begin with, on top of any developing cell malfunction during pregnancy. Women whose GDM develops later in pregnancy, on the other hand, exhibit signs of β -cell malfunction as early as the first trimester, suggesting that their bodies have burned through their compensatory mechanisms. The development of better methods for screening for metabolic disorders during pregnancy and for performing an accurate risk classification might benefit from the discovery of pathophysiological anomalies linked to the timing of beginning of GDM.

In a review an outline of diabetes mellitus in pregnant ladies with corpulence, directed by Mohammed Bashir et al⁵⁹, GDM risk increases with increased BMI in a portion subordinate example.

Positive correlation of abnormal HbA1c with reference to BMI

Name of the study	Normal BMI	Abnormal BMI
Mohammed Bashir et al	---	Yes
Marie parfait et al	Yes	No
Bozkurt et al	---	Yes
Mehraz valadan et al	---	Yes
Paula Breitenbah Renz.	---	Yes
Jyothi balani et al	---	Yes

In our study Positive correlation of abnormal HbA1c with 2hr OGTT at 24-28wks of gestation found to be 100%.

In a study conducted by Mehrnaz valadan, zeinab bahramnezhad, Fatemeh Golshahi & Elham feizabad⁴⁷, BMC pregnancy and child birth, prevalence of GDM was 16.4%. High prevalence in the above study is due to increased high risk referrals in view of tertiary hospital, and also using carpenter couston criteria, which has lower threshold for detection of diabetes mellitus. In this study, AUC value was found to be of 0.84, which has shown excellent predictive value of HbA1c for diagnosis of GDM. In this study, HbA1c with value more than 6% were with high risk for GDM, and GDM was not seen in women with HbA1c less than 4.55%. Using HbA1c declined OGTT IN 40% OF Patients. (18)

In concentrate on drove by jyoti balani et al⁶⁰ early pregnancy Of the 160 women, 39 made GDM, giving ordinariness in this undertaking people of 24.4%. The median (interquartile range) for HbA1c was 33(31-36)mmol/mol. Women with GDM (n=39) had a higher BMI, were even more often of Asian identity (26%), had more pervasive hypertension in pregnancy, had higher first trimester HbA1c, and a higher fasting and 2-hour glucose at OGTT contrasted and women without GDM. HbA1c more imperative than 5.7% had a high expressness near 100 percent and a high certain likelihood extent (9.31), and low responsiveness 7.7%. This study

shows that those with high HbA1c were found to have GDM at an early gestation. High HbA1c didn't show association with macrosomia. HbA1c level more than 5.9% showed association with blood poisoning. The AUC of first trimester HbA1c for predicting GDM was 0.708 (95% CI 0.631-0.777; $p < 0.001$) indicating incredible farsighted incentive for GDM (Figure 1). The Youden index (mindfulness + unequivocality - 1) was maximal at HbA1c of ≥ 33 mmol/mol (5.2%) with 69.2% responsiveness and 63.6% identity. Nevertheless, using this model, the positive farsighted worth (PPV) for GDM was simply 38.1 (95% CI 31-45.7).

In study conducted by Xiaoxiao Peng et al.⁶¹, AUC for predictive value of HbA1c was 0.67, which showed there is high reliability of HbA1c in predicting GDM. Other parameters used are Fasting blood glucose, One hour post prandial blood glucose, two hours post prandial blood glucose. Among this, one hour post prandial blood glucose showed high predictive value.

According to research by Rajesh Rajput et al.⁶² Compared to women with GDM, those without the condition had a much lower mean HbA1C level. With an area under the curve (AUC) of 0.805, a specificity of 97.2%, and a sensitivity of 28.6%, the HbA1c cutoff value was $> 5.95\%$. The specificity for identifying gestational diabetes mellitus type 2 in women was 61.1% and the sensitivity was 85.7% when the HbA1c cutoff value was between 5.45% and 5.95%. The researchers found that using HbA1c in

conjunction with OGTT eliminated the requirement for GTT on its own. Patients with gestational diabetes mellitus had greater mean HbA1c levels than those without the condition.

Sophia Amulidi et al. 56 tracked down a genuinely critical relationship between's hba1c levels and the beginning of GDM in their investigation. The research discovered that test execution had unfortunate responsiveness and explicitness at the 4.5% HbA1c cutoff edge. The AUC of first trimester HbA1c and irregular blood glucose levels showed a huge improvement when maternal qualities such age, BMI, earlier GDM, and family background of diabetes mellitus were considered.

As per Osmundson study, 63 Foreseeing GDM with a 13% responsiveness and 94% particularity was connected to a HbA1c in the prediabetic range during the principal trimester. One hundred 26 ladies (9.6%) were prediabetic as indicated by A1c, out of 1438 who satisfied incorporation measures. More Asian ladies (62.7%) and more seasoned (34.1 versus 32.5, $p < 0.001$) were prediabetic than more youthful ladies (52.5), with a p -worth of 0.001. The main trimester OGTT recognized GDM in 3.6% of prediabetic ladies. The probability of prediabetic ladies being determined to have gestational diabetes mellitus at any phase of the pregnancy was 31.8% contrasted with typical ladies' 12.3% ($p < 0.001$). No genuinely tremendous

changes were found in the level of individuals with GDM who required insulin (7.9% versus 4.9%).

One examination found that stood out from an ordinary benchmark bunch, those with GDM had much higher HbA1c levels (In a little while Sung Kwon et al., 2016). At a cut-off worth of 5.05% (32 mmol/mol), HbA1c displayed 91.3% responsiveness and 62% expressness for GDM end, including the 100-g OGTT as a wellspring of viewpoint. The responsiveness was 73.6% and the unequivocality was 77.2% at a taken out worth of 5.25% (34 mmol/mol). Those with pre-diabetic metabolic condition had higher HbA1c levels generally through pregnancy stood out from those without PDM (5.91 [41 mmol/mol] versus 5.44% [36 mmol/mol], $p\} 0.001$). At a cut-off worth of 5.55% (37 mmol/mol), ROC twist assessment studied the insightful worth of HbA1c for PDM, and the results showed a responsiveness of 78.6% and a distinction of 72.5%.

GDM was recognized in 152 ladies (13.1%) out of 1195 ladies (David Benaiges et al.,²⁶). The ROC region for HbA1c as a GDM symptomatic instrument was 0.679 (95%CI 0.631-0.727). Utilizing a HbA1c cutoff of 4.8% (29 mmol/mol), the responsiveness was 96.7% (95%CI 93.9-99.5), the explicitness was 10.1% (95%CI 8.3-12.0), and the negative prescient worth was 95.3% (95%CI 91.3-99.3). The positive prescient worth, particularity, and responsiveness were all 89.3%, 25.4%, and 25.4%, separately, for a

standard in worth of 5.6% (38 mmol/mol), with a 95% certainty span (CI) of 24.4-38.9. Despite the fact that the standard in limit can't be utilized to analyze GDM because of its unfortunate positive prescient worth, it very well may be utilized to recognize ladies who are at high gamble of fostering the sickness and can be analyzed utilizing a one-step system. The recommended approach would save 6.5% of the absolute expense contrasted with the standard method. Also, this investigation discovered that first-trimester HbA1c levels probably won't be explicit or delicate enough to analyze GDM. Notwithstanding, by changing the edges, both higher and lower, the analytic cycle can be made simpler, prompting less oral glucose resilience tests and less issue for pregnant patients. (27). Significant focuses from the previously mentioned research incorporate

HbA1c in first trimester may be an attractive choice for GDM screening.

First trimester HbA1c didn't have adequate ability to analyze GDM.

Utilization of lower and higher limits of HBA1C improves on the indicative cycle ,of GDM.

Ribal Kattini et al.⁶ state We investigated 121 summaries that were pertinent. Out of the 32 tests that were qualified for a complete overview, 11 satisfied the necessities as a whole. A higher gamble of GDM was associated with HbA1c esteems more than 5.7, as indicated by all surveys. People with GDM-delivering levels >6.0 were recognized. Pre-eclampsia, constrained

work, shoulder dystocia, cesarean area, huge for gestational age child weight, macrosomia, characteristic irregularities, and perinatal demise were among the ominous pregnancy results connected with raised HbA1c levels in four out of six tests. Two examinations neglected to track down a relationship with upsetting occasions. The HbA1c readings in this exploration were somewhere in the range of 5.7% and 6.4%. Shown a serious level of consistency for the determination of gestational diabetes mellitus in the intense phases of pregnancy. Higher HbA1c values in the primary trimester of pregnancy were likewise decidedly associated with the improvement of gestational diabetes mellitus, as per this focus. HbA1c cut off . High expressness levels of 95% to 98% were viewed as in 5.7% or over 5.9% of the cases.¹³

Hatice kansu-celik et al.⁶⁶ found that women who were likely to acquire gestational diabetes mellitus had much higher median hba1c and fasting plasma glucose concentrations. The diagnostic accuracy was 83.55%, specificity was 89.8%, and sensitivity was 34.78% for the highest youndex cutoff with a cutoff value greater than 5.6%. The prediction reliability for GDM was enhanced when fasting plasma glucose and hba1c were combined. GDM was viewed as in 67.3% of pregnant ladies, as per Paula Breitenbach renz and Gabriella Cavagnoli. Displayed a responsiveness level of simply 7% and an explicitness of 100 percent with an end of 6.5% and above. With a

HbA1c worth of 5.8% as the end, utilizing the ROC bend and the OGTT as the reference standards, a similar exploration tracked down a low responsiveness of 26.4% and a particularity of 94.9%. For GDM, the positive probability proportion was 5.14 and the negative probability proportion was 0.78. A 5% HbA1c cutoff was ready to determine GDM to have a proper responsiveness of 89.7 percent however an unfortunate explicitness of 32.6 percent. Since 38% of people with GDM might be related to a HbA1c test alone, an end worth of 5.8 for this test delivers the drawn-out glucose resistance test superfluous in close to 33% of cases. The ROC bend examination in this exploration showed that HbA1c performed likewise when contrasted with the WHO 1999 and WHO 2013 reference measures for GDM determination (AUC Upsides of 0.714 and 0.756, separately). the year (19). This recommends that, no matter what the measures used, there are no varieties in the exhibition of OGTT against hba1c. The consequences of this exploration propose that OGTT, related to other HbA1c endpoints, might be more useful in the analysis of GDM. The reason for the review was to decide if high-risk pregnant ladies could be distinguished involving non-diabetic reaches for first-trimester fasting plasma glucose and HbA1c readings. The specialists L. Mañé, J. A. Flores-le Roux, N. Gómez, and others teamed up on the request. The information used to assess an impending buddy came from April 2013 to September 2015. All

pregnant ladies had their hemoglobin A1c and fasting plasma glucose levels checked during the principal prenatal blood test. The gestational age at which gestational diabetes mellitus was recognized was 24-28 weeks. The most widely recognized complexities were macrosomia and toxemia, whereas different results including unexpected labor, cesarean area, and enormous for gestational age were viewed as discretionary. To find a relationship, specialists looked at different fasting plasma glucose and HbA1c cut-off levels. The objective of the result study was to incorporate 1,228 pregnancies. In the wake of considering every single pertinent variable, we found no connection between's fasting plasma glucose levels and pregnancy results. Hemoglobin A1c was a more exact predictor of entanglements during pregnancy in a multiethnic model contrasted with fasting plasma glucose levels in the main trimester. High hemoglobin A1c levels (39.9 mmol/mol or above) are associated with an increased gamble of macrosomia (OR2.69), toxemia (a three-get more than risk; 95% CI1.03-9.9), and enormous for-gestational age (a four-crease risk; 95% CI 1.49-11.07). Early HbA1c levels of 5.8% or higher (39.9 mmol/mol) are associated with a higher gamble of macrosomia.

An investigation was finished by Çetin, N.D. Güngör et al.⁶⁸ to determine the reasonability of hemoglobin A1c in the early phases of pregnancy for the distinguishing proof of GDM. All through the span of the essential trimester,

the levels of hemoglobin A1c (ftHbA1c) were continued in 195 pregnant women participating in this assistant exploration. Between the eleventh and fourteenth extended lengths of incubation, individuals had their blood tests taken. Starting there forward, each participant, developed 24-28, was given a standardized 75 g OGTT. An amount of 195 pregnant women gave the information used to make these determinations. The examination included 32 women who had been determined to have GDM. While comparing individuals with and without GDM, the commonplace ft-HbA1c level was 5.52% ($p \leq 0.000$).

Just seven women (3.6% of the total) had fasting hemoglobin A1c esteems more prominent than as far as possible. Preceding acquiring gestational diabetes, these ladies were prediabetic. The ft-HbA1c cut-off incentive for differentiating GDM was 5.33%. Both responsiveness and identity were 71.9% and 82.8%, independently, at this edge. Just ft-HbA1c stood up as a bet factor for GDM. The analysis of the ROC twist (Fig. 1) uncovered that ft-HbA1c had a district under the twist of 0.839 ($p \leq 0.000$). In solicitation to distinguish GDM, a cutoff worth of 5.33% was used for ft-HbA1c. Both mindfulness and unequivocality were 71.9% and 82.8%, independently, at this cutoff. Perinatal issues of the patients and the mean ft-HbA1c. A confident sign for the determination of gestational diabetes mellitus, according to the survey's makers, was fasting hemoglobin A1c levels. An

increased bet of developing GDM is associated with ft-HbA1c readings of 5.33 percent or higher.

The reason for this study was to determine in case there is a link between developing GDM and having an early-stage hemoglobin A1c level under 6.5%. The assessment has been formed by countless individuals, among them being Claire B. and Sharon H.⁶⁹ During the essential trimester of pregnancy, is it safeguarded to test for unseen diabetes using hemoglobin A1c? Glycosylated hemoglobin (HbA1c) is an encouraging indicator as it shows the commonplace blood glucose level over the past a couple of months. Focuses on that used many markers, notwithstanding, found incongruous findings.²⁶ Even directly following controlling for maternal age and weight index, a fasting glucose (FG) level some place in the range of 5.6 and 7 mmol/L at the essential pre-birth visit was significantly associated with blood poisoning (OR2.7; 95% CI:1.2-5.9) and shoulder dystocia/birth injury (OR24.5; 95% CI:2.8-214.8). The usage of hemoglobin A1c for gestational diabetes mellitus risk delineation is appealing a consequence of its wide use in the finding of pregestational diabetes during the main typical pre-birth visits.

In created by D. Benaiges and J. A. Flores-le Roux, Region I.²⁶ applies. Marcelo et al. zeroed in on the manner that first trimester hemoglobin A1c testing is one likely method for screening for gestational diabetes

mellitus. First trimester HbA1c is insufficient for the disclosure of gestational diabetes mellitus. Possible definite rearrangements could come about in view of using upper and lower HbA1c rules. We determined whether hemoglobin A1c values measured during the essential trimester were helpful and reliable for diagnosing gestational diabetes mellitus. Pregnant women who had the choice to maintain a predictable weight gain all through their pregnancies were the subjects of this planned observational investigation. Both pre-birth screenings for gestational diabetes (24-28 weeks) and first-trimester HbA1c testing were coordinated on all pregnant women as an element of a two layered framework. To find out how sensitive and unequivocal HbA1c is for identifying GDM, we made a ROC twist and proposed a standard in, block definite system. An expense is associated with the suggested estimation. It has been shown that a first-trimester HbA1c doesn't have adequate unequivocality or repugnance for perceive GDM, whether or not that could deal with the determination procedure by changing the edge for oral glucose tolerance tests, which would moreover reduce related expenses and patient pain.

SUMMARY

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at the right end of the horizontal line. The horizontal line extends from the left edge of the page towards the right, and the vertical line extends from the bottom edge of the page upwards, crossing the horizontal line.

STRENGTH OF MY STUDY

Strength of my study include covering all singleton pregnant women, older than 18 yrs so that bias is reduced.

This is a prospective study, which helps in follow-up, eliminating disadvantages of retrospective studies.

This study is done in a single centre, with mixed population.

All the subjects underwent same type of assessment including laboratory workup and diagnostic procedures, which reduces confusion and bias.

Hba1C testing was done which does not require any special pretest preparation.

As fasting is not required for testing HBA1C levels, pregnant patients are not subjected to any fasting.

High risk women were excluded so that they might influence the prediction of HbA1c, removing confounding factors.

LIMITATIONS OF THE STUDY

Limitations of the study are more no. of participants could be included in the study.

Nutritional and life style characteristics are not studied, which would have given more information about predictability of HbA1c.

Predictive accuracy of HbA1c for GDM in correlation with FBS would have more supported my hypothesis.

Family history or genetic factors are not included in the study, hence possible variations in degree of HbA1c could o

Association between HbA1c and maternal and neonatal outcomes could have been assessed.

Integrated model which covers predictive factors combining maternal age, pre pregnancy, maternal medical and obstetric history and pregnancy biomarkers, would have yielded more results.

Cost effective analysis of HbA1c has not been done.

CONCLUSION

CONCLUSION

Gestational Diabetes mellitus has many complications in pregnancy, for both mother and baby. Few complications noted in mother are development of Pre-Eclampsia, increased incidence of macrosomia to baby, leading to increased operative interference, like instrumental delivery and lower segment caesarean section v shoulder dystocia leading to maternal injuries.

Fetal complexities incorporate respiratory trouble disorder, macrosomia, metabolic inconveniences like hypoglycemia, hypocalcemia, hypomagnesemia.

As rate and predominance of GDM are increasing in India, it is vital for early discovery and mediating with appropriate treatment. The ladies with GDM ought to be taught in regards to slim down, way of life change, and entanglements after conveyance. An itemized concentrate on hereditary parts and furthermore biomarkers assume a vital part in early expectation.

BIBLIOGRAPHY

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at the right end of the horizontal line. The horizontal line is slightly offset from the bottom of the page, and the vertical line is positioned to the right of the word 'BIBLIOGRAPHY'.

REFERENCES

1. Macintosh MC, Fleming KM, Bailey JA et al. Perinatal mortality and natural idiosyncrasies in offspring of women with type 1 or type 2 diabetes in England, Wales, and Northern Ireland: population based study. *BMJ* 2006;333:177.
2. Tennant PW, Glinianaia SV, Bilous RW, Rankin J, Toll R. Preexisting diabetes, maternal glycated hemoglobin, and the risks of fetal and child destruction: a population based study. *Diabetologia* 2014;57:285-294.
3. Dabelea D. The tendency to weight and diabetes in successors of diabetic mothers. *Diabetes Care* 2007;30 (Suppl 2):S169-S174.
4. HAPO Study Pleasant Investigation Social occasion. Hyperglycemia and negative pregnancy results. *N Engl J Solution* 2008;358:1991-2002.
5. Confidential Enquiry into Maternal and Youngster Prosperity. Pregnancy in Women with Type 1 and Type 2 Diabetes, 2002-2003. England, Wales and Northern Ireland. London: CEMACH, 2005
6. Diabetes care and investigation in Europe: the Blessed individual Vincent declaration. *Diabetes Solution* 1990;7:36
7. Ring R, Bailey K, Cresswell T, Hawthorne G, Critchley J, Lewis-Barned N. Designs in transcendence and consequences of pregnancy in women with pre-existing type I and type II diabetes. *BJOG* 2008;115:445–452.

-
- 8.Fadl HE, Simmons D. Designs in diabetes in pregnancy in Sweden 1998-2012. *BMJ Open Diabetes Res Care* 2016;4: e000221.
 - 9.Davenport MH, Campbell MK, Mottola MF. Extended recurrence of glucose issues during pregnancy isn't figured out by pre-pregnancy strength in London , Canada. *BMC Pregnancy Work* 2010 ; 10:85.
 - 10.Lawrence JM, Contreras R, Chen W, Sacks DA.Trends in the inescapability of earlier diabetes and gestational diabetes mellitus among a racially/ethnically unique people of pregnant women, 1999-20.05. *Diabetes Care* 2008;31:899-904.
 - 11.D. Keith Edmonds, Christoph Remains and Tom Bourne. Dewhurst's Perusing material of Obstetrics & Gynecology, tenth Delivery. Modified by 2018 John Wiley and Kids Ltd. Conveyed 2018 .
 12. Catalano PM, McIntyre HD, Cruickshank JK et al. The hyperglycemia and ominous pregnancy result study: relationship of GDM and chubbiness with pregnancy results. *Diabetes Care* 2012;35:780-786.
 13. Public Association for Prosperity and Care Significance. Diabetes in Pregnancy: The chiefs from Inclination to the Post pregnancy Time period. Lovely Bearing NG3. London: NICE, 2015. Available at nice.org.uk/heading/ng3.
 - 14.Plants JL. Irregularities in children of diabetic mothers. *Birth Defects Res A Clin Mol Teratol* 2010;88:769-778.

-
15. Tobias DK, Hu FB, Chavarro J, Rosner B, Mozaffarian D, Zhang C. Bracing dietary models and type 2 diabetes mellitus risk among women with a foundation set apart by gestational diabetes mellitus. *Bend Partner Medication* 2012;172:1566-1572.
 16. Mitanchez D, Yzydorczyk C, Siddeek B, Boubred F, Benahmed M, Simeoni U. The any kind of future family of the diabetic mother: short- and long term ideas. *Best Pract Res Clin Obstet Gynaecol* .2015;
 17. Gentle CL, Lewis HB, Patient C, Murphy HR, Simmons D. Assurance of gestational diabetes mellitus: falling through the net. *Diabetologia* 2015;58:2003-2012.
 18. Clausen TD, Mathiesen trauma center, Hansen T et al. Overweight and the metabolic condition in grown-up successors of women with diet-treated gestational diabetes mellitus or type 1 diabetes. *J Clin Endocrinol Metab* 2009;94:2464-2470
 19. Lo JC, Feigenbaum SL, Escobar GJ, Yang J, Crites YM, Ferrara A. Expanded routineness of gestational diabetes mellitus among ladies with separated polycystic ovary mix: a population-based study. *Diabetes Care* 2006;29:1915-1917,
 20. Weiss Father, Scholz HS, Haas J, Tamussino KF, Seissler J, Borkenstein MH. Long-term follow-up of offspring of moms with type 1 diabetes:

-
- confirmation for gained and non hereditary transmission of diabetes and ancestors. *Diabetes Care* 2000;23:905-911.
21. Guerin A, Nisenbaum R, Shaft JG. Utilization of maternal GHb fixation to review the bet of intrinsic eccentricities in the replacements of ladies with prepregnancy diabetes. *Diabetes Care* 2007;30:1920-1925.
22. World Success Association, "Coherent models and depiction of hyperglycaemia perceived in pregnancy: a World Flourishing Connection rule," *Diabetes Examination and Clinical Practice*, vol. 103, no. 3, pp. 341–363, 2014.
23. L. Mañé, J. A. Flores-le Roux, N. Gómez et al., " Relationship of firsttrimester HbA1c levels with contradicting pregnancy achieves different ethnic social events," *Diabetes Examination and Clinical Practice*, vol. 150, pp. 202–210, 2019.
24. E. Vounzoulaki, K. Khunti, S. C. Abner, B. K. Tan, M. J. Davies, and C. L. Gillies, "Improvement to type 2 diabetes in ladies with a known history of gestational diabetes: exact audit and metaanalysis," *BMJ: English Clinical Diary*, vol. 369, 2020.
25. B. Baz, J. P. Riveline, and J. F. Gautier, "Endocrinology of pregnancy: gestational diabetes mellitus: definition, aetiological and clinical focuses," *European Diary Of Endocrinology*, vol. 174, no. 2, pp. R43–R51, 2016

-
- 26.D. Benaiges, J. A. Flores-le Roux, I. Marcelo et al., " Is firsttrimester HbA1c obliging in the assessment of gestational .diabetes?," Diabetes Examination and Clinical Practice, vol. 133, pp. 85–91, 2017. [10] L. R. Nielsen, P. Ekb
- 27.L. R. Nielsen, P. Ekbom, P. Damm et al., " HbA1c levels are endlessly out lower in right on time and late pregnancy," Diabetes Care, vol. 27, no. 5, pp. 12001201, 2004
- 28,Sacks DA, Hadden DR, Maresh M, Deerochanawong C, Dyer AR, Metzger BE, et al. HAPO center around obliging .research pack. Rehash of gestational diabetes mellitus at working together sparkles considering IADPSG course of action board proposed rules: the hyperglycemia and awful pregnancy result (HAPO) study. Diabetes Care. 2012;35(3):526–8.
29. Dhulkotia JS, Ola B, Fraser R, Farrell T. Oral hypoglycemic experts versus insulin in relationship of gestational diabetes: a decided study and metaanalysis. Am J Obstet Gynecol 2010;203:457.e1-9.
30. Rowan JA, Hague WM, Gao W, Battin MR, Moore MP. Metformin versus insulin for the treatment of gestational diabetes. N Engl J Fix 2008;358:2003-2015.

-
31. Feig DS, Moses RG. Metformin treatment during pregnancy: incredible for the goose and really extraordinary for the gosling besides? *Diabetes Care* 2011;34:2329-2330.
 32. Gilbert C, Valois M, Koren G. Pregnancy result after first-trimester responsiveness to metformin: a meta-analysis. *Fertil Steril* 2006;86:658-663.
 33. Klingensmith GJ, Pyle L, Nadeau KJ et al. Pregnancy accomplishes youth with type 2 diabetes: the TODAY Study Understanding. *Diabetes Care* 2016;39:122-129.
 34. American Diabetes Association. (2) party and certification of diabetes. *Diabetes Care*. 2015;38(Suppl):S8–S16.
 35. Popova PV, Grineva EN, Gerasimov AS, Kravchuk EN, Ryazantseva EM, Shelepova ES. The new mix of chance factors picking a high bet of gestational diabetes mellitus. *Minerva Endocrinol*. 2015;40(4):239-47
Epub 2014 Oct 7
 36. Hughes RC, Rowan J, Florkowski CM. Is there a task for HbA1c in pregnancy? *Curr Diab. Rep*. 2016;16(1)5
 37. Politi S, D'emidio L, Cignini P, Giorlandino M, Giorlandino C. Shoulder dystocia: an evidence- based approach. *J Prenat Game plan* 2010;4:35-42.

-
38. Athukorala C, Crowther CA, Willson K. Women with gestational diabetes mellitus in the ACHOIS fundamental: risk factors for shoulder dystocia. *Aust NZ J Obstet Gynaecol* 2007;47:37-41.
 39. Roeder HA, Moore TR, Ramos GA. Changes in post pregnancy insulin necessities for patients with well-controlled type 1 diabetes. *Am J Perinatol* 2016;33:683-687
 40. Bellamy L, Casas JP, Hingorani Movement, Williams D. Type 2 diabetes mellitus after gestational diabetes: an exact survey and meta-analysis. *Lancet* 2009;373:1773-1779.
 41. Sharp AJ, Duncan E, McKillop-Smith A, Evans ND, Gold AE. Segment Change for Normal Eating (DAFNE) in routine clinical practice: who benefits? *Diabet Fix* 2012;29:670-676.
 42. Wahabi HA, Alzeidan RA, Bawazeer GA, Alansari LA, Esmaeil SA. Actually settled tendency idea for diabetic people for chipping away at maternal and fetal outcomes: a purposeful review and meta-analysis. *BMC Pregnancy Work* 2010;10:63.
 43. Crowther CA, Hiller JE, Vegetation JR, McPhee AJ, Jeffries WS, Robinson JS. Effect of treatment of gestational diabetes mellitus on pregnancy results. *N Engl J Solution* 2005;352:2477-2486.
 44. Mary C M Macintosh, Kate M Fleming, Jaron A Bailey et al. Perinatal mortality and regular fanciful notions in successors of women with Type I

-
- or Type II Diabetes in England, Grains, and Northern Ireland. *BMJ* 2006 July 22; 333(7560): 177.
45. Cooper WO, Hernandez-Diaz S, Arbogast PG et al. Significant normal mutilations after first-trimester responsiveness to Rule inhibitors. *N Engl J Solution* 2006;354:2443-2451.
46. Rohini HN, Punita P, Santhekadur PK, Ravishankar MV. Gestational Diabetes Mellitus - The Top tier Indian Perspective. *Indian J Endocrinol Metab.* 2023 Sep-Oct;27(5):387-393. doi: 10.4103/ijem.ijem_147_23. Epub 2023 Oct 30. PMID: 38107727; PMCID: PMC10723610 1
47. Mehrnaz Valadin, Zeinab Bahramnezhad, Elham Feizabad et al. The control of first trimester HbA1c in the early end GDM. *BMC Pregnancy and Work* Vol.22, Article No.77, 2022
48. Howard Berger, Robert Gagnon, Mathew Sermer et al. Diabetes in Pregnancy *Journal of Obstetrics and Gynecology Canada* 38(7) May 2016
49. American Diabetes Association. Classification and Assessment of Diabetes: Rules of Clinical Thought. *Diabetes care* 2016: 39(Suppl 1: S13-S22
50. Knight M, Kenyon S, Brocklehurst P, Neilson J, Shakespeare J, Kurinczuk JJ (eds) taking into account a genuine concern for MBRRACE- UK. Saving Lives, Further making Mothers'Care. Models sorted out a feasible strategy for enlightening future maternity care from

-
- the UK and Ireland Coordinated Enquiries into Maternal Passings and Dreariness 2009-12. Oxford: Public Perinatal The assessment of defilement transmission Unit, School of Oxford, 2014.
51. Diabetes care and Assessment in Europe: The Sanctified individual Vincent articulation. Diabet Medication 1990;7:360
52. Diabetes and pregnancy(MF Hivert ,portion supervisor), voi 17,article number 12,(2017) Early signs of Gestational Diabetes Mellitus.
53. Junnei Wang ,Li Zhang,Yubai et al Diaetes 2023 jan. The effect of more restricted red platelet life length on the speed of HbA1c target achieved in type 2 Diabetes patients with HbA1c assertion regard lower than 7%.
54. Pavitra R,Sreelatha S,Pooja p et al As a rule Of Reproduction,Contraception,Obstetrics and Gynaecology,vol.12.no 2 (2023) .
55. Madleen Lemaitre, Camille Ternynck, Julien Bourry, Florence Baudoux, Damien Subtil, and Anne Vambergue .J Clin Endocrinol Metab. 2022 Mar; (3107e1117):- .
56. Conveyed e1125online 2021 Oct Relationship Between HbA1c Levels on Poorly arranged Pregnancy Results During Pregnancy in Patients With Type 1 Diabetes
57. Sofia Amylidi, Beatrice Mosimann, Christoph Stettler, Georg Martin Fiedler, Daniel Surbek, Luigi Raio.

Marie Parfaite uwimana muhuza et al. Front Endocrinol (Lausanne) 2023 The Relationship between maternal HbA1c and awful outcomes in gestational diabetes.

58. Latife Bozkurt , Christian S. Gobi , Lisa Pfigl. et al. Pathophysiological characteristics and effects of enormity in women with early and late appearances of Gestational diabetes isolated by the general relationship of Diabetes and pregnancy base on Gatherings rules.

The journal of clinical Endocrinology and Metabolism, volume 100, Issue 3, 1 walk 2015, pages 1113-1120.

59. Bashir M, E Abdel - Rahman M, Aboufotouh M, Eltaher F, Omar K et al Undeniability of really seen diabetes in pregnancy in Qatar, using general screening. PLoS ONE 13(8) : e 0201247

60. J Balani . S L Hyer, H Shehata , F Mohareb Expecting gestational diabetes mellitus by first trimester HbA1c ; a review revolve around in women with moderate to over the top power. Sensible DIABETES 31 JULY 2023.

61. XiaoXiao peng et al. Curve Gynecol obstet. 2023 may Usage of oral glucose versatility testing and HbA1c at 6-14 gestational weeks to expect gestational diabetes mellitus in high - risk women.

62. Rajesh Rajput , Yogesh Yadav RM , Nanda S. et al . Diabetes res clin pract . 2012. oct Utility of HbA1c for finish of gestational diabetes mellitus.

-
63. Sarah S. Osmundson ,Benain S.Zhao. Liza Kunz et al. First Trimester HemoglobinA1c notion for gestational diabetes mellitus. Am J perinatal 2016; 33 (10): 977-982
64. Soon sung kwon et Ja - young kwon,yong - won park ,young - han kim e al..Diabetes Res Clin pract. 2015 Oct; 110 (1) : 38-43
- 65.Ribal kattini ,Ruben Hummelen MD,Ph D,Len Kelly MD, MClinsci Early gestational diabetes mellitus screening with Glycated Hemoglobin; A Purposeful layout. Journal of obstetrics and gynecology Canada volume 42,issue 11,November 2020,pages 1379-1384.
- 66.Hatice Kansu - celik ,A. seval ozgu - Erdnic et al. The Journal of maternal - fetal &Neonatal drug volume 34,2021-issue 12 Maternal serum glycosylated hemoglobin and fasting plasma glucose predicts gestational diabetes at the fundamental trimester in a long time with a low - risk pregnancy and its relationship with fetal birthweight : an outline buddy study.
- 67.Paula Breitenbach Renz,Gabriela Cavagnolli ,Leticia Schwerz Weinert , Sandra Pinho Silveiro et al.
- HbA1c test asa contraption in the finding of Gestational Diabetes Mellitus PLoS One.2015; 10 (8): e0135989.2015 Aug 20.

-
- 68.Cihan cetin et al. Nur Dokuzeylul Gungor,Melike Yavuz First trimester glycosylated hemoglobin for gestational diabetes mellitus screening,Taiwan J obstet Gynecol.2021
- 69.Claire Beynon .Hiller Sharon etal journal of general flourishing 42 february 2020 Should HbA1C be used to study pregnant individuals for disguised diabetes in the first triester/A Review of the attestation.
70. Text Book Williams Obstetrics 24th Edition Page no.1160-1170.”**on**

ANNEXURE

A decorative graphic consisting of a thick horizontal black line and a thick vertical black line intersecting at a right angle. The horizontal line extends from the left edge of the page towards the right, and the vertical line extends from the bottom edge of the page upwards. The intersection point is located to the right of the word 'ANNEXURE'.

PROFORMA

NAME:

AGE:

ADDRESS:

UHID NO:

I.P NO:

DATE/ TIME OF ADMISSION: DATE/ TIME OF DISCHARGE:

CHIEF COMPLAINTS:

OBSTETRICAL HISTORY: BOOKED/ UNBOOKED/ REFERRED

MARRIED LIFE: CONSANGUINOUS MARRIAGE: YES/ NO
OBSTETRICAL SCORE: : ANC : REGULAR / IRREGULAR

NO OF VISITS :

INJ .TT DOSES: IRON AND FOLIC ACID:

MENSTRUAL HISTORY: LMP: EDD: POG:

PAST HISTORY:

FAMILY HISTORY:

PERSONAL HISTORY: TREATMENT HISTORY :

INVESTIGATIONS: 1) LEVELS OF HBA1C AND FASTING BLOOD
SUGAR DURING FIRST TRIMESTER :

FASTING BLOOD SUGAR : HBA1C :

2) DURING 24-28 WKS :

FBS: HBA1C: HB%: BLOOD GROUP AND RHTYPE:

GENERAL PHYSICAL EXAMINATION:

PALLOR/ ICTERUS/ CYANOSIS/ CLUBBING/ LYMPHADENOPATHY/
EDEMA

PULSE:

BP:

RR:

TEMP:

CNS: CVS: RS:

P/A:

PROVISIONAL DIAGNOSIS:

INFORMED CONSENT FORM

I Mr./Mrs. _____ have been explained in my own understandable language, that I will be included in a study which is “THE ROLE OF HBA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY- A PROSPECTIVE STUDY.

I have been explained that my clinical findings, investigations, will be assessed and documented for study purpose.

I have been explained my participation in this study is entirely voluntary, and I can withdraw from the study any time and this will not affect my relation with my doctor or the treatment for my ailment.

I have been explained about the interventions needed possible benefits and adversities due to interventions, in my own understandable language.

I have understood that all my details found during the study are kept confidential and while publishing or sharing of the findings, my details will be masked.

I have principal investigator mobile number for enquiries.

I in my sound mind give full consent to be added in the part of this study.

Signature of the patient:

Name:

Date:

Place:

signature of witness

Relation to patient:

ನೀಡಿದ ಒಪ್ಪಿಗೆ ಮಾಹಿತಿನಮೂನೆ

ನಾನು ಶ್ರೀ/ಶ್ರೀಮತಿ _____ ನನ್ನದೇ ಆದ ಅರ್ಥವಾಗುವ ಭಾಷೆಯಲ್ಲಿ ವಿವರಿಸಲಾಗಿದೆ, ಇದು ಒಂದು ಅಧ್ಯಯನದಲ್ಲಿ ನನ್ನನ್ನು ಸೇರಿಸಿಕೊಳ್ಳಲಾಗುವುದು ಅದು "ಪೂರ್ವಭಾವಿ ಗರ್ಭಾವಸ್ಥೆಯಲ್ಲಿ ಗರ್ಭಾವಸ್ಥೆಯ ಮಧುಮೇಹ ಮೆಲ್ಲಿಟಸ್‌ನ ಆರಂಭಿಕ ಮುನ್ನೂಚಕವಾಗಿ HBA1C ಪಾತ್ರ - ಒಂದು ನಿರೀಕ್ಷಿತ ಅಧ್ಯಯನ.

ನನ್ನ ಕ್ಲಿನಿಕಲ್ ಸಂಶೋಧನೆಗಳು, ತನಿಖೆಗಳು, ಶಸ್ತ್ರಚಿಕಿತ್ಸೆಯ ನಂತರದ ಸಂಶೋಧನೆಗಳನ್ನು ಮೌಲ್ಯಮಾಪನ ಮಾಡಲಾಗುತ್ತದೆ ಮತ್ತು ಅಧ್ಯಯನ ಉದ್ದೇಶಕ್ಕಾಗಿ ದಾಖಲಿಸಲಾಗುತ್ತದೆ ಎಂದು ನನಗೆ ವಿವರಿಸಲಾಗಿದೆ.

ಈ ಅಧ್ಯಯನದಲ್ಲಿ ನನ್ನ ಭಾಗವಹಿಸುವಿಕೆಯು ಸಂಪೂರ್ಣವಾಗಿ ಸ್ವಯಂಪ್ರೇರಿತವಾಗಿದೆ ಎಂದು ನನಗೆ ವಿವರಿಸಲಾಗಿದೆ ಮತ್ತು ನಾನು ಯಾವುದೇ ಸಮಯದಲ್ಲಿ ಅಧ್ಯಯನದಿಂದ ಹಿಂದೆ ಸರಿಯಬಹುದು ಮತ್ತು ಇದು ನನ್ನ ವೈದ್ಯರೊಂದಿಗಿನ ನನ್ನ ಸಂಬಂಧ ಅಥವಾ ನನ್ನ ಕಾಯಿಲೆಯ ಚಿಕಿತ್ಸೆಯ ಮೇಲೆ ಪರಿಣಾಮ ಬೀರುವುದಿಲ್ಲ.

ನನ್ನ ಸ್ವಂತ ಅರ್ಥವಾಗುವ ಭಾಷೆಯಲ್ಲಿ ಮಧ್ಯಸ್ಥಿಕೆಗಳಿಂದಾಗಬಹುದಾದ ಪ್ರಯೋಜನಗಳು ಮತ್ತು ಪ್ರತಿಕೂಲತೆಗಳ ಅಗತ್ಯವಿರುವ ಮಧ್ಯಸ್ಥಿಕೆಗಳ ಬಗ್ಗೆ ನನಗೆ ವಿವರಿಸಲಾಗಿದೆ.

ಅಧ್ಯಯನದ ಸಮಯದಲ್ಲಿ ಪತ್ತೆಯಾದ ನನ್ನ ಎಲ್ಲಾ ವಿವರಗಳನ್ನು ಗೌಪ್ಯವಾಗಿ ಇರಿಸಲಾಗಿದೆ ಮತ್ತು ಸಂಶೋಧನೆಗಳನ್ನು ಪ್ರಕಟಿಸುವಾಗ ಅಥವಾ ಹಂಚಿಕೊಳ್ಳುವಾಗ, ನನ್ನ ವಿವರಗಳನ್ನು ಮರೆಮಾಚಲಾಗುತ್ತದೆ ಎಂದು ನಾನು ಅರ್ಥಮಾಡಿಕೊಂಡಿದ್ದೇನೆ.

ವಿಚಾರಣೆಗಾಗಿ ನಾನು ಪ್ರಧಾನ ತನಿಖಾಧಿಕಾರಿಯ ಮೊಬೈಲ್ ಸಂಖ್ಯೆಯನ್ನು ಹೊಂದಿದ್ದೇನೆ.

ಈ ಅಧ್ಯಯನದ ಭಾಗದಲ್ಲಿ ಸೇರಿಸಲು ನನ್ನ ಉತ್ತಮ ಮನಸ್ಸಿನಲ್ಲಿ ನಾನು ಸಂಪೂರ್ಣ ಒಪ್ಪಿಗೆಯನ್ನು ನೀಡುತ್ತೇನೆ. ರೋಗಿಯ ಸಹಿ ಸಾಕ್ಷಿಯ ಸಹಿ

ಹೆಸರು:

ದಿನಾಂಕ:

ಹೆಸರು

ರೋಗಿಗೆ ಸಂಬಂಧ

PATIENT INFORMATION SHEET

STUDY TITLE: THE ROLE OF HBA1C AS AN EARLY PREDICTOR OF GESTATIONAL DIABETES MELLITUS IN EARLY PREGNANCY-

A PROSPECTIVE STUDY.

Department of OBG

STUDY SITE: R.L Jalappa Hospital and Research Centre, Tamaka, Kolar.

Please read the following information and discuss with your family members. You can ask any question regarding the study.

If you agree to participate in the study we will collect information (as per proforma) from you or from a person responsible for you or both. Relevant history will

be taken. This information collected will be used only for dissertation and publication.

Blood samples will be collected between 12-14 weeks HBA1C and 24-28 weeks for gestational age for HBA1C to assest the risk of GDM in the Present pregnancy.

The relevant investigations which are required others than regular investigations will

be funded by me. All information collected from you will be kept confidential and will not be disclosed to any outsider. Your identity will not be revealed.

This study has been reviewed by the Institutional Ethics Committee .

There is no compulsion to agree to this study. The care you will get will not change if you don't wish to participate. You are required to sign/ provide thumb impression only if you voluntarily agree to participate in this study.

You are free to contact Dr. SOMAVARAPU DIVYA or any other member of the above research team for any doubt or clarification you have.

Dr. SOMAVARAPU DIVYA

Mobile no: 7680887208

E-mail id: drdivyarao1994@gmail.com

ರೋಗಿಯ ಮಾಹಿತಿ ಹಾಳೆ

ಅಧ್ಯಯನದ ಶೀರ್ಷಿಕೆ: ಆರಂಭಿಕ ಗರ್ಭಾವಸ್ಥೆಯಲ್ಲಿ ಗರ್ಭಾವಸ್ಥೆಯ ಮಧುಮೇಹ ಮೆಲ್ಲಿಟಸ್‌ನ ಮುಂಚಿನ ಪೂರ್ವಭಾವಿಯಾಗಿ HBA1C ಯ ಪಾತ್ರ-ಒಂದು ಪ್ರಾಸ್ಪೆಕ್ಟಿವ್ ಸ್ಟಡಿ.

ಸ್ವಡಿ ಸ್ಟೈಟ್: ಆರ್.ಎಲ್ ಜಾಲಪ್ಪ ಆಸ್ಪತ್ರೆ ಮತ್ತು ಸಂಶೋಧನಾ ಕೇಂದ್ರ, ಟಮಕ, ಕೋಲಾರ.

ದಯವಿಟ್ಟು ಕೆಳಗಿನ ಮಾಹಿತಿಯನ್ನು ಓದಿ ಮತ್ತು ನಿಮ್ಮ ಕುಟುಂಬದ ಸದಸ್ಯರೊಂದಿಗೆ ಚರ್ಚಿಸಿ. ಅಧ್ಯಯನಕ್ಕೆ ಸಂಬಂಧಿಸಿದಂತೆ ನೀವು ಯಾವುದೇ ಪ್ರಶ್ನೆಯನ್ನು ಕೇಳಬಹುದು. ನೀವು ಅಧ್ಯಯನದಲ್ಲಿ ಭಾಗವಹಿಸಲು ಸಮ್ಮತಿಸಿದರೆ ನಾವು ನಿಮ್ಮಿಂದ ಅಥವಾ ನಿಮ್ಮಿಂದ ಅಥವಾ ಇಬ್ಬರಿಗೂ ಜವಾಬ್ದಾರರಾಗಿರುವ ವ್ಯಕ್ತಿಯಿಂದ ಮಾಹಿತಿಯನ್ನು (ಪ್ರೌಢಾರ್ಥ ಪ್ರಕಾರ) ಸಂಗ್ರಹಿಸುತ್ತೇವೆ. ಸಂಬಂಧಿತ ಇತಿಹಾಸ ಕಾಣಿಸುತ್ತದೆ ತೆಗೆದುಕೊಳ್ಳಲಾಗುವುದು. ಸಂಗ್ರಹಿಸಿದ ಈ ಮಾಹಿತಿಯನ್ನು ಪ್ರಬಂಧ ಮತ್ತು ಪ್ರಕಟಣೆಗೆ ಮಾತ್ರ ಬಳಸಲಾಗುತ್ತದೆ. ನಿಯಮಿತ ತನಿಖೆಗಳಿಗಿಂತ ಇತರ ಅಗತ್ಯವಿರುವ ಸಂಬಂಧಿತ ತನಿಖೆಗಳು ನನ್ನಿಂದ ಧನಸಹಾಯ ಮಾಡಿ. ನಿಮ್ಮಿಂದ ಸಂಗ್ರಹಿಸಲಾದ ಎಲ್ಲಾ ಮಾಹಿತಿಯನ್ನು ಗೌಪ್ಯವಾಗಿ ಇರಿಸಲಾಗುತ್ತದೆ ಮತ್ತು ಯಾವುದೇ ಹೊರಗಿನವರಿಗೆ ಬಹಿರಂಗಪಡಿಸಲಾಗುವುದಿಲ್ಲ. ನಿಮ್ಮ ಗುರುತನ್ನು ಬಹಿರಂಗಪಡಿಸಲಾಗುವುದಿಲ್ಲ. ಈ ಅಧ್ಯಯನವನ್ನು ಸಾಂಸ್ಥಿಕ ನೀತಿಶಾಸ್ತ್ರ ಸಮಿತಿಯು ಪರಿಶೀಲಿಸಿದೆ ಮತ್ತು ನೀವು ಸಾಂಸ್ಥಿಕ ನೀತಿಶಾಸ್ತ್ರ ಸಮಿತಿಯ ಸದಸ್ಯರನ್ನು ಸಂಪರ್ಕಿಸಲು ಮುಕ್ತರಾಗಿದ್ದೀರಿ. ಈ ಅಧ್ಯಯನವನ್ನು ಒಪ್ಪಿಕೊಳ್ಳಲು ಯಾವುದೇ ಒತ್ತಾಯವಿಲ್ಲ. ನೀವು ಭಾಗವಹಿಸಲು ಬಯಸದಿದ್ದರೆ ನೀವು ಪಡೆಯುವ ಕಾಳಜಿಯು ಬದಲಾಗುವುದಿಲ್ಲ. ಈ ಅಧ್ಯಯನದಲ್ಲಿ ಭಾಗವಹಿಸಲು ನೀವು ಸ್ವಯಂಪ್ರೇರಣೆಯಿಂದ ಸಮ್ಮತಿಸಿದರೆ ಮಾತ್ರ ನೀವು ಸಹಿ/ಹೆಬ್ಬರಳಿನ ಗುರುತನ್ನು ಒದಗಿಸಬೇಕಾಗುತ್ತದೆ. ನಿಮ್ಮಲ್ಲಿರುವ ಯಾವುದೇ ಸಂದೇಹ ಅಥವಾ ಸ್ಪಷ್ಟೀಕರಣಕ್ಕಾಗಿ ಡಾ. ಸೋಮವರಪು ದಿವ್ಯಾ ಅಥವಾ ಮೇಲಿನ ಸಂಶೋಧನಾ ತಂಡದ ಇತರ ಸದಸ್ಯರನ್ನು ಸಂಪರ್ಕಿಸಲು ನೀವು ಮುಕ್ತರಾಗಿದ್ದೀರಿ.

ಡಾ.ಸೋಮವರಪು ದಿವ್ಯಾ

ಮೊಬೈಲ್ ಸಂಖ್ಯೆ: 7680887208

ಇ-ಮೇಲ್ ಐಡಿ: drdivyaraol1994@gmail.com

MASTER CHART

A decorative graphic consisting of a thick horizontal line and a thick vertical line intersecting at a right angle. The intersection is located to the right of the text 'MASTER CHART'. The lines are black with a slight gray shadow or offset, giving them a three-dimensional appearance.

ANNEXURE III

KEY TO MASTER CHART

KEY	AGE	NO			PERCENTAGE		
A	<25	40			53		
	25-35	32			42		
	>35	3			4		
B	Distribution of study population with reference to HbA1c values	N	ABNORMAL		76 24		
		57	18 0				
C	Distribution of patients with reference to HbA1c values	6	10	2	8	14.28	2.8
D	Age wise distribution of sample with reference to glycemic (HbA1c) control	NO	<6.5	6.5-7.9	40 30 3		
		<25	37	10			
		26-35	22	2			
		<35	1	15			
E	Incidence of GDM in different age groups.						
		<25	25-35	>35			
		3	12	3			
F	Parity Distribution of Total sample tested		63				
	PRIMI	MULTI					
	46	29	38				
G	Parity distribution with reference glycemic control		PRIMI		MULTI		
			<6.5		6.5-7.9		
			53		8		
			7		7		

H	Incidence of GDM with reference to parity	<3	>3	
		61 14.28%	14 24%	
I	Abnormal HbA1c correlates with excessive weight gain Abnormal HbA1c-	P	A	
		18 100%	0 0%	
J	Glycemic control basing on Booked/Un booked status	UNBOOKED	BOOKED	
		36	24	
K	Glycemic control basing on social status	36	9	
		24	6	
L	HbA1c distribution with oral glucose tolerance test at 24-28 weeks			
	1st TRIMISTER		2 6- 28 WKS	100%
	NORMAL	ABNORMAL	ABNORMAL	
	57	18	18	

	ROC CURVE IN FIRST TRIMISTER IN PREDICTING GDM	P value-0.0001
		Area under the curve=1

MASTER CHART

SNO	Age	B/UB	Social status	Parity	HbA1c	OGTT
1	25	B	LM	1	6.4	200
2	26	UB	LM	3	7.2	220
3	24	UB	LM	1	6.5	180
4	30	B	LM	3	6.4	188
5	22	B	M	1	5.4	139
6	28	UB	LM	3	7.4	180
7	32	B	LM	1	5	126
8	20	B	M	1	5.1	135
9	22	UB	M	1	4.6	111
10	19	B	LM	1	4.8	112
11	25	UB	M	2	5	136
12	20	B	LM	1	5.2	125
13	24	B	LM	2	5.4	128
14	36	UB	LM	4	5.8	234
15	22	UB	M	1	5	130
16	20	UB	M	1	4.7	110
17	21	B	LM	2	5.1	131
18	23	B	M	1	5.3	138
19	24	B	LM	3	5.4	136
20	22	UB	LM	1	5	130
21	36	B	M	4	7.4	262
22	38	UB	LM	3	7.1	192
23	28	B	LM	3	7.6	194
24	27	B	LM	2	4.6	110
25	22	UB	M	1	4.8	115
26	20	UB	M	1	5	128
27	22	B	LM	1	4.9	126
28	26	UB	LM	2	5.3	135
29	30	B	LM	3	7.1	188
30	21	B	M	1	4.9	115
31	20	UB	LM	1	5.3	138
32	21	UB	LM	1	5.2	129
33	24	B	M	1	5	120
34	28	UB	M	2	7.2	178
35	20	B	LM	1	5.1	132
36	25	UB	LM	2	5.4	130
37	28	UB	LM	2	5.3	136
38	25	B	LM	1	5.2	129
39	21	B	LM	1	4.9	120
40	20	B	M	1	4.6	111

SNO	Age	B/UB	Social status	Parity	HbA1c	OGTT
41	22	UB	M	1	4.8	100
42	21	B	M	1	5	130
43	29	UB	LM	3	6.8	182
44	24	B	LM	1	6.9	220
45	20	B	M	1	5	132
46	23	B	LM	1	5.1	131
47	20	UB	M	1	5.2	136
48	28	UB	M	3	4.9	123
49	22	UB	LM	1	4.6	120
50	30	UB	LM	3	4.9	126
51	24	B	M	2	4.5	122
52	25	B	M	2	4.2	120
53	24	UB	LM	1	4	98
54	26	B	LM	2	5	130
55	29	B	M	3	5.1	133
56	23	UB	LM	1	5.3	136
57	24	UB	M	1	7.1	198
58	22	B	M	1	5.2	132
59	20	UB	LM	1	5	134
60	26	B	M	2	4.5	120
61	22	B	M	1	4.9	122
62	25	UB	LM	1	4.6	128
63	26	B	LM	2	5	130
64	22	UB	LM	1	5.1	132
65	26	B	LM	2	5.2	128
66	23	UB	LM	1	5.4	134
67	26	B	LM	1	4.5	120
68	27	UB	M	1	6.7	188
69	28	B	M	1	6.5	184
70	26	UB	LM	1	6.8	200
71	28	B	LM	1	5	132
72	27	UB	M	1	7.2	208
73	22	B	LM	1	5.1	136
74	28	B	LM	2	5.3	132
75	26	B	M	3	5.2	130