DISSERTATION ON A PRE-EXPERIMENTAL STUDY TO EVALUATE THE EFFECTIVENESS OF STRUCTURED TEACHING PROGRAMME ON KNOWLEDGE REGARDING EARLY IDENTIFICATION OF HIGH RISK PREGNANCY AMONG ANTENATAL MOTHERS ATTENDING AT INSTITUTE OF OBSTETRICS AND GYNAECOLOGY, CHENNAI-08.

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In Partial Fulfillment of the Requirement for the Award of

DEGREE OF MASTER OF SCIENCE IN NURSING

OCTOBER – 2017

CERTIFICATE

This is to certify that this dissertation titled **A Pre-Experimental** study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High **Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08** is a bonafide work done by Mrs.J.Dhanalakshmi, M.Sc (N) II year student, College of Nursing, Madras Medical College, Chennai submitted to The Tamilnadu Dr.M.G.R Medical University, Chennai-32 towards the partial fulfillment of the requirements for the award of degree of Master of Science in Nursing, Branch – III, Obstetrics and Gynecological Nursing, under our guidance and supervision during the academic year 2015 -2017.

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"A PRE-EXPERIMENTAL STUDY TO EVALUATE THE EFFECTIVENESS OF STRUCTURED TEACHING PROGRAMME ON KNOWLEDGE REGARDING EARLY IDENTIFICATION OF HIGH RISK PREGNANCY AMONG ANTENATAL MOTHERS ATTENDING AT INSTITUTE OF OBSTETRICS AND GYNAECOLOGY,CHENNAI-08".

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ABSTRACT

Title:

A Pre-experimental study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08.

Antenatal period is amazing experience of each and every women. Even though this period is going smooth for many mothers some of them may suffer with some high risk during pregnancy may endanger the life of the mother and fetus. The main aim of the study particularly this group of antenatal mother is early identification of high risk pregnancy, prevention of complications , and management of these conditions.

Need for the study

The researcher had seen various kinds or pregnancy associated complications in obstetric department. The women who are all attending the antenatal outpatient department do not have an adequate knowledge regarding complications of high risk pregnancy. The researcher felt that to increase the knowledge and awareness about high risk pregnancy and selected this topic.

Objectives

- 1) The assess the knowledge regarding high risk pregnancy among antenatal women
- To assess the effectiveness of structured teaching programme on knowledge regarding high risk pregnancy among antenatal women.

3) To associate the findings with selected demographic variables and obstetrical variables.

Key words

High risk pregnancy, Structured Teaching Programme, Identification of high risk pregnancy.

Methodology

Research approach	Quantitative research approach		
Duration of the study	Four weeks (20.11.2016 to 18.12.2016)		
Study setting	Antenatal OPD at IOG		
Research design	Pre- experimental design		
Study Population	Antenatal mothers with gestational age 8-28 weeks.		
Sampling Technique	Purposive Sampling Technique		
Sample size	60		

Data collection procedure

After obtaining informed and written consent, approximately three to five samples were selected every day and pretest questionnaire was assessed by means of interview method. Structured Teaching Programme was given to the participants for 30 minutes. After pre-test, post -test was conducted after one week to assess the knowledge of women regarding early identification of high risk pregnancy.

Data analysis

The data were tabulated and analyzed using descriptive statistics like mean, standard deviation, frequency distribution and percentage.

Inferential statistics like paired t-test, McNemars test and chi-square test.

Results

The findings of the study revealed that the Structured Teaching Programme had improved the knowledge of women regarding early identification with paired t –test P value is 0.001. There is statistical significance in knowledge attainment on early identification of high risk pregnancy which shows the effectiveness of Structured teaching programme.

Discussion

Hypothesis was proved by the great statistically significance occurs after Structured Teaching Programme. The chi square test shows that there is a statistically significant association between the post- test level of knowledge and demographic variables among women.

Recommendation

A comparative study can be conducted between the knowledge of the patients with high risk pregnancy and their practices.

A follow up study can be recorded to determine the participation in screening practice for the prevention and early detection of high risk pregnancy.

Conclusion

The result of the study shows that the Structured Teaching Programme was effective in improving the knowledge regarding early identification of high risk pregnancy among antenatal mothers.

INDEX

CHAPTER NO.	TITLE	PAGE NO.
Ι	INTRODUCTION	1
	1.1 Need for the study	3
	1.2 Statement of the problem	6
	1.3 Objectives of the study	6
	1.4 Operational definitions	7
	1.5 Assumptions	7
	1.6 Research Hypothesis	8
	1.7 Delimitations	8
II	REVIEW OF LITERATURE	9
	2.1. Literature review related to study	9
	2.2 Conceptual framework	23
III	RESEARCH METHODOLOGY	25
	3.1 Research approach	25
	3.2 Study design	25
	3.3 Setting of the study	26
	3.4 Duration of the study	26
	3.5 Study Population	27
	3.5.1 Target Population	27
	3.5.2 Accessible Population	27
	3.6 Sample	27
	3.7 Sampling Criterion	27
	3.7.1 Inclusion criteria	28
	3.7.2 Exclusion criteria	28
	3.8 Sample size	28
	3.9 Sampling technique	28
	3.10 Research Variables	28
	3.10.1 Independent Variable	28
	3.10.2 Dependent Variable	29
	3.10.3 Influencing Variable	29

CHAPTER NO.	TITLE	PAGE NO.
	3.11 Development and description of the tool	30
	3.11.1Development of the tool	30
	3.11.2Description of the tool	30
	3.11.3Teaching Module	31
	3.12. Content Validity	31
	3.13 Protection of Human Subjects	32
	3.14 Reliability of the tool	32
	3.15 Pilot study	32
	3.16 Data collection procedure	33
	3.17 Intervention protocol	34
	3.18 Data entry and analysis	34
IV	DATA ANALYSIS AND INTERPRETATION	36
V	SUMMARY OF RESULTS	54
VI	DISCUSSION	59
VII	CONCLUSION AND RECOMMENDATIONS	66
	7.1 Implications of the study	66
	7.2 Limitations	68
	7.3 Recommendations for further study	68
	REFERENCES	
	APPENDICES	

LIST OF TABLES

TABLE NO	TITLE	PAGE.NO		
1.1	Maternal Death at Institute of Obstetrics and Gynaecology	6		
4.1	Demographic and Obstetrical profile in the study group.	38		
4.2	Percentage of pretest knowledge score in the study group	40		
4.3	Pretest level of knowledge score in the study group	40		
4.4	Percentage of posttest knowledge score in the study group	41		
4.5	Posttest level of knowledge score in the study group	41		
4.6	Comparison of pre and posttest knowledge score on high risk pregnancy	42		
4.7	Comparison of overall pre and posttest knowledge score on high risk pregnancy	44		
4.8	Pretest and posttest level of knowledge score in the study group	45		
4.9	Percentage of knowledge gain score on high risk pregnancy	46		
4.10	Effectiveness of structured teaching programme on high risk pregnancy	47		
4.11	Association between pretest level of knowledge score with demographic and obstetrical variables	48		
4.12	Association between posttest level of knowledge score with demographic and obstetrical variables.	50		
4.13	3 Association between post-test level of knowledge score with demographic and obstetrical variables			

LIST OF FIGURES

FIGURE NO	TITLE			
1.1	Maternal Mortality Ratio			
1.2	Time trend in Maternal Mortality Ratio-Tamilnadu			
2.3	Modified Pender's Health promotion model.			
3.4	Schematic representation of the variables			
3.5	Schematic representation of the study			
4.6	Age wise distribution of antenatal mothers			
4.7	Religion wise distribution of antenatal mothers.			
4.8	Type of family system of antenatal mothers.			
4.9	Educational status of antenatal mothers.			
4.10	Occupational status of the antenatal mothers			
4.11	Monthly income status of antenatal mothers.			
4.12	Age at marriage of antenatal mothers.			
4.13	Gravida wise distribution of antenatal mothers.			
4.14	Pretest level of knowledge score of antenatal mothers.			
4.15	Posttest level of knowledge score of antenatal mothers.			
4.16	Mean pretest and posttest knowledge score of antenatal mothers.			
4.17	Pretest and posttest level of knowledge score.			
4.18	Pre and posttest percentage of knowledge score of antenatal mothers.			
4.19	Percentage of knowledge gain.			
4.20	Association between posttest level of knowledge and antenatal women age.			
4.21	Association between posttest level of knowledge and antenatal women type of family system.			
4.22	Association between posttest level of knowledge score and antenatal women gravida.			
4.23	Association between knowledge gain score and demographical variables.			

LIST OF APPENDICES

S. NO	DESCRIPTION			
1.	Certificate approval by Institutional Ethics Committee			
2.	Certificate of content validity by experts			
3.	Letter seeking permission to conduct the study			
4.	Study tool			
	Section A – Demographic Information			
	Section B – Obstetric data			
	Section C– Structured questionnaire on anemia, pre- eclampsia, Gestational disbetes mellitus and multiple pregnancy.			
	Scoring key			
	Lesson plan			
4.	Informed consent			
5.	Coding sheet			
6.	Certificate for English Editing			
7.	Certificate for Tamil Editing			

ABBREVIATION

WHO	World Health Organization				
MMR	Maternal Mortality Rate				
OPD	Outpatient Department				
IOG	Institute of Obstetrics and Gynaecology				
STP	Structured Teaching Programme				
GDM	Gestational Diabetes Mellitus				
RCT	Randomized controlled trial				
SSC	Skin to Skin Contact				
MNMR	Maternal and Neonatal Mortality Rate				
RBC	Red Blood Cells				
PUFA	Poly Unsaturated Fatty Acid				
DIC	Disseminated Intravascular Coagulation				
NTD	Neural tube Defects				
USA	United States of America				
НАРО	Hyperglycemia and Adverse Pregnancy Outcome				
OGTT	Oral Glucose Tolerance Test				
ICU	Intensive Care Unit				
NICU	Neonatal Intensive Care Unit				
ART	Assisted Reproductive Technology				
BF	Breast Feeding				

CHAPTER –I INTRODUCTION

A women is the full circle. Within her is the power to create, nurture and transform.

-Diane Mariechild.

Women form the centre of the family and their health is of prime importance to the well-being of the whole family. With the emergence of nuclear family type, health of the women has increased in its relevance. Moreover, women's health is of cardinal importance to the health of the society.

It implies that if the mother is healthy, then the child will also be healthy. "Child health" is based on the "Mother's health".

"A happy child is a Nation's pride" and every child will be happy with his or her mother's bonding and family care. Health is wealth of the family. Preserving this health during the antenatal period becomes quite imperative to guard the mother from ill health.

Pregnancy is a unique experience in every woman's life. The thought of growing fetus in the mother's womb indeed is nature's way of expressing the attributes of motherhood. Most of the pregnant mothers suffer with many high risk conditions in their journey towards motherhood.

Identification of risk factors related to pregnancy is important because "A pregnant woman is like a ship on stormy sea, out of balance" seeking a new equilibrium in waves of physical and physiological change. Therefore nurses as women, midwives and health care providers have special responsibility to reduce the annual loss of life through early identification of high risk pregnancy related to maternal mortality and morbidity. The prevalence of medical problems in pregnancy is increasing because of a complex interplay between demographic and lifestyle factors, and developments in modern medicine. Maternal mortality and morbidity resulting from treatable medical conditions, such as anemia, gestational diabetes mellitus and hypertensive disorders have not decreased in recent years. It is vital that the health care professionals must acquire a basic knowledge and understanding of medical problems in pregnancy. This includes pre pregnancy measures such as counseling and optimization of medical therapy, as well as multidisciplinary management throughout the pregnancy and the postpartum period. (Narayan B, Nelson-Piercy C, 2016)¹

During pregnancy, a woman's body undergoes complex physiological changes, still which may not be understood. Every system in a woman's body adapts to the demand of the growing fetus. A great deal of attention during pregnancy is focused on ensuring minimum risk at delivery and maximum health of the women and her fetus (**Cooper, 2012**)²

Prenatal care is essential for ensuring the overall health of the mother and the infant. It is a major strategy for helping to reduce complications of pregnancy. Prenatal care should include collection of obstetric history, measuring height and weight, taking blood pressure to check preclampsia, screening for anemia and giving education regarding complications during pregnancy. So that the signs of complications can be identified and also measures can be taken to modify, control and reduce and high risk pregnancy (**Barley.E.Rose, 2010**)³

Several organizations have mentioned a large number of high risk maternal conditions which need to be identified during pregnancy. The maternal risk conditions can be classified as maternal characteristics of physical conditions ,medical illness or disorder, obstetrical conditions and warning signs.

Teaching is an important nursing intervention because a woman with illness must make modifications in her usual therapy to adjust to pregnancy. Pregnancy often stimulates woman to learn more about primary diseases. (**Pillitteri, 2003**)⁴

1.1 Need for the study

"Prevention is better than cure"

All pregnant women by virtue of their pregnant status face some level of maternal risk. Data suggest that amount all pregnant women around 40% of them have some complications. About 15% of the pregnant women need obstetric care to manage complications, which is potentially the threatening to mother and fetus. (Samiya M, Samina M, 2015)⁵

Approximately 10-20 % of all pregnancies in the United states are labeled as high risk. High risk pregnancies are account for more than half of all fetal and prenatal deaths. Nurses are in unique position to educate and empower women, through the phases of child birth, for them to achieve a healthy pregnancy with optimum outcome of a healthy baby. Despite various measures taken to reduce maternal mortality rate and morbidity rate, it remains very high in India (<u>http://www.faqs.org/</u> health/topics/ 3/High-risk pregnancy.html)⁶

Global statistics (WHO)

- Every day, approximately 830 women die from preventable causes related to pregnancy and childbirth.
- ✤ 99% of all maternal deaths occur in developing countries.

- Between 1990 and 2015, maternal mortality worldwide dropped by about 44%.
- Between 2016 and 2030, as part of the Sustainable Development Goals, the target is to reduce the global maternal mortality ratio to less than 70 per 100 000 live births.

Indian Scenario

556/one lakh live births-during 1990

398/one lakh live births - during 1997-1998

301/one lakh live births - during 2001-2003

178/ one lakh live births- during 2010-2012

At present 174/ one lakh live births (WHO, UNICEF, UNFPA, WORLD BANK AND UNITED NATIONS POPULATION DIVISION, 2015)

Tamilnadu Scenario

No State in India, except Tamil Nadu is able to calculate MMR reliably. In TamilNadu, the district level MMR has been calculated every year from 1994. Around 1100 maternal deaths occur every year which is approxi-mately 10% of maternal deaths occur in India and 0.2% of maternal deaths in the world. This gives an estimate MMR of 100 in TamilNadu while the SRS estimate is 134 (2001-03) During the year 2007-08, 1059 maternal deaths has reported which gives an estimate of 95 per one lakh live births. The district level MMR varies from 226 in Nilgiris district (hill area) to 32 in Chennai (100% urban areas) district. About 22% of deaths occur in ante natal Period, 6% in natal period and the remaining 72% on post natal period. The maternal deaths where as the

share of SC / ST population in total population is 20%. Among the 11% of maternal deaths occur in home deliveries, the share of SC / ST community is 50%. The major causes of maternal morbidity are pregnancy induced hypertension, anaemia, hemorrages and diabetic induced pregnancy. Cause of maternal deaths shown that 20% of mothers die due to PPH followed by PIH with severe anaemia. Maternal morbidity is another major cause for concern.

V.C. Subash Gandhi, coordinator of the Tamil Nadu Health System Project, said the staff nurses should play a key role in bringing down the maternal mortality rate. A close observation and surveillance would go a long way in achieving the desired results, he added.

Institutional Statistics

2012- 13	Causes	2013- 144	2014- 15	2015- 16	2016- 17
3	Haemorrhage	1	2	1	2
4	Eclampsia	3	4	5	3
1	DIC	1	1	2	1
-	Cardiac failure	1	-	1	4
1	Amniotic fluid embolism	1	1	2	1
-	Heart diseae	1	1	-	2
9	Total	8	9	11	13

Table 1.1: Maternal Death at Institute of Obstetrics and Gynaecology

The cost of prenatal care is modest compared to the expense of managing serious, preventable complications in the mother and fetus. It is clear that prenatal education on danger signals of pregnancy is a cost effective means for improving health of the mother and infant. $(Arulkumaran, 2012)^7$

A descriptive study conducted at Belgaum in selected hospitals, to evaluate the prenatal care among 100 risk mothers, revealed that they did not receive adequate prenatal care even though they had attended prenatal clinics. They were not informed about simple measure to prevent high risk conditions. This study recommends imparting education to high risk mothers on various aspects of prenatal care and signs of complications. This study also opens an avenue for research on the effect to maternal education for prevention of complications during pregnancy (**Reddy SR, 2014**)⁸

The researcher had seen various kinds or pregnancy associated complications in obstetric department. The women who are all attending

the antenatal outpatient department do not have an adequate knowledge regarding complications of high risk pregnancy. The researcher felt that in order to increase the knowledge and awareness about high risk pregnancy this topic has been selected.

1.2. Statement of the problem:

A Pre-experimental study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08.

1.3. Objectives of the study

- 1) The assess the knowledge regarding high risk pregnancy among antenatal women
- To assess the effectiveness of structured teaching programme on knowledge regarding high risk pregnancy among antenatal women.
- 3) To associate the findings with selected demographic variables and obstetrical variables.

1.4. Operational definitions

Effectiveness- refers to the extent to which the structured teaching programme on high risk pregnancy has achieved the desired effects in improving the knowledge of antenatal women as evident from gain in knowledge scores.

Structured teaching programme- refers to a systematically organized teaching programme for antenatal women to provide information regarding high risk pregnancy.

7

Knowledge- refers to understanding level of antenatal women regarding high risk pregnancy, as determined by their sources based on their responses.

Antenatal mother- refers to pregnant women between 8-28 weeks or gestational age.

High risk pregnancy- refers to pregnancy which is complicated by conditions such as Anemia, Pre-eclampsia, Gestational diabetes mellitus and Multiple pregnancy that adversely affect the maternal and fetal outcome.

1.5. Assumptions

- Antenatal mothers have lack of knowledge regarding high risk pregnancy.
- Structured teaching programme on high risk pregnancy will improve their knowledge and they can prevent further complications.

1.6. Research hypothesis

Antenatal women who receive structured teaching programme on high risk pregnancy will show a significant increase in the level of knowledge in post- test than that in pre-test.

1.7. Delimitations

- 1) Mother who come for antenatal checkup in antenatal outpatient department at Institute of Obstetrics and Gynaecology.
- 2) Study period is limited to 4 weeks.
- 3) Antenatal mothers with gestational age of 8-28 weeks.

CHAPTER-II REVIEW OF LITERATURE

The material gathered in the literature review should be treated as an integral part of the research data, since what is found in the literature cannot only have an important influence of the problem, but also find the design of the research, and provide useful comparative material when the data collected in the research is analysed.

Literature review is defined as a broad, comprehensive, in depth, systematic and critical review of scholarly publication, unpublished printed audio or visual materials and personal communications. (Suresh.K Sharma 2015)⁹

This chapter consists of two parts.

Part I : 2.1 Literature review related to study.

Part II: 2.2 Conceptual framework

2.1. Literature review related to study

This part is divided into two sections.

Section-A

2.1.1 Studies related to identification of high risk pregnancy.

Section -B

2.1.2 Studies related to Anemia.

2.1.3 Studies related to Pregnancy Induced Hypertension.

2.1.4 Studies related to Gestational Diabetes Mellitus.

2.1.5 Studies related to Multiple Pregnancy.

Section-A:

2.1.1 Studies related to Identification of high risk pregnancy:

(Samar k et al., 2014), conducted an exploratory descriptive study aimed to identify profile of high risk pregnancy among Saudi pregnant women in Taif. A convenient sample of two hundreds were included in the study subjects, an interview questionnaire also was used for data collection. The results revealed that 44% of Saudi women with high risk pregnancy and about two-thirds (66%) of them had previous complicated pregnancies and 68.7% of them complained of different medical associated conditions during their current pregnancies such as: anemia (25.3%), gestational diabetes (16.2%), pregnancy induced hypertension (15.2%). 40% of them reported exposure to smoking while 22% exposed to pesticide during their current pregnancies.¹⁰

(*Dutta S &Das 2012*), conducted a descriptive study regarding identification of high risk pregnancy at Srinagar. The study concludes that perinatal mortality rate was 46.1/ 1000 live birth. The perinatal mortality rate in women with no risk factor was 5.02 per 1000 live births, which rose with the level of risk from 27 per 1000 in low risk to 96 in moderate risk to 222 in the high risk group. The women with one or more risk factors were 17.1 times more likely to lose their baby during the perinatal period than those with no risk.¹¹

(*Shahzada J Malik 2010*),conducted a prospective study to identify the preventable factors in high risk pregnancy in India. Over a period of one year, of total 1,600 deliveries, 1,107 were considered to be at high risk, 33 fetal and 31 early neonatal deaths with an overall perinatal mortality rate of 40/1,000 births. The mortality was higher in mothers who had received inadequate antenatal care or with bad obstetric history.¹²

10

(*Dr.Mahmuda Mubasher 2010*), conducted a study to find out prevalence and outcome of high risk pregnancies in Pakistan. The sample was composed of 226 pregnancies. The study shows that high risk pregnancies were 69.5% as compared to 30.5% of low risk category. This indicates high prevalence of high-risk pregnancies in the area. It also shows that 73.24% of females with high-risk pregnancy were illiterate as compared to 62.31% with low risk pregnancy. Literate women were of 26.75% in high-risk pregnancies and of 37.68% in low risk.¹³

(*Taehan Kanho 2010*), conducted a descriptive study regarding the identification of high-risk pregnancy, using a simplified risk-scoring system at Chung Ang Medical Center, Korea. This study was based on the 1300 pregnant women. The results of the study show that 560 infants (42.7%) were born to mothers with risk-scores greater than 7, and 753 infants (57.3%) were born to mothers with risk scores less than 7. 2. Maternal age, parity, educational level,demographic factors was significant related statistically to identify the high risk pregnancies.¹⁴

(Yara Almerie et al.,2010), conducted a prospective study regarding near-miss and maternal mortality in maternity university hospital, Damascus, Syria revealed that there were 28025 deliveries, 15 maternal deaths and 901 near-miss cases. The study showed a MNMR of 32.9/1000 live births, a MMR of 54.8/1000 live births. Hypertensive disorders (52%) and haemorrhages (34%) were the top causes of nearmisses.¹⁵

Section B:

2.1.2 Studies related to anemia

(Jimenez K, Lang M.2016), quoted that anemia affects a fourth of the global population, with iron deficiency being the primary cause. It is associated with diminished work capacity, fatigue, impaired cognitive function, and can negatively impact the course of diseases like chronic heart failure or chronic kidney disease.¹⁶

(Shander A 2014), stated that anemia means haemoglobin below 13 g/dl in males, 12 g/dl in non-pregnant women, and 11 g/dl in pregnant women. Anemia is prevalent in various patient populations. The detrimental effect of anemia is present in all of this population of patients.¹⁷

(*Breymann C,2015*), emphasized a meta-analysis study that anemia is a common problem in obstetrics and perinatal care. Any hemoglobin below10.5 g/dL can be regarded as true anemia regardless of gestational age. Reasons for anemia in pregnancy are mainly nutritional deficiencies, parasitic and bacterial diseases, and inborn red blood cell disorders such as thalassemias. The main cause of anemia in obstetrics is iron deficiency, which has a world wide prevalence between estimated 20%-80% and consists of a primarily female population.¹⁸

(*Rahman MM et al.,2016*), conducted a systematic review and meta-analysis on maternal anemia and risk of adverse birth and health outcomes in low- and middle-income countries. Overall, 42.7% (95% CI: 37.0%, 48.4%) of women experienced anemia during pregnancy in low- and middle-income countries. There were significantly higher risks of low birth weight (RR: 1.31; 95% CI: 1.13, 1.51), preterm birth (RR: 1.63; 95% CI: 1.33, 2.01), perinatal mortality (RR: 1.51; 95% CI: 1.30, 1.76), and neonatal mortality (RR: 2.72; 95% CI: 1.19, 6.25) in pregnant women with anemia.¹⁹

(Burke RM, Leon JS, Suchdev PS 2014), stated that iron deficiency is a global problem across the life course. Maternal iron deficiency during pregnancy can predispose offspring to the development of iron deficiency during infancy, with potentially lifelong sequelae. This review explores iron status throughout these "first 1000 days" from pregnancy through two years of age, covering the role of iron and the epidemiology of iron deficiency.²⁰

(*De-Regil LM et al.,2015*), conducted all randomised or quasirandomised trials evaluating the effect of periconceptional folate supplementation alone, or in combination with other vitamins and minerals, in women independent of age and parity. Five trials involving 7391 women (2033 with a history of a pregnancy affected by a NTD and 5358 with no history of NTDs) were included. Four comparisons were made: 1) supplementation with any folate versus no intervention, placebo or other micronutrients without folate (five trials); 2) supplementation with folic acid alone versus no treatment or placebo (one trial); 3) supplementation with folate plus other micronutrients versus other micronutrients without folate (four trials); and 4) supplementation with folate plus other micronutrients versus the same other micronutrients without folate (two trials). Concluded that Folic acid, alone or in combination with vitamins and minerals, prevents NTDs, but does not have a clear effect on other birth defects.²¹

(*Taylor S, Rampton D,2015*), conducted a comparative study was on the indications for oral and Intravenous iron therapy. While most patients respond well to oral iron preparations, a substantial minority have side effects that make them adhere poorly to their treatment. For oral iron-intolerant patients, those responding poorly despite good adherence, and those with severe and/or symptomatic anemia, intravenous iron is an excellent alternative.²²

(Orlov IuP, Lukach VN, Govorova NV.2014), analysed a descriptive study with a modern view of iron exchange in general and during pregnancy. According to a large number of studies assigning to pregnant iron on the one hand, contributes to excessive activation of free radical oxidation, the accumulation of lipid peroxidation products and

demonstrations of eclampsia, and from the other potentiates the bacterial aggression and development of purulent-septic diseases that generally leads to the development of complications in pregnancy.²³

(Shi Q et al.,2015), made a meta-analysis of randomised controlled trials comparing patients treated with intravenous iron sucrose (intravenous group) with those treated with oral iron (oral group) for IDA during pregnancy was performed. Significant increases in haemoglobin [meandifference (MD), 0.85; 95% confidence interval (CI), 0.31-1.39; p = 0.002] and ferritin levels (MD, 63.32; 95% CI, 39.46-87.18; p < 0.00001) were observed in the intravenous group. Compared with the oral group, there were fewer adverse events in the intravenous group (risk ratio, 0.50; 95% CI, 0.34-0.73; p =0.0003). There was no significant difference in birth weight between the two groups.²⁴

(*Hlimi T,2015*), adopts a systematic approach to investigate the maternal outcome of anemia and eclampsia in relation to seasonality. A review of 23 published studies, shows a statistically significant link between these maternal disorders and seasonality in developing countries in Sub-Saharan Africa and Central and South Asia. Anemia and eclampsia tend to decrease during the dry season, only to increase with greater rainfall, low and cold temperatures. More research is required to identify the seasonal link between malaria and eclampsia particularly as climate change may exacerbate the rate of the disorders in tropical and sub-tropical areas.²⁵

(*Kidson-Gerber G et al., 2015*), conducted an observational study that demonstrated an association between refusal of blood products in major obstetric haemorrhage and increased morbidity and mortality. This review draws upon evidence in the literature, physiological principles and expert opinion for strategies and guidance to optimise the outcomes of pregnant women in whom blood transfusion is either refused or impossible. The importance of a multidisciplinary antenatal and perinatal management plan, including optimisation of haemoglobin and iron stores pre-delivery, blood loss minimisation, early haemorrhage control and postpartum anaemia treatment, is discussed.²⁶

(*Rukuni R 2015*), conducted a study to appraise whether a national screening programme could reduce the prevalence of iron deficiency anaemia and/or iron deficiency in pregnancy and improve maternal and fetal outcomes. Many studies evaluated haematological outcomes of anaemia, but few analysed clinical consequences. Haemoglobin and ferritin appeared the most suitable screening tests, although future options may follow recent advances in understanding iron homeostasis. The clinical consequences of iron deficiency without anaemia are unknown. Oral and intravenous iron are effective in improving haemoglobin and iron parameters. There have been no trials or economic evaluations of a national screening programme for iron deficiency anaemia in pregnancy.²⁷

2.1.3 Studies related to pre-eclampsia

(Dhariwal NK, Lynde GC, 2017), quoted that hypertensive disorders of pregnancy complicate approximately 10% of all deliveries in the United States and are a leading cause of maternal and fetal morbidity and mortality. Preeclampsia is defined as hypertension in association with proteinuria, thrombocytopenia, impaired liver function, renal insufficiency, pulmonary edema, or new-onset cerebral or visual disturbances.²⁸

(Spradley FT, Palei AC, Granger JP, 2015), made a systemic review summarized the current experimental evidence supporting the concept that obesity and metabolic factors like lipids, insulin, glucose, and leptin affect placental function and increase the risk for developing hypertension in pregnancy by reducing placental perfusion; enhancing placental release of soluble factors; and by increasing the sensitivity of the maternal vasculature to placental ischemia-induced soluable factors.²⁹

(Singh MD et al., 2015), researched a systematic search strategy to identify citations in electronic databases for the following terms: folic acid supplementation AND pre-eclampsia, folic acid supplementation AND genome stability, folate AND genome stability AND preeclampsia, folic acid supplementation AND DNA methylation, and folate AND DNA methylation AND pre-eclampsia. Forty-three articles were selected according to predefined selection criteria. The present review highlights associations between folate deficiency and certain biomarkers observed in various tissues of women at risk of preeclampsia.³⁰

(*Mudaliar and Menon 2015*), revealed that Pregnancy induced hypertension is defined as hypertension that develops for the first time in pregnancy after 20 weeks of gestation .Hypertension has been defined as blood pressure of 140/90 mm of Hg or greater, on at least two occasions, four to six hours apart.³¹

(Cormick G et al., 2016), assessed the risk of recurrent preeclampsia in the immediate subsequent pregnancy according to different birth intervals. Meta-analysis of adjusted odds ratios (aOR) with 95 % confidence intervals (CI) was used to measure the association between various interval lengths and recurrent pre-eclampsia or eclampsia. We identified 1769 articles and finally included four studies with a total of 77,561 women. The meta-analysis of two studies showed that compared to inter-pregnancy intervals of 2-4 years, the aOR for recurrent pre-eclampsia was 1.01 [95 % CI 0.95 to 1.07, I(2) 0 %] with intervals of less than 2 years and 1.10 [95 % CI 1.02 to 1.19, I(2) 0 %] with intervals longer than 4 years.³²

(*Vigil-De Gracia P, Ludmir J 2015*), made a randomized controlled trials comparing anticonvulsants with placebo or no anticonvulsant for prevention (a) of eclampsia in women with severe preeclampsia diagnosed during the postpartum period or diagnosed before delivery but without previous treatment and (b) prevention of seizures recurrence in women with eclampsia postpartum. There was no clear difference between the groups reporting eclampsia (relative risk: 0.54, 95% confidence interval: 0.16-1.80). For seizure recurrence, magnesium sulfate was superior to diazepam, but there was no significant difference compared with phenytoin. No conclusion can be drawn on the role of magnesium sulfate post partum as established in antepartum pre-eclampsia/eclampsia management because of lack of powered randomised controlled trials.³³

(Brown CM, Garovic VD 2014), conducted study that compared and contrasted the recommendations from different treatment guidelines and outlined some newer perspectives on management, aims to provide a clinically oriented guide to the drug treatment of hypertension in pregnancy. A recent report highlighted hypertensive disorders as one of the major causes of pregnancy-related maternal deaths in the USA, accounting for 579 (12.3 %) of the 4,693 maternal deaths that occurred between 1998 and 2005. In low-income and middle-income countries, preeclampsia and its convulsive form, eclampsia, are associated with 10-15 % of direct maternal deaths.³⁴

(*Meher S et al., 2017*), researched a meta-analysis of individual participant data including 32,217 women and 32,819 babies recruited to 31 randomized trials comparing low-dose aspirin or other antiplatelet agents with placebo or no treatment for the prevention of preeclampsia.

There was no significant difference in the effects of antiplatelet therapy for women randomized before 16 weeks' gestation compared with those randomized at or after 16 weeks for any of the prespecified outcomes.³⁵

2.1.4 Studies related to gestational diabetes mellitus

(*Taschereau-Charron A et al., 2017*),conducted a comparative study to compare FA profiles in different blood lipid fractions and the influence of dietary fat intake in women with GDM or normoglycemic pregnancies. Results show that women with GDM have more saturated and less polyunsaturated FA (PUFA) in their red blood cell (RBC) membranes than normoglycemic pregnant women. Further research is required to determine whether FA profiles are altered prior to the diagnosis of GDM and can be prevented by diet.³⁶

(*Rozenberg P, 2016*), conducted a randomized control trial was for induction of labour and expectant management in macrosomic babies. The mean birth weight (\pm SD) was 3831 (\pm 324) g in the induction group 4118(\pm 392) g in the expectant group. Induction of labor significantly reduced the risk of shoulder dystocia or associated morbidity (8/407; 2 %) compared with expectant management (25/411; 6 %); P=0.004.³⁷

(Harrison AL et al., 2016), conducted a randomised trials in Pregnant women diagnosed with gestational diabetes mellitus. This systematic review identified eight randomised, controlled trials involving 588 participants; seven trials (544 participants) had data that were suitable for meta-analysis. Five trials scored ≥ 6 on the PED scale, indicating a relatively low risk of bias. Meta-analysis showed that exercise, as an adjunct to standard care, significantly improved postprandial glycaemic control (MD-0.33mmol/L, 95% CI -0.49 to -0.17) and lowered fasting blood glucose (MD -0.31mmol/L, 95% CI - 0.56 to -0.05) when compared with standard care alone, with no increase in adverse events.³⁸

(*McCance DR, 2013*), made a randomised trials have offered new insight of hyperglycaemia to adverse pregnancy outcome was clearly demonstrated by the Hyperglycaemia and Adverse Pregnancy Outcome (HAPO) study, but translation of these data into clinical practice has proved challenging because of the continuum of risk. Long-term metabolic and cardiovascular implications of hyperglycaemia during pregnancy for mother and child are now generally recognised with major implications for public health.³⁹

(Valentina Palkova, 2011), conducted a quasi experimental study to assess the effectiveness of an educational program on gestational diabetes mellitus was done among 30 patients with mild GDM and between 19 and 41 years of age. The women were divided into two groups-for group 1 (n=15) education was given group II (n=15) was considered as control group. One month educational programmed was conducted. The clinical data and metabolic control for the two groups were evaluated. Group I showed better improvement in comparison with Group II, and also mild improvement in the blood glucose level. The results proved that the educational approach has the patients to improve patient's quality of life.⁴⁰

Tieu J et al., (2011), analysed a randomised controlled trials (RCTs) and quasi-RCTs to assess the effects of dietary advice interventions compared with no intervention (standard care), or to different dietary advice interventions. Very low-quality evidence from five trials suggests a possible reduction in GDM risk for women receiving dietary advice versus standard care, and low-quality evidence from four trials suggests no clear difference for women receiving low-versus moderate- to high-GI dietary advice. A possible reduction in

pregnancy-induced hypertension for women receiving dietary advice was observed and no clear differences were seen for other reported primary outcomes.⁴¹

Farrar D et al (2016), conducted a systematic review and metaanalysis on hyperglycaemia and risk of adverse perinatal outcomes. the odd ratios is large for gestational age per 1 mmol/L increase of fasting and two hour post-load glucose concentrations (after a 75 g OGTT) were 2.15 (95% confidence interval 1.60 to 2.91) and 1.20 (1.13 to 1.28), respectively. This review and meta-analysis identified a large number of studies in various countries. There was a graded linear association between fasting and post-load glucose concentration across the whole glucose distribution and most adverse perinatal outcomes in women without pre-existing or gestational diabetes.⁴²

2.1.5 Studies related to multiple pregnancy

(Moldenhauer JS, Johnson MP, 2015), quoted that although monochorionic (MC) twins comprise only 20% of all twin pregnancies, the risk for fetal loss and long-term morbidity is significantly higher than dichorionic twins due to the presence of placental vascular anastomoses.⁴³

(*Tran et al., 2015*), conducted a study on delayed- interval delivery in multifoetal pregnancy. Among 18 relevant cohort studies 391 twin and 39 triplet pregnancies could be analyzed. In case of delayed delivery, the survival rate of the second twin or higher order multiple was respectively 44.8% and 82.7% when the first twin was born before or after 24 weeks of gestation. The later was the delivery of the first twin, the higher was the second twin's survival rate, but the shorter was the interval between births(14 Vs 26 days).⁴⁴

(Joshua, p. Vogel, 2013), conducted multilevel logistic regression to determine the association between twins and adverse maternal and perinatal outcomes.279.425 mothers gave birth to 276,187(98.8%) singletons and 6,476(1.2%) twins. Odds of severe adverse maternal outcomes (Death, blood transfusion, ICU admission or hysterectomy) (AOR 1.85, 95% CI 1.60-2.14) and perinatal mortality (AOR 2.46, 65% CI 1.40-4.35) in twin pregnancies were higher, however early neonatal deaths didn't reach significance. ⁴⁵

(*Lee, K.E,2012*), conducted a study to develop a greater understanding of the grief parents of a multi-fetal pregnancy experience when one or more of their infants are lost either prior to delivery or in the neonatal period. Recommendations are put forth as to how the NICU nurse can provide care for bereaved parents of multiples as aligned with the human science paradigm of nursing. ⁴⁶

(*Richards, J et al.,2015*), carried out a qualitative study with mothers who had experienced a loss in pregnancy or the neonatal period and had a surviving twin on the neonatal unit. The analysis identified three key themes in the accounts mothers gave of their experiences: the status of 'special'; the importance of trust; and control and empowerment. Where the surviving co-twin remained in hospital for many weeks, mothers described the emotional support of health professionals as crucial to their wellbeing.⁴⁷

(Whitford HM et al., 2017), made a quasi-randomised trial to assess effectiveness of breastfeeding education and support the women with twins or higher order multiples. We searched the Cochrane Pregnancy and Childbirth Group's Trials Register (30 June 2016), ClinicalTrials.gov (30 June 2016), We found no evidence from randomised controlled trials about the effectiveness of breastfeeding education and support for women with twins or higher order multiples, or the most effective way to provide education and support.⁴⁸ (*Kushnir VA et al., 2017*), conducted a Systematic review of worldwide trends in assisted reproductive technology. SETs and utilization of frozen-thawed embryos increased worldwide over the study period. In 2012 SET utilization in all ART cycles was highest in Japan and Australia/New Zealand (82.6% and 76.3% respectively) and lowest in Latin America (16.0%).⁴⁹

(*Lumbiganon P et al., 2016*), analysed all identified published, unpublished and ongoing randomised controlled trials (RCTs) assessing the effect of formal antenatal breast feeding education or comparing two different methods of formal antenatal breast feeding education, on the duration of BF. Assessed all potential studies identified as there was no conclusive evidence supporting any antenatal BF education for improving initiation of BF, proportion of women giving any BF or exclusively BF at three or six months or the duration of BF. There is an urgent need to conduct a high-quality, randomised controlled study to evaluate the effectiveness and adverse effects of antenatal BF education, especially in low- and middle-income countries. ⁵⁰

(Moore ER et al., 2016), emphasized a randomized controlled trial that compared immediate or early Skin to Skin Contact with usual hospital care. Quality of the evidence was assessed using the GRADE approach. Evidence supports the use of SSC to promote breastfeeding. Studies with larger sample sizes are necessary to confirm physiological benefit for infants during transition to extra-uterine life and to establish effects possible dose-response and optimal initiation time. Methodological quality of trials remains problematic, and small trials reporting different outcomes with different scales and limited data limit our confidence in the benefits of SSC for infants. ⁵¹

22

2.2. Conceptual frame work

A conceptual framework deals with abstraction, which is assembled by nature of their relevance to a common theme. It describes the mental image of phenomenon and integrates them into a meaningful configuration. It is a visual diagram by which the researcher explains the specific area of interest.

The study was aimed at determining the effectiveness of structured teaching programme on knowledge regarding early identification of high risk pregnancy among antenatal women.

The investigator adopted "Modified Pender's Health Promotion Model (1984)

The model focuses on aspects of individuals configuration to perceptional factors, modifying factors and participation on health promoting behavior. The model also identifies factors that influence health promotion activities.

The Model focuses on the following areas :

- 1) Cognitive perceptual factors
- 2) Perceived health status
- 3) Health promoting services
- 4) Perceived benefits of health promotions
- 5) Barriers to health promoting behavior

Cognitive perceptual factors

In this modified model, the investigator finds out the cognitive perceptual factors of antenatal women on high risk pregnancy. The investigator assessed their knowledge regarding the causes, signs and symptoms, prevention, management and complications and high risk pregnancy by using structured questionnaire.

Perceived health status

The antenatal women have inadequate knowledge regarding early identification of high risk pregnancy and such women should undergo the next step that involves health promoting services.

Health promoting services

Structured teaching programme on high risk pregnancy is given as health promoting service to the antenatal women irrespective of their knowledge.

Perceived benefits of health promotion

Health promoting behavior is the desired behavioral outcome. In this study health promoting behavior developed by the teaching programme will result in improved knowledge on high risk pregnancy among antenatal women.

Barrier to health promoting behavior

In this study, if the antenatal women have inadequate knowledge on high risk pregnancy, reassessment of the antenatal women knowledge has to be done. But it is not included in this study.

CHAPTER-III RESEARCH METHODOLOGY

Introduction

Methodology is the most important part of any research study, which enables the researcher to form a blue print for the study undertaken. This chapter deals with the methodology related to evaluate the effectiveness of Structured Teaching Programme on knowledge regarding early identification of high risk pregnancy among antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08.

3.1. Research approach

Research approach involves the description of the plan to investigate the phenomenon under study in a structured, unstructured or a combination of the two methods. For this research study the investigator selects the quantitative research approach.

3.2. Study design

Research design incorporates the most important methodology called decisions that the researcher make in conducting a research study. It helps the researcher in the selection of subjects and the manipulation of independent variables to be studied.

The investigator proposed Pre- experimental research design for this Study.

In pre-experimental research design one group pre-test post-test design was used.

Pre Test	Structured Teaching Programme	Post Test
01	Х	O2

In the present study pretest is administered by means of structured questionnaire and then planned teaching programme will be provided to the group .Post test will be conducted by using the same structured questionnaire.

 $O_{1=}$ Assess the pretest level of knowledge regarding Early identification of High risk pregnancy among antenatal mothers.

X= Structured Teaching Programme on Early identification of High risk pregnancy.

 O_2 = Assess the posttest level of knowledge among the same group, one week after STP.

3.3. Setting of the study

The study was conducted in Institute of Obstetrics and Gynaecology and Government Hospital for Women and Children, Chennai. This hospital was started in the year 1844 for public service. It is a 1075 bedded maternity hospital, tertiary care centre and referral centre. The hospital is renowned for its excellence in medical experts, nursing care and quality diagnostic services. All facilities are provided for conducting normal, high risk and instrumental deliveries. IOG has departments like neonatal intensive care unit, family planning services, oncology department, endocrinology, human milk bank and genetic department which are rendering comprehensive care for entire Tamilnadu and for neighboring state like Andhra Pradesh also.

3.4. Duration of the study

The study was conducted for a period of 4 weeks from 20.11.16 to 18.12.2016.

3.5. Study population

Population is the aggregation of all the unit in which the researcher is interested.

All antenatal mothers attending antenatal outpatient Department are the population for my study.

3.5.1 Target population

A target population consists of the total number of people or objects which are meeting the designated set of criteria.

Antenatal mothers who are in 8-28 weeks of pregnancy are target population of this study.

3.5.2 Accessible population

It is the aggregate of cases that conform to designated criteria and also accessible as study subjects for study.

Antenatal mothers between 8-28 weeks and who have attended Antenatal Outpatient Department during the data collection period are the accessible population for this study.

3.6. Sample

The sample comprised of all antenatal women with gestational age of 8-28 weeks, and who met the inclusion criteria .

3.7.Sampling criterion

The researcher specified certain inclusion and exclusion characteristics for the population to be considered as a sample. Accordingly the population was studied and those come under inclusion were selected as the sample and the other elements were excluded from the study.

27

3.7.1. Inclusion criteria

- Antenatal women with gestational age of 8-28 weeks.
- Antenatal women who attended antenatal outpatient department..
- Antenatal women who are willing to participate.
- Antenatal women who can able to read and speak Tamil and or English.

3.7.2. Exclusion criteria

- Antenatal women who have visual and hearing impairment.
- ✤ Antenatal women with mental illness.
- Antenatal women having related obstetrical complications like severe Anemia, Pre- eclampsia, Gestational Diabetes Mellitus, and Multiple pregnancy.

3.8. Sample size

Sample size of the study is 60.

3.9. Sampling technique

Purposive sampling technique was adopted for this study.

3.10. Research variables

Variables are concept at different levels of abstraction that are concisely defined to promote their measurement and manipulation within the study.

3.10.1 . Independent variable

Structured Teaching Programme on knowledge regarding high risk pregnancy is the independent variable of the study.

3.10.2. Dependent variable

Knowledge of antenatal mother is the dependent variable of the study.

3.10.3 . Influencing variable

Age, education, occupation, type of family, age at marriage, and obstetrical score of antenatal mother is the influencing variable.

3.11. DEVELOPMENT AND DESCRIPTION OF TOOL

3.11.1 development of tool

A structured questionnaire was developed on the basis of objectives of the study. Tool was developed after extensive review of literature from various text books, journals, internet search and discussion and guidance from the experts in the field of Nursing, and medical experts from the Institute of Obstetrics and Gynaecology. The tool was developed in English and translated into Tamil. Congruency was maintained in translation.

3.11.2 . Description of the tool

The tool consists of three section.

Section-A

It consists of demographic variables of the sample such as age, religion, type of family, education, occupation and family monthly income.

Section-B

It consists of obstetrical variables of the sample such as age at marriage, last menstrual period, expected date of delivery, obstetrical score, age during first child birth and pre- pregnant health status.

Section-C

Structured questionnaire is to assess the knowledge of antenatal women regarding high risk pregnancy. It consists of 30 multiple choice questions regarding high pregnancy. It consists of Part A,B ,C and D.Part A and B each consists of 8 questions and part C and D each consists of 7 questions.

- **Part B:** Questions related to Pre-eclampsia
- Part C: Questions related to gestational diabetes mellitus
- **Part D:** Questions related to multiple pregnancy

Scoring key

Each correct answer carries - One Mark

Each wrong answer carries – Zero Mark

<50% - Inadequate knowledge

51-75% - Moderately adequate knowledge

76-100% - Adequate knowledge

3.11.3 Teaching Module

The teaching module was developed by literature review and by obtaining expert's opinion. The aid used for this study was a booklet on high risk pregnancy. The structured teaching programme consists of introduction, definition, causes, signs and symptoms, preventive measures, management and complications of high risk pregnancy. The method of teaching programme will be given by lecture cum discussion.

3.12. Content validity

Validity of the tool was assessed using content validity. Content validity was determined by experts from Nursing and Medical. They suggested certain modifications in tool. After the modifications, they agreed this tool to evaluate the effectiveness of Structured Teaching Programme on knowledge regarding early identification of High risk pregnancy among antenatal mothers attending at Institute of Obstetrics and gynaecology, Chennai.

3.13. Protection of human subjects

Following submission of study proposal, the permission was obtained from Institutional Ethics committee. Permission for conducting the study was obtained from the Director of Institute of Obstetrics & gynaecology and Hospital for Women and Children. Thus the investigator followed the guidelines which were issued by the Institutional Ethics committee. Confidentiality of the results and anonymity will be assured to the subjects. Throughout the study period the respect of the mother and the family members will be maintained. The participants, were assured that at anytime they can withdraw from the study.

3.14. Reliability of the tool

Reliability of the tool was assessed by using Test-retest method and its correlation coefficient r value was 0.84. This correlation coefficient is very high and it is a good tool to evaluate the effectiveness of Structured Teaching Programme on knowledge regarding early identification of High risk pregnancy among antenatal mothers attending at Institute of Obstetrics and gynaecology, Chennai.

3.15. Pilot study

Pilot study is a trial run for the major study to test the reliability, practicability, appropriateness and flexibility of the tool for the study. The investigator conducted a pilot study with 10 samples in Institute of Obstetrics and Gynaecology, Egmore. The data was analysed by using descriptive statistics that is percentage, mean and standard deviation of the variables and were calculated and compared. In the pilot study the researcher had no difficulties and the study was found to be feasible and practicable. The investigator proceeded for the main study.

3.16. Data collection procedure

The study was conducted in Antenatal outpatient department, after obtaining permission from the Director of IOG. Before the data collection, the researcher introduced herself, explained the purpose of the study to the OPD staff nurse and antenatal mothers between the 8-28 weeks of gestational age.The confidentiality was assured and consent was obtained from the participants.Three to five participants were selected every day and assured that at anytime they can withdraw from the study.

Data collection was done using questionnaire method. In pretest the researcher administered structured questionnaire to each participant to assess the knowledge of antenatal women on high risk pregnancy. The subjects took 20-30 minutes to answer the questionnaire by interview method. After that a structured teaching programme was conducted on the same day approximately for 30 minutes per subject and booklets were provided. After that the researcher clarified many doubts asked by the participants regarding my study and also in general, according to the need of each participant. Then each mother was thanked and instructed to come on seventh day after the pretest. The researcher had given a reminder through phone on previous day. Post test was conducted after one week by using the same questionnaire to find out the effectiveness of teaching programme. The data collection process was terminated after thanking the participants for their co-operation. After collecting the posttest questionnaire, every now and then, coding sheet were entered.

The data collection period was 4 weeks.(20.11.16 to 18.12.2016). Data was collected for 6 days in a week.

33

3.17. Intervention protocol

Place	: Antenatal OPD, IOG, Chennai.	
Intervention tool	: Lecture cum discussion	
Duration	: 30 minutes	
Frequency	: One-time teaching	
Time	: 7am-1pm	

3.18. Data entry analysis

The obtained data were analyzed on the basis of objectives and hypothesis by using the descriptive and inferential statistics.

- 1) Organize the data.
- Descriptive statistics was used to analyse the frequency, percentage, mean and standard deviation of the following variables.
 - a. Demographic variables were analysed in terms of frequency and percentage.
 - b. Obstetrical variables were analysed in terms of frequency and percentage.
 - c. Knowledge on high risk pregnancy among antenatal women before and after administration of structured teaching programme was analysed in terms of frequency, percentage and standard deviation.
- 3) Inferential statistics was used to determine the comparison and association.

- a. Difference between pre-test and post-test was calculated using McNemars test.
- b. Paired 't' test was used to find out the effectiveness of structured teaching programme by comparing the pre-test and post- test knowledge scores.
- c. Chi-square was used to associate the knowledge with selected demographic and obstetrical variables.
- 4) P<0.05 was considered statistically significant.

CHAPTER-IV DATA ANALYSIS AND INTERPRETATION

Analysis is categorizing, ordering, manipulation and summarizing of data to obtain the answers for hypothesis.

This chapter deals with the analysis and interpretation of data collected from 60 antenatal mothers between the gestational age of 8-28 weeks at Institute of Obstetrics and Gynaecology, Egmore, to evaluate the effectiveness of structured teaching programme on high risk pregnancy.

The purpose of analysis was to reduce the collected data to an intelligible and interpretable form so that the relation of the research problem can be studied and tested.

The collected data was analyzed by using descriptive and inferential statistics.

Organization of data

Section-A: Demographic and Obstetric information of antenatal mothers those who participated in the study.

Section-B: Assess the pre-test knowledge regarding high risk pregnancy in the study group.

Section-C: Assess the post-test knowledge regarding high risk pregnancy in the study group.

Section-D: Evaluating the effectiveness of structured teaching programme on knowledge regarding high risk pregnancy in the study group.

Section-E: Association of findings with selected demographic and Obstetric variables.

Statistical analysis

- Demographic variables in categorical/ dichotomous were given in frequencies with their percentage.
- 2) Knowledge score was given in mean and standard deviation.
- 3) Difference between pre-test and post-test was analysed using student paired t-test.
- 4) Categorical variables difference between pre-test and post-test was calculated using McNemars test.
- 5) Differences between pre-test and post-test score was analysed using percentage with 95% CI and mean difference with 95% CI.
- 6) Association between post-test knowledge score with demographic variables are analysed using Chi-square test.
- 7) Association between knowledge gain score with demographic variables are analysed using oneway analysis of variance and student independent t-test.
- 8) P<0.05 was considered statistically significant.

Section-A : Demographic and obstetric information of antenatal mothers those who participated in the study

Dem	ographic variables	Antenatal women	%
Age	< 20 years	1	1.7%
	20 - 28 years	50	83.3%
	> 28 years	9	15.0%
Religion	Hindu	54	90.0%
	Muslim	3	5.0%
	Christian	3	5.0%
Family type	Nuclear family	24	40.0%
	Joint family	36	60.0%
Education	Elementary education	13	21.7%
	Middle education	11	18.3%
	Higher secondary education	22	36.7%
	Graduate	14	23.3%
Job	Home maker	54	90.0%
	Government job	1	1.7%
	Private job	2	3.3%
	Self job	3	5.0%
Income	< Rs.10000	37	61.7%
	Rs.10001 -20000	23	38.3%
Age at	< 20 years	19	31.7%
marriage	21 -25 years	33	55.0%
	> 25 years	8	13.3%
Gravida	Primi	32	53.3%
	Multi	28	46.7%

 Table-4.1:Demographic
 and
 Obstetrical
 profile
 in
 the
 study
 group:

Table 4.1 shows the demographic and obstetrical information of antenatal mothers who are participated in the study.

In considering the **age** wise distribution of antenatal mothers,83.3% were 20-28 years of age, 15% were in more than 28 years of age group, and remaining 1.7% belongs to less than 20 years of age.

In the **religion** wise distribution , 90% are Hindus , 5% belongs to Christian religion and remaining 5% are of in Muslim religion.

In considering the **family type**, 60% are in joint family and remaining 40% belongs to nuclear family.

In **education** wise distribution, 36.7% (22) had Higher secondary education, 23.3% (14) were graduates, 21.7%(13) had elementary education and 18.3%(11) were of middle education status.

In considering the **job**, 90%(54) of them were home makers, 5%(3), of them had self job, and 3.3%(2) of them were working in private sector and only 1.7%(1) in Government jobs.

In the **monthly income** status, 61.1%(37) family monthly income was below Rs.10,000 and remaining 38.8%(23) monthly income was Rs. 10001-20000.

In considering the Obstetrical variables, in **age at marriage**, 55% (55) were married at the age of 21-25 years, and 31.7%((19) were married at the age of below 20 years and remaining 13.3%(8) were married at more than 25 years of age.

In **gravida**, 53.3%(32) were primi and remaining 46.7%(28) were multi.

39

Section-B : Assess the pre-test knowledge regarding high risk pregnancy in the study group.

		Mean	SD	% of knowledge
Anemia	8	3.35	1.89	41.9%
Pre-eclampsia	8	3.10	1.66	38.8%
Gestational Diabetes Mellitus	7	2.57	1.81	36.7%
Multiple pregnancy	7	2.45	1.73	35.0%
Total	30	11.47	4.64	38.2%

Table-4.2: Percentage of pre-test knowledge score in the study group:

Table 4.2 represents the percentage of pretest knowledge score each domain wise. They are having more score in anemia(41.9%), and moderate score of 38.8% in pre-eclampsia and 36.7% in gestational diabetes mellitus, and minimum score in multiple pregnancy(35.0%) Overall they are having 38.2% of score.

Table 4.3: Pre-test level of knowledge score in the study group:

Level of knowledge	No. of antenatal women	%
Inadequate	46	76.7%
Moderate	14	23.3%
Adequate	0	0.0%
Total	60	100%

Table 4.3 shows the level of pretest knowledge regarding early identification of high risk pregnancy among antenatal women before administration of structured teaching programme. 76.7%(46) of the antenatal women had inadequate knowledge score, 23.3%(14) of them had moderate knowledge score, and none of them are having adequate level of knowledge score.

Section-C: Assess the post-test knowledge regarding high risk pregnancy in the study group.

		Mean	SD	% of knowledge
Anemia	8	6.78	1.01	84.8%
Pre-eclampsia	8	6.52	.97	81.5%
Gestational Diabetes Mellitus	7	5.93	.86	84.7%
Multiple pregnancy	7	5.80	.88	82.9%
Total	30	25.03	2.22	83.4%

Table-4.4: Percentage of post-test knowledge score in the study group:

Table 4.4 shows each domain wise post-test percentage of knowledge score regarding early identification of high risk pregnancy among antenatal women. They are having more score in anemia(84.8%), and minimum score in preeclampsia(81.5%), score in gestational diabetes mellitus(84.7%), and score in multiple pregnancy (82.9%). Overall they are having 83.4% of score.

Table 4.5: Post-test level of knowledge score in the study group:

Level of knowledge	No. of antenatal women	%
Inadequate	0	0.0%
Moderate	13	21.6%
Adequate	47	78.4%
Total	60	100%

Table 4.5 represents the level of posttest knowledge regarding early identification of high risk pregnancy among antenatal women after administration of structured teaching programme. 78.4%(47) of them are having adequate level of knowledge score, 21.6%(13) of them are having moderate knowledge score, None of the antenatal women are having inadequate knowledge score.

Section-D: Evaluating the effectiveness of structured teaching programme on knowledge regarding high risk pregnancy in the study group.

	Knowledge score						
	Pre-test Post-		Post-	test	Mean Difference	Student's paired t-test	
	Mean	SD	Mean	SD		T	
Anemia	3.35	1.89	6.78	1.01	3.43	t=12.41, P=0.001*** significant	
Pre-eclampsia	3.10	1.66	6.52	.97	3.42	t=16.05, P=0.001*** significant	
Gestational Diabetes Mellitus	2.57	1.81	5.93	.86	3.37	t=15.01 P=0.001*** significant	
Multiple pregnancy	2.45	1.73	5.80	.88	3.35	t=14.25, P=0.001*** significant	

Table-4.6: Comparison of pre and post-test knowledge score on high risk pregnancy:

* significant at P≤0.05

** highly significant at P≤0.01

*** very highly significant at $P \le 0.001$.

Table 4.6 shows comparison of Pretest and Posttest mean knowledge score.

Considering anemia aspects, in pretest antenatal mothers are having 3.45 score where as in posttest they are having 6.78 score, so the difference is 3.43. This difference between pretest and posttest is large and is statistically significant. Considering preeclampsia aspects, in pre-test antenatal mothers are having 3.10 score where as in posttest they are having 6.52 score, so the difference is 3.42. This difference between pretest and posttest is large and it is statistically significant.

Considering Gestational Diabetes Mellitus aspects, in pretest, antenatal mothers are having 2.57 score where as in posttest they are having 5.93, so the difference is 3.37. This difference between pretest and posttest is large and it is statistically significant.

Considering Multiple Pregnancy aspects, in pretest antenatal mothers are having 2.45 score where as in posttest they are having 5.80 score, so the difference is 3.35. This difference between pretest and posttest is large and it is statistically significant.

Statistical significance was calculated by using student's paired 't' test.

Table-4.7: Comparison of overall pre and post-test knowledge score on high risk pregnancy:

	No. of antenatal women	Knowledge score Mean ± SD	Mean Difference	Student's paired t-test
Pretest	60	11.47±4.64	10.55	t=21.88 P=0.001***
Posttest	60	25.03±2.22	13.57	significant

** highly significant at P≤0.01

*** very highly significant at $P \le 0.001$

Table 4.7 represents the comparison of overall pretest and posttest knowledge score. Considering overall knowledge score, in pretest antenatal women are having 11.47 score where as in posttest they are having 25.03, so the difference is 13.57.The difference between pretest and post-test score is large and it is statistically significant. Differences between pretest and posttest score was analysed using students paired 't' test

	L	evel of k				
	Pre-test		test Post-test		Extended McNemar's test	
	Ν	%	n	%	Wieneman's test	
Inadequate	46	76.7%	0	0.0%		
Moderate	14	13.3%	13	21.6%	$\chi 2=53.83$ p=0.001*** DF= 2	
Adequate	0	0.0%	47	78.4%	p=0.001*** DF= 2 significant	
Total	60	100%	60	100%	6	

Table4. 8: Pre-test and post-test level of knowledge score in the study
group:

** highly significant at P≤0.01

*** very highly significant at $P \le 0.001$

Table 4.8 shows the pre-test and post-test level of knowledge score regarding early identification of high risk pregnancy among antenatal women, before and after administration of Structured Teaching Programme. In pretest, 76.7% of antenatal women are having inadequate knowledge score, 13.3% of them are having moderate knowledge score, none of them are having adequate level of knowledge score.

In posttest, none of the antenatal women are having inadequate knowledge score, 21.6% of them are having moderate knowledge score, 78.4% of them are having adequate level of knowledge score.

Statistically there is a significant difference between pretest and posttest knowledge score. It was confirmed using extended McNemar's test.

Table-4.9: Percentage of knowledge gain score on high risk pregnancy:

	Max score	knowledge score Mean ± SD	Mean Difference in score with 95% Confidence interval	Percentage of gain score with 95% Confidence interval
Pretest	30	11.47±4.64	13.57(12.33 – 14.81)	45.2%(41.1% – 49.4%)
Posttest	30	25.03±2.22		

Table 4.9 shows the comparison of overall knowledge score between pretest and posttest. On an average, in posttest, antenatal women gained 45.2% of knowledge score after having Structured Teaching Programme regarding early identification of high risk pregnancy. Differences between pretest and posttest score was analysed using proportion with 95% confidence interval and mean difference with 95% confidence interval. This 42.5% of knowledge gains shows the effectiveness of Structured Teaching programme on early identification of high risk pregnancy among antenatal women.

Domains	Pre-test Knowledge	Post-test Knowledge	% of knowledge Gain	
Anemia	41.9%	84.8%	42.9%	
Pre-eclampsia	38.8%	81.5%	42.7%	
Gestational Diabetes Mellitus	36.7%	84.7%	48.0%	
Multiple pregnancy	35.0%	82.9%	47.9%	
Overall	38.2%	83.4%	45.2%	

Table-4.10: Effectiveness of structured teaching programme on high risk pregnancy:

Table 4.10 represents each domain wise percentage of knowledge gain. 42.9% of knowledge gain anemia aspects, 42.7% knowledge gain in preeclampsia aspects, and 48% knowledge gain in Gestational diabetes mellitus and 47.9% of knowledge gain in Multiple pregnancy aspect. Overall they gained 45.2% of knowledge score when comparing pretest and posttest after having STP. This shows the effectiveness of STP on knowledge regarding high risk pregnancy among antenatal women.

Section-E: Association of findings with selected demographic and obstetric variables.

Demographic variables			Pretest knowl		Total	Chi square test		
		Ina	dequate	Moderate				
		n	%	n	%			
Age	< 20 years	1	100.0%	0	0.0%	1	χ2=2.85	
	20 - 28 years	40	80.0%	8	20.0%	50	P=0.24 DF=2 NS	
	> 28 years	5	55.6%	4	44.4%	9		
Religion	Hindu	41	75.9%	13	24.1%	54	χ2=1.09	
	Muslim	3	100.0%	0	0.0%	3	P=0.57 DF=2 NS	
	Christian	2	66.7%	1	33.3%	3		
Family type	Nuclear family	19	79.2%	5	20.8%	24	χ2=0.14 P=0.71	
	Joint family	27	75.0%	9	25.0%	36	DF=1 NS	
Education	Elementary education	10	76.9%	3	23.3%	13	χ2=0.92 P=0.81	
	Middle education	8	72.7%	3	27.3%	11	DF=3 NS	
	Higher secondary education	16	72.7%	6	23.3%	22		
	Graduate	12	85.7%	2	14.3%	14		
Job	Home maker	41	75.9%	13	24.1%	54	χ2=1.09	
	Government job	1	100.0%	0	0.0%	1	P=0.78 DF=3NS	
	Private job	2	100.0%	0	0.0%	2		
	Self job	2	66.7%	1	33.3%	3		

Table4. 11: Association between pretest level of knowledge score with demographic and obstetrical variables:

Demographic variables			Pretest knowl			Chi		
		Inadequate		Moderate		Total	square test	
		n	%	n	%			
Income	< Rs.10000	28	75.7%	9	24.3%	37	χ2=0.06	
	Rs.10001 - 20000	18	78.3%	5	21.7%	23	P=0.82 DF=1 NS	
Age at marriage	< 20 years	13	68.4%	6	31.6%	19	χ2=1.33	
	21 - 25 years	26	78.8%	7	21.2%	33	P=0.51 DF=2 NS	
	> 25 years	7	87.5%	1	12.5%	8		
Gravida	Primi	27	84.4%	5	15.6%	32	χ2=2.27	
	Multi	19	67.9%	9	32.1%	28	P=0.13 DF=1 NS	

** highly significant at P≤0.01

*** very highly significant at $P \le 0.001$

Table 4.11 shows the association between pretest level of knowledge score with clients demographic variables. None of the demographic variables are significantly associated with their pretest level of knowledge. It was confirmed using Chi square test.

Demographic variables			Posttest know			Chi		
		Moderate		Adequate		Total	square test	
		n	%	n	%		iesi	
Age	< 20 years	1	100.0%	0	0.0%	1	χ2=6.61	
	20 - 28 years	12	24.0%	38	76.0%	50	P=0.03*	
	> 28 years	0	0.0%	9	100.0%	9	DF=2 S	
Religion	Hindu	12	22.2%	42	77.8%	54	χ2=1.08	
	Muslim	1	33.3%	2	66.7%	3	P=0.58	
	Christian	0	0.0%	3	100.0%	3	DF=2 NS	
Family type	Nuclear family	9	37.5%	15	62.5%	24	χ2=5.90 P=0.02*	
	Joint family	4	11.1%	32	88.9%	36	DF=1 S	
Education	Elementary education	3	23.1%	10	76.9%	13	χ2=0.37 P=0.94	
	Middle education	3	27.3%	8	72.7%	11	DF=3 NS	
	Higher secondary education	4	18.2%	18	81.8%	22		
	Graduate	3	21.4%	11	78.6%	14		
Job	Home maker	13	24.1%	41	75.9%	54	χ2=1.84	
	Government job	0	0.0%	1	100.0%	1	P=0.61 DF=3NS	
	Private job	0	0.0%	2	100.0%	2		
	Self job	0	0.0%	3	100.0%	3		
Income	< Rs.10000	9	24.3%	28	75.7%	37	χ2=0.40	
	Rs.10001 - 20000	4	17.3%	19	82.7%	23	P=0.52 DF=1 NS	
Age at	< 20 years	6	31.6%	13	68.4%	19	χ2=3.31	
marriage	21 - 25 years	7	21.2%	26	78.8%	33	P=0.19	
	> 25 years	0	0.0%	8	100.0%	8	DF=2 NS	
Gravida	Primi	10	31.3%	22	68.7%	32	χ2=3.96	
	Multi	3	10.7%	25	89.3%	28	P=0.05* DF=1 S	

Table 4.12:Association between posttest level of knowledge score withdemographic and obstetric variables:

** highly significant at P≤0.01

*** very highly significant at $P \le 0.001$

Table 4.12 shows the association between post-test level of knowledge score and antenatal women demographic variables. When considering the age 76.0% of 20-28 years age group gained adequate knowledge and 100% of above 28 years age group gained adequate knowledge. This is statistically significant(P value=0.03).

When taking family type 62.5% of nuclear family women have gained adequate knowledge and 88.9% of joint family women have gained adequate knowledge. This is also statistically significant.(P value= 0.02).

When considering the gravid 68.7% of primi gravid women have gained adequate knowledge and 89.3% of multi gravid women have gained adequate knowledge. This is also statistically significant(P value= 0.05).

So, elder, joint family and multi gravid women gained more knowledge than others after STP. It was confirmed using Chi square test.

		Knowledge gain score							0	
Demographic variables		Pretest		Posttest		Knowledge gain= post- pre		Total	Oneway ANOVA F-test/ t- test	
		Ν	%	n	%	Ν	%		lesi	
Age	< 20 years	14.00	0.00	23.00	0.00	9.00	0.00	1	F=3.12	
	20 - 28 years	11.26	4.45	23.28	2.31	12.02	4.68	50	P=0.05*	
	> 28 years	11.11	5.86	27.09	1.09	15.98	5.88	9	S	
Religion	Hindu	11.65	4.72	25.00	2.27	13.35	4.87	54	F=0.71	
	Muslim	8.00	1.73	24.67	2.08	16.67	2.31	3	P=0.49	
	Christian	11.67	4.73	26.00	2.00	14.33	5.51	3	NS	
Family type	Nuclear family	11.33	4.45	23.54	2.12	12.21	4.49	24	t=1.99 P=0.05*S	
	Joint family	11.56	4.83	26.31	2.28	14.75	5.06	36		
Education	Elementary education	7.54	3.10	25.31	2.14	17.77	2.89	13	F=1.76 P=0.14	
	Middle education	10.73	5.24	25.48	1.94	14.75	4.93	11	NS	
	Higher secondary education	11.05	4.80	26.12	2.63	15.07	2.30	22		
	Graduate	13.21	2.69	27.30	1.91	14.09	2.97	14		
Job	Home maker	11.50	4.72	24.89	2.23	13.39	4.87	54	F=0.31 P=0.81	
	Government job	13.00		30.00		17.00	•	1	NS	
	Private job	10.00	5.66	25.50	.71	15.50	6.36	2		
	Self job	11.33	5.03	25.67	.58	14.33	4.51	3		
Income	< Rs.10000	11.05	4.70	25.05	2.12	14.00	4.91	37	t=0.88	
	Rs.10001 - 20000	12.13	4.57	25.00	2.43	12.87	4.64	23	P=0.33 NS	
Age at marriage	< 20 years	12.63	5.18	24.53	2.22	11.89	5.58	19	F=2.28	
	21 - 25 years	11.21	4.47	25.18	2.38	13.97	4.28	33	P=0.11	
	> 25 years	9.75	3.73	25.63	1.41	15.88	4.02	8	NS	
Gravida	Primi	12.00	4.24	23.30	2.33	11.30	4.39	32	t=2.05	
	Multi	11.00	5.08	25.95	2.12	14.95	4.30	28	P=0.05* S	

Table-4.13 Association between post-test level of knowledge score with demographic and obstetric variables:

** highly significant at $P \leq 0.001$.

Table 4.13 represents the association between post-test level of knowledge score with antenatal women demographic variables.

In considering the **age**, 20-28 years age group gained knowledge score of 4.68%, more than 28 years gained knowledge score of 5.88%. P value is 0.05. This is statistically significant.

In **family type**, 4.49% knowledge gain score in nuclear family type, and 5.06% knowledge gain score in joint family type. P value is 0.05. This is also statistically significant.

In considering the **Gravida**, 4.39% of knowledge gain score in primi gravida, and 4.30% of knowledge gain score in multi gravida mothers.P value is 0.05. This is also statistically significant.

CHAPTER-V SUMMARY OF RESULTS

The study was done to determine the effectiveness of the effectiveness of Structured Teaching Programme on knowledge regarding early identification of high risk pregnancy among antenatal mothers attending at Institute of Obstetrics and Gynaecology.

5.1. Based on demographic findings

The study findings reveal the following demographic and obstetric characteristic features of 60 antenatal mothers who participated in the study.

- Age wise distribution ,83.3% were 20-28 years of age, 15% were in more than 28 years of age group, and remaining 1.7% belonged to less than 20 years of age.
- In religion, 90% are Hindus, 5% belongs to Christian religion and remaining 5% are of Muslim religion.
- In family type, 60% were in joint family and remaining 40% nuclear family.
- In education wise, 36.7% (22) had Higher secondary education, 23.3% (14) were graduates, 21.7%(13) had elementary education and 18.3%(11) were of middle education status.
- Job wise, 90%(54) of them are home makers, 5%(3), of them had self job, and 3.3%(2) of them working in private sector and only 1.7%(1) were in Government jobs.
- In the monthly income, 61.1%(37) family monthly income was below Rs.10,000 and remaining 38.8%(23) monthly income was Rs. 10001-20000.

- In age at marriage, 55% (55) were married at the age of 21-25 years, and 31.7%((19) were married at the age of below 20 years and remaining 13.3%(8) were married more than 25 years of age.
- In gravida, 53.3%(32) were primi and remaining 46.7%(28) were multi.

5.2. Based on Pre-test knowledge of antenatal mothers in the study group

The percentage of pretest knowledge score in each domain wise. They are having more score in anemia(41.9%), and moderate score of 38.8% in pre-eclampsia and 36.7% in gestational diabetes mellitus, and minimum score in multiple pregnancy(35.0%) Overall they are having 38.2% of score.

The level of pretest knowledge regarding early identification of high risk pregnancy among antenatal women before administration of structured teaching programme. 76.7%(46) of the antenatal women had inadequate knowledge score, 23.3%(14) of them had moderate knowledge score, and none of them are having adequate level of knowledge score.

5.3. Based on post-test knowledge of antenatal mothers on high risk pregnancy in the study group

Each domain wise post-test percentage of knowledge score regarding early identification of high risk pregnancy among antenatal women. They are having more score in Anemia(84.8%), and minimum score in preeclampsia(81.5%), score in Gestational Diabetes Mellitus(84.7%), and score in Multiple pregnancy (82.9%). Overall they are having 83.4% of score.

The level of posttest knowledge regarding early identification of high risk pregnancy among antenatal women after administration of

55

Structured teaching programme. 78.4%(47) of them having adequate level of knowledge score , 21.6%(13) of them are having moderate knowledge score, None of the antenatal women are having inadequate knowledge score.

5.4. Based on the effectiveness of structured teaching programme on knowledge regarding high risk pregnancy in the study group

Considering anemia aspects, in pretest antenatal mothers are having 3.45 score where as in posttest they are having 6.78 score, so the difference is 3.43. This difference between pretest and posttest is large and is statistically significant.

Considering preeclampsia aspects, in pre-test antenatal mothers are having 3.10 score where as in posttest they are having 6.52 score, so the difference is 3.42. This difference between pretest and posttest is large and it is statistically significant.

Considering Gestational Diabetes Mellitus aspects, in pretest, antenatal mothers are having 2.57 score where as in posttest they are having 5.93, so the difference is 3.37. This difference between pretest and posttest is large and it is statistically significant.

Considering Multiple Pregnancy aspects, in pretest antenatal mothers are having 2.45 score where as in posttest they are having 5.80 score, so the difference is 3.35. This difference between pretest and posttest is large and it is statistically significant.

Statistical significance was calculated by using student's paired 't' test.

The comparison of overall pretest and posttest knowledge score. Considering overall knowledge score, in pretest antenatal women are having 11.47 score where as in posttest they are having 25.03, so the difference is 13.57.The difference between pretest and posttest score is large and it is statistically significant. Differences between pretest and posttest score was analysed using students paired 't' test.

In post-test, none of the antenatal women are having inadequate knowledge score, 21.6% of them having moderate knowledge score, 78.4% of them are having adequate level of knowledge score.

Statistically there is significant difference between pretest and posttest knowledge score.

Each domain wise percentage of knowledge gain. 42.9% of knowledge gain anemia aspects, 42.7% knowledge gain in preeclampsia aspects, and 48% knowledge gain in Gestational diabetes mellitus and 47.9% of knowledge gain in Multiple pregnancy aspect. Overall they gained 45.2% of knowledge score when comparing pretest and posttest after having STP. This shows the effectiveness of STP on knowledge regarding high risk pregnancy among antenatal women.

5.5. Based on association between posttest score with selected demographic and Obstetrical variables

In considering the **age**, 20-28 years age group gained knowledge score of 4.68%, more than 28 years gained knowledge score of 5.88%. P value is 0.05. This is statistically significant.

In **family type**, 4.49% knowledge gain score in nuclear family type, and 5.06% knowledge gain score in joint family type. P value is 0.05. This is also statistically significant.

In considering the **Gravida**, 4.39% of knowledge gain score in primi gravida, and 4.30% of knowledge gain score in multi gravida mothers.P value is 0.05. This is also statistically significant.

CHAPTER-VI DISCUSSION

In this study the researcher has made an attempt to identify the common high risk among antenatal mothers who are attending antenatal outpatient department. Prevention of many high risk pregnancies and its complications is very essential for successful outcome of healthy mother and healthy child.

The purpose of the study was to evaluate the effectiveness of Structured teaching programme on high risk pregnancy among antenatal women. The sample size for this study was 60 antenatal mothers who were labeled into one group. Pretest was conducted in each participants by interview method with the use of Structured questionnaire. Structured teaching programme on early identification of high risk pregnancy was given nearly 30 minutes. One week after STP posttest was conducted to assess the effectiveness of teaching programme.

The results of the study were based on the statistical analysis. The effectiveness of teaching programme was assessed by using Paired 't' test. Chi-Square was used to find the association between the level of knowledge with selected demographic and obstetric variables. The results were formulated according to the stated objectives.

The details of demographic characteristics of 60 antenatal mothers who participated in the study were as follows.

In considering the **age** wise distribution of antenatal mothers,83.3% were 20-28 years of age, 15% were in more than 28 years of age group, and remaining 1.7% belongs to less than 20 years of age.

In the **religion** wise distribution, 90% are Hindus, 5% belongs to Christian religion and remaining 5% are of Muslim religion.

In considering the **family type**, 60% were of joint family and remaining 40% belonged to nuclear family.

In **education** wise distribution, 36.7% (22) had Higher secondary education, 23.3% (14) were graduates, 21.7%(13) had elementary education and 18.3%(11) only in middle education status.

In considering the **job**, 90%(54) of them are home makers, 5%(3), of them had self job, and 3.3%(2) of them working in private sector and only 1.7%(1) in Government jobs.

In the **monthly income** status, 61.1%(37) family monthly income was below Rs.10,000 and remaining 38.8%(23) monthly income was Rs. 10001-20000.

In considering the Obstetrical variables, in **age at marriage**, 55% (55) were married at the age of 21-25 years, and 31.7%((19) were married at the age of below 20 years and remaining 13.3%(8) were married more than 25 years of age.

In **gravida**, 53.3%(32) were primi and remaining 46.7%(28) were multi.

To assess the Knowledge regarding high risk pregnancy among antenatal women.

The level of knowledge of antenatal women on high risk pregnancy was assessed by using Structured questionnaire. It denotes the percentage of pretest knowledge score each domain wise. They are having more score in anemia(41.9%), and moderate score of 38.8% in pre-eclampsia and 36.7% in Gestational diabetes mellitus, and minimum score in multiple pregnancy(35.0%). Overall they are having 38.2% of score.

60

The percentage of pretest knowledge regarding early identification of high risk pregnancy among antenatal women before administration of structured teaching programme. 76.7%(46) of the antenatal women had inadequate knowledge score, 23.3%(14) of them had moderate knowledge score, and none of them are having adequate level of knowledge score.

(Singh. N.J 2010) A study was conducted at the urban centers in Bangalore, to assess the effectiveness of a structured teaching programme on warning signs in pregnancy among 60 primi gravid women. The results indicated that the mean percentage of pretest knowledge score was 25.58%, which was considered to be lower of mean percentage.⁵²

To assess the effectiveness of Structured teaching programme on knowledge regarding high risk pregnancy among antenatal women.

A Structured teaching programme on high risk pregnancy was conducted for 30 minutes. Posttest was conducted with the same questionnaire on 7th day.

It was evident that the percentage of posttest knowledge scores on level of knowledge among antenatal women on high risk pregnancy. It denotes that in post-test 84.8% of them adequate knowledge on anemia, 81.5% of them are having adequate knowledge on preeclampsia, whereas in gestational diabetes mellitus 84.7% of them are having adequate knowledge, and in multiple pregnancy, 82.9% of them are having adequate knowledge.They are having more score in anemia (84.8%) and minimum score in Pre-eclampsia(81.5%). Overall they are having 83.4% of knowledge score.

It shows the posttest knowledge score among 60 antenatal women, 13 of them (21.6%) are having moderate knowledge and 47 of them (78.4%) are having adequate knowledge, and none of them are having inadequate knowledge on high risk pregnancy.

It shows the comparison of overall knowledge score between pretest and posttest. On an average, in posttest, antenatal women gained 45.2% of knowledge score after having structured teaching programme regarding early identification of high risk pregnancy. Differences between pre-test and post-test score was analysed using proportion with 95% confidence interval and mean difference with 95% confidence interval. This 42.5% of knowledge gain shows the effectiveness of structured teaching programme on early identification of high risk pregnancy among antenatal women.

The findings of the present study reveals the domain wise percentage of knowledge gain. Overall they gained 45.2% of knowledge score when compared pre-test and post-test after having STP. This shows the effectiveness of STP on knowledge regarding high risk pregnancy among antenatal women.

To evaluate the effectiveness of structured teaching programme the pretest and posttest values are compared. The calculated 't' value is greater than the tabulated value at 5% level of significance. It shows that the teaching was effective.

(**Irbaihat. Jamil2009**) A study was conducted at the antenatal clinic of St. John's Medical College Hospital in Bangalore, to assess the effectiveness of a Structured teaching programme on high risk pregnancy, among 100 antenatal women. The results indicated that the mean percentage of pretest knowledge score was 45.6%, which was considered to be the average. The mean post-test knowledge score(t=44.04). It could be presumed that the teaching programme on high risk

62

conditions in pregnancy was effective in enhancing the knowledge of the mother.⁵³

To associate the findings with selected demographic and Obstetric variables.

It revealed the association between the level of knowledge on high risk pregnancy with selected demographic and obstetrical variables such as age, religion, family type, education, job, income, age at marriage and gravida.

It shows the association between pretest level of knowledge score with clients demographic variables. None of the demographic variables are significantly associated with their pretest level of knowledge. It was confirmed using Chi square test.

The findings of the present study shows the association between posttest level of knowledge score with antenatal women demographic variables. When considering the age 76.0% of 20-28 years of age group gained adequate knowledge and 100% of above 28 years age group gained adequate knowledge. This is statistically significant(P value=0.03).

When taking family type 62.5% of nuclear family ,women have gained adequate knowledge and 88.9% of joint family women have gained adequate knowledge. This is also statistically significant.(P value= 0.02).

When considering the gravid 68.7% of primi gravid women have gained adequate knowledge and 89.3% of multi gravid women have gained adequate knowledge. This is also statistically significant(P value= 0.05).

63

So, elder, joint family and multi gravid women gained more knowledge than the others after STP. It was confirmed using Chi square test.

(Christ Anees 2008) A study was conducted in Ludhiana, to assess the knowledge of urban mothers about high risk conditions in pregnancy and relationship of knowledge with education, age group and occupation. 40 samples were selected for this study. The findings revealed that the mothers possessed high level of mean percentage score(70.62%). The mothers with highest level of education obtained the highest level of mean percentage knowledge scores(88.4%). Those with VIII pass had low level of mean percentage knowledge scores(36.25%). Younger group had (17-26 years) had high level of mean percentage of knowledge scores (48%). In the same way, the mothers who had been working had a higher knowledge score than the housewives.⁵⁴

(Mahalingam G, Venkatesan M 2014)A descriptive design with cross sectional survey approach was conducted in Dehradun to assess the mothers knowledge on warning signs of pregnancy, labour and puerperium. Data was collected from 50 antenatal mothers by purposive sampling technique and questionnaire was used as a data collection tool. Result of this study shows that 36% of the mothers has poor knowledge, 24% has average knowledge,34% has good knowledge and only 6% had excellent knowledge about warning signs of pregnancy, labour, and puerperium. The overall knowledge percentage mean which was 50.73% revealed that the antenatal mothers has average knowledge about warning signs of pregnancy,labour and puerperium. The knowledge of antenatal mothers is statistically , not associated with their age, educational qualification, parity, occupation, family income, type of family, religion, residential areas and previous source of information. They concluded to conduct the study in larger samples.⁵⁵

Hypothesis

H: Antenatal women who received structured teaching programme on high risk pregnancy shows a significant increase in the level of knowledge in post-test than pre-test.

From the findings of the present study it was concluded that the structured teaching programme on early identification of high risk pregnancy improves the knowledge of antenatal mothers who participated in the study. **Thus the hypothesis was proved statistically.**

CHAPTER-VII CONCLUSION AND RECOMMENDATIONS

7.1 Implications of the study

The investigator has drawn the following implication for the study which is vital concern in the field of nursing practice, nursing education, nursing administration and nursing research.

Some of the implications for the present study in various areas as follows:

Nursing practice

- The nurses working in the health care services should update the knowledge about current trends and treatments in high risk pregnancy.
- Structured teaching programme on high risk pregnancy is to be scheduled in the obstetrical department.
- Self instructional module can be distributed to the high risk groups who are visiting the hospital.

Nursing education

- Conferences, workshops and seminars can be held for impart, to update the knowledge and positive attitudes.
- In-service education updating their knowledge and skill in various health care settings can be given.
- Nursing curriculum has to focus on enabling the nursing students to develop skill in identifying risk groups and prevent the complications.

- Every student can be motivated to plan and give health education for antenatal women in high risk pregnancy.
- Short term courses can be organized for the nurses, who are working in obstetrical unit.

Nursing administration

The present study is proposed to help the health administrators to strategically plan and meet the health needs of the antenatal women.

- The administrators in both private and government sectors should take initiative actions to update the knowledge of risk groups on high risk pregnancy.
- The administrator can encourage the nurses for conducting research in various aspects high risk pregnancy.
- The administrator can organize conference, workshop and seminars for nurses working in the community.
- The administrator should support the staffs to conduct programme on high risk pregnancy.
- The nursing administration should awaken to the fact that patient education is a necessity and should provide resources in terms of manpower, money and material.

Nursing research

- The study will be a valuable reference material for further researchers.
- This study is a preliminary set up for exploring the concept of knowledge on high risk pregnancy.

The results of the study encourage the antenatal women to adopt healthy life.

7.2 Limitations

- The sample size is limited to 60 antenatal mothers.
- The study was limited to mothers only attending antenatal out patient department at Institute of Obstetrics and Gynaecology.

7.3 Recommendations for further study

The study recommends the following,

- A similar study can be undertaken for a large sample in different settings.
- A comparative study can be conducted between urban and rural antenatal women.
- A similar study can be conducted among antenatal women in community area.
- ✤ A similar study can be conducted using pre-test and post- test with control group.
- Experimental study can be conducted on anemia, pregnancy induced hypertension, gestational diabetes mellitus and multiple pregnancy and its management for high risk women, in terms of gain in knowledge and control of symptoms.

Conclusion

Nursing personnel must have holistic knowledge regarding the different aspects of high risk pregnancy among antenatal mothers. Nurses play a vital role in the teaching aspects of high risk pregnancy.The present study had been supported by a series of other studies which confirmed that the knowledge on high risk pregnancy among antenatal women is important to get healthy baby and healthy mother. From the analysis and results, it was found that Structured teaching programme on high risk pregnancy is an essential intervention to identify the high risk pregnancy in the earlier stage and prevent from complications of high risk pregnancy.

REFERENCES

- Narayan B, Nelson-Piercy C. Medical problems in pregnancy. Clin Med (Lond). 2016 Dec;16(Suppl 6):s110-s116.
- 2) Cooper, (2012).*Pregnancy*, 5th edition. London Pen Book publishers.
- Barley.E.Rose (2010). Obstetrics and Gynaecology Nursing, 10th edition, Baltimore Willium and Wilkens company.
- 4) Pilliteri Adele.(2003). *Maternal and Child Health Nursing*, 4th edition, Philadelphia Lippincot Williams & Wilkims.
- 5) Samiya M, Samina M. Identification of high risk pregnancy, Indian Journal for the Practicing Doctor, Vol. No. 1 (2008-03 – 2008-04).
- 6) http://www.faqs.org/health/topics/3/High-risk pregnancy.html
- Arulkumaran.S. Retnam (2012). *The management of labour*, 4th edition, Chennai Orient Longman Ltd.
- 8) Reddy, S. R. Study to evaluate the perinatal care among high risk mothers in selected hospitals of Belgaum. The Nursing Journal of India.
- Suresh k. Sharma (2015). Nursing Research and Statistics, 2nd edition, Published by Reed Elsevier India Private Limited.
- Samar K. Hafez, Laila Sh. Dorgham and Suheir A.M. Sayed. Profile of High Risk Pregnancy among Saudi Women in Taif-KSA. World Journal of Medical Sciences 11 (1): 90-97, 2014 ISSN 1817-3055.

- 11) Dutta S and Das XS. Identification of high risk mothers by a scoring system and it's correlation with perinatal outcome, Journal of Obstet Gynaecol India. 2012; 40: 181-190.
- 12) Shahzada J. Malik and Nisar A. Mir. Identify the preventable factors in high risk pregnancy, Journal of obstetrics and gynecology research, May 2010.
- Dr. MahmudaMubasher. To find out prevalence and outcome of high risk pregnancies, OBG Department, MAROOF International Hospital in Pakistan.
- Taehan Kanho. Identification of high-risk pregnancy, using a simplified risk-scoring system at Chung Ang Medical Center, Korea", Journal of obstetrics and gynecology, 2010 Aug; 30(3):49-65.
- 15) Yara Almerie et al., Near-miss and maternal mortality in maternity university hospital, Syria, BMC Pregnancy and Childbirth 2010, 10.1186/1471-2393-10-65.
- 16) Jimenez K, Lang M. Diagnostic approach to iron deficiency anemia. Wien Med Wochenschr. 2016 Oct;166(13-14):402-410. Epub 2016 Aug 26.
- 17) Shander A, Goodnough LT, Javidroozi M, Auerbach M, Carson J, Ershler W, et al., *Iron deficiency anemia--bridging the knowledge* and practice gap. Transfus Med Rev. 2014 Jul;28(3):156-66. doi: 10.1016/j.tmrv.2014.05.001. Epub 2014 May 15.
- 18) Breymann C. Iron Deficiency Anemia in Pregnancy. Semin Hematol. 2015 Oct;52(4):339-47. doi: 10.1053/j.seminhematol.2015.07.003. Epub 2015 Jul 10.

- 19) Rahman MM et al., Maternal anemia and risk of adverse birth and health outcomes in low- and middle-income countries:. Am J Clin Nutr. 2016 Feb;103(2):495-504. doi: 10.3945/ajcn.115.107896. Epub 2016 Jan 6.
- 20) Burke RM, Leon JS, Suchdev PS. Identification, prevention and treatment of iron deficiency during the first 1000 days. Nutrients.
 2014 Oct 10;6(10):4093-114. doi: 10.3390/nu6104093.
- 21) De-Regil LM, Peña-Rosas JP, Fernández-Gaxiola AC, Rayco-Solon P, et al., *Effects and safety of periconceptional oral folate supplementation for preventing birth defects. Cochrane Database Syst Rev.* 2015Dec14;(12): CD007950. doi:10.1002/ 14651858. CD007950. pub3.
- Taylor S, Rampton D. Treatment of iron deficiency anemia: practical considerations. Pol Arch Med Wewn. 2015;125(6):452-60. Epub 2015 Apr 29.
- 23) Orlov IuP, Lukach VN, Govorova NV. Iron metabolism in women with anemia and eclampsia (Part I). Anesteziol Reanimatol. 2014 Nov-Dec;59(6):67-72.
- Shi Q, Leng W, Wazir R, Li J, Yao Q, Mi C, et al., Intravenous Iron Sucrose versus Oral Iron in the Treatment of Pregnancy with Iron Deficiency Anaemia. Gynecol Obstet Invest. 2015;80(3):170-8. doi: 10.1159/000376577. Epub 2015 Mar 25.
- 25) Hlimi T. Association of anemia, pre-eclampsia and eclampsia with seasonality Epub 2014 Dec 31.
- 26) Kidson-Gerber, Kerridge I, Farmer S, Stewart CL, Savoia H, Challis D. Caring for pregnant women for whom transfusion is

not an option. Aust N Z J Obstet Gynaecol. 2016 Apr;56(2):127-36. doi: 10.1111/ajo.12420. Epub 2015 Nov 17.

- 27) Rukuni R, Knight M, Murphy M, Roberts D, Stanworth SJ. Screening for iron deficiency and iron deficiency anaemia in pregnancy BMC Pregnancy Childbirth. 2015 Oct 2 0;15:269. doi: 10.1186/s12884-015-0679-9.
- 28) Dhariwal NK, Lynde GC. Update in the Management of Patients with Preeclampsia. Anesthesiol Clin. 2017 Mar;35(1):95-106. doi: 10.1016/j.anclin.2016.09.009. Epub 2016 Dec 12.
- 29) Spradley FT, Palei AC, Granger JP. Increased risk for the development of preeclampsia in obese pregnancies: weighing in on the mechanisms Am J Physiol Regul.
- 30) Singh MD, Thomas P, Owens J, Hague W, Fenech M. Potential Role of folate in pre-eclampsia. Nutr Rev. 2015 Oct;73(10):694-722. doi: 10.1093/nutrit/nuv028. Epub 2015 Sep 10.
- Mudaliar and Menon. (2015), Clinical Obstetrics, 12th edition
 Chennai Orient Longman Pvt ltd.
- 32) Cormick G, Betrán AP, Ciapponi A, Hall DR, Hofmeyr GJ.calcium and Pre-eclampsia Study Group. Inter-pregnancy interval and risk of recurrent pre-eclampsia. Reprod Health. 2016 Jul 18;13(1):83. doi: 10.1186/s12978-016-0197-x.
- 33) Vigil-De Gracia P, Ludmir J. The use of magnesium sulfate for women with severe preeclampsia or eclampsia diagnosed during the postpartum period. J Matern Fetal Neonatal Med. 2015;28(18):2207-9. doi: 10.3109/14767058.2014.982529. Epub 2014 Nov 27.

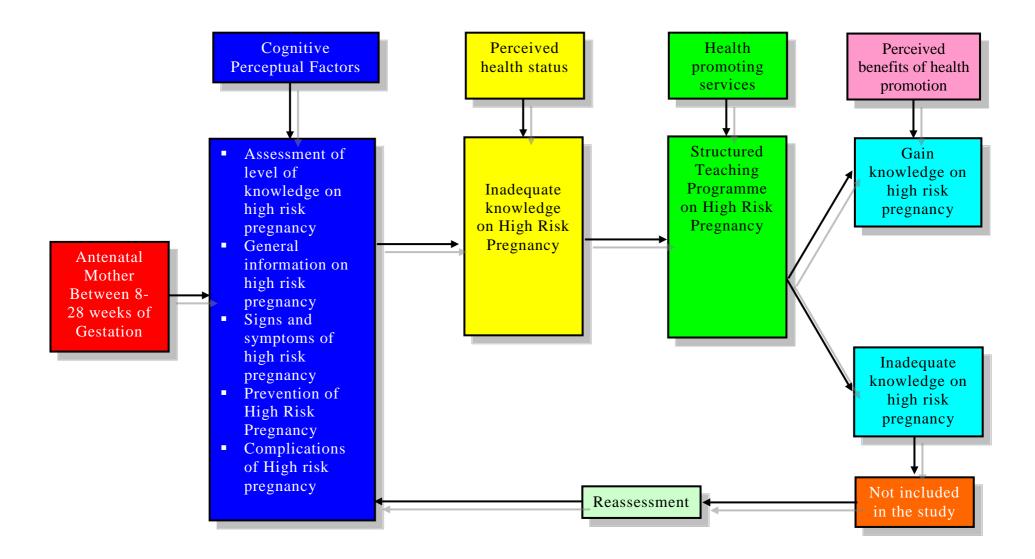
- 34) Brown CM, Garovic VD. Drug treatment of hypertension in pregnancy. Drugs. 2014 Mar;74(3):283-96. doi: 10.1007/s40265-014-0187-7.
- 35) Meher S, Duley L, Hunter K, Askie L. Antiplatelet therapy before or after 16 weeks' gestation for preventing preeclampsia. Am J Obstet Gynecol. 2017 Feb;216(2):121-128.e2. doi: 10.1016/j.ajog.2016.10.016. Epub 2016 Nov 1.
- 36) Taschereau-Charron A, Da Silva MS, Bilodeau JF, Morisset AS, Julien P, Rudkowska I. Alterations of fatty acid profiles in gestational diabetes and influence of the diet. Maturitas. 2017 May;99:98-104. doi: 10.1016/j.maturitas.2017.01.014. Epub 2017 Jan 27.
- 37) Rozenberg P. J Gynecol Obstet Biol Reprod (Paris). 2016
 Nov;45(9):1037-1044. doi: s10.1016/j.jgyn.2016.09.001. Epub
 2016 Oct 19.
- 38) Harrison AL, Shields N, Taylor NF, Frawley HC. Exercise improves glycaemic control in women diagnosed with gestational diabetes mellitus. J Physiother. 2016 Oct;62(4):188-96. doi: 10.1016/j.jphys.2016.08.003. Epub 2016 Aug 22.
- 39) McCance DR. Diabetes in pregnancy. Best Pract Res Clin Obstet Gynaecol. 2015 Jul;29(5):685-99. doi: 10.1016/j. bpobgyn. 2015.
 04.009. Epub 2015 Apr 28.
- 40) Valentina Palkova J. Effectiveness of an educational program on gestational diabetes mellitus. Cochrane Database Syst Rev. 2011;12:CD004736.
- 41) Tieu J, Shepherd E, Middleton P, Crowther CA. Dietary advice interventions in pregnancy for preventing gestational diabetes

mellitus .*Cochrane Database Syst Rev.* 2017 Jan 3;1:CD006674. doi: 10.1002/14651858.CD006674.pub3.

- Farrar D, Simmonds M, Bryant M, Sheldon TA, Tuffnell D, Golder S, et al., *Hyperglycaemia and risk of adverse perinatal outcomes*. BMJ. 2016 Sep 13;354:i4694. doi: 10.1136/bmj.i4694.
- Moldenhauer JS, Johnson MP. Diagnosis and Management of Complicated Monochorionic Twins Clin Obstet Gynecol. 2015 Sep;58(3):632-42. doi: 10.1097/GRF.000000000000127.
- 44) Tran S, Whitworth M, Bricker L, Mullan C. Survival rate of delayed- interval delivery in multifoetal pregnancy. Cochrane Database Syst Rev. 2015 Jul 14;(7):CD007058. doi: 10.1002/14651858.CD007058.pub3.
- 45) Joshua, p. Vogel. Association between twins and adverse maternal and perinatal outcomes. Reprod Health. 2013 Sep 26;11
 Suppl 3:S2. doi: 10.1186/1742-4755-11-S3-S2. Epub 2013 Sep 26.
- 46) Lee, K.E. Critical Review of the Literature: Parental Grief after the Loss of a Multiple. Journal of Neonatal Nursing, 18(6): 226-31. doi:10.1016/j.jnn.2011.12.002
- Richards, J,Graham, R. H. Embleton, N. D., & Rankin, J. Health Professionals' Perspectives on bereavement following Loss from a Twin Pregnancy. Journal of Perinatology. doi:10.1038/jp.2016.13
- 48) Whitford HM, Wallis SK, Dowswell T, West HM, Renfrew MJ. Breastfeeding education and support for women with twins or higher order multiples. Cochrane Database Syst Rev. 2017 Feb 28;2:CD012003. doi: 10.1002/14651858.CD012003.pub2.

- 49) Kushnir VA, Barad DH, Albertini DF, Darmon SK, Gleicher N. Systematic review of worldwide trends in assisted reproductive technology 2004-2013. Reprod Biol Endocrinol. 2017 Jan 10;15(1):6. doi: 10.1186/s12958-016-0225-2.
- 50) Lumbiganon P, Martis R, Laopaiboon M, Festin MR, Ho JJ, Hakimi M. Antenatal breastfeeding education for increasing breastfeeding duration. Cochrane Database Syst Rev. 2016 Dec 6;12:CD006425. doi: 10.1002/14651858.CD006425.pub4.
- 51) Moore ER, Bergman N, Anderson GC, Medley N. Early skin-toskin contact for mothers and their healthy newborn infants. Cochrane Database Syst Rev. 2016 Nov 25;11:CD003519.
- 52) Singh.N.J .Knowledge of urban mothers about high risk conditions in pregnancy. The Nursing Journal of India 5(2), 108-110.
- 53) Irbaihat.Jamil. The Effect of a Health Education Programme among pregnant women American Journal of Obstetrics and Gynaecology, 189, 934-938.
- 54) Christ Anees. Safe Motherhood .The Nursing journal of India 2008 14-71.
- 55) Mahalingam G, Venkatesan M. Mother's knowledge of warning signs of pregnancy, labour and puerperium.Int/ med sci/ public health 2014;3:720-722

FIG: 2.3: MODIFIED PENDER'S HEALTH PROMOTION MODEL



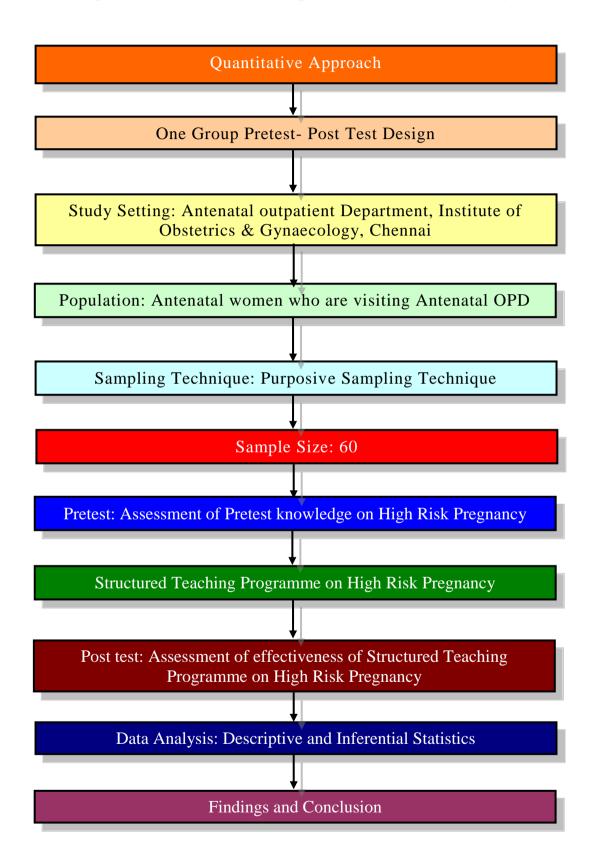


Figure: 3.5: Schematic Representation of the study

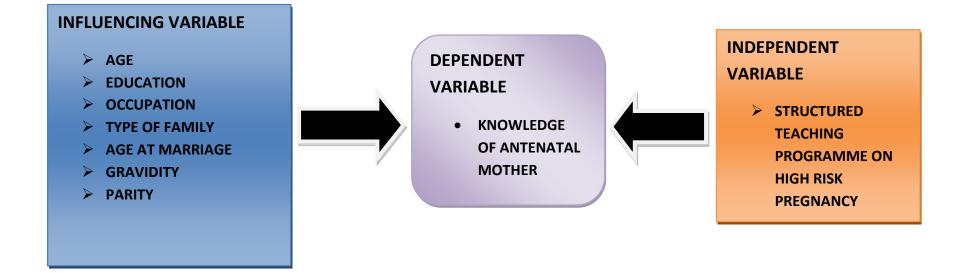


Figure 3.4: Schematic representation of relationship between variables

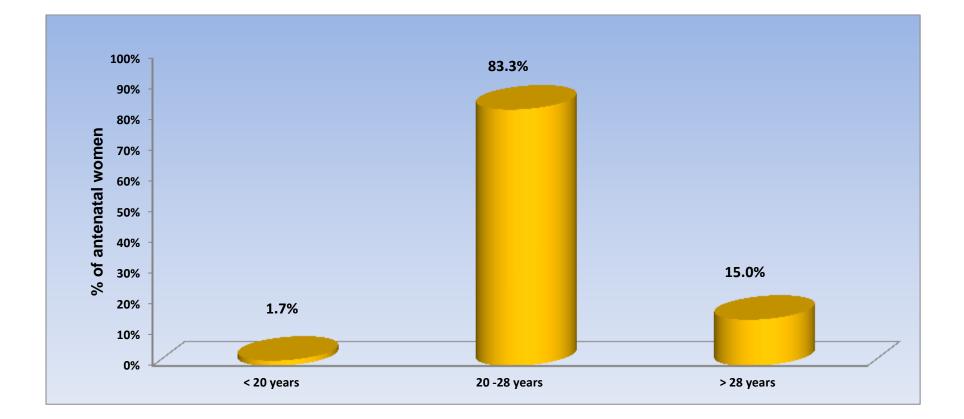


Figure 4.6: Age wise distribution of antenatal mothers

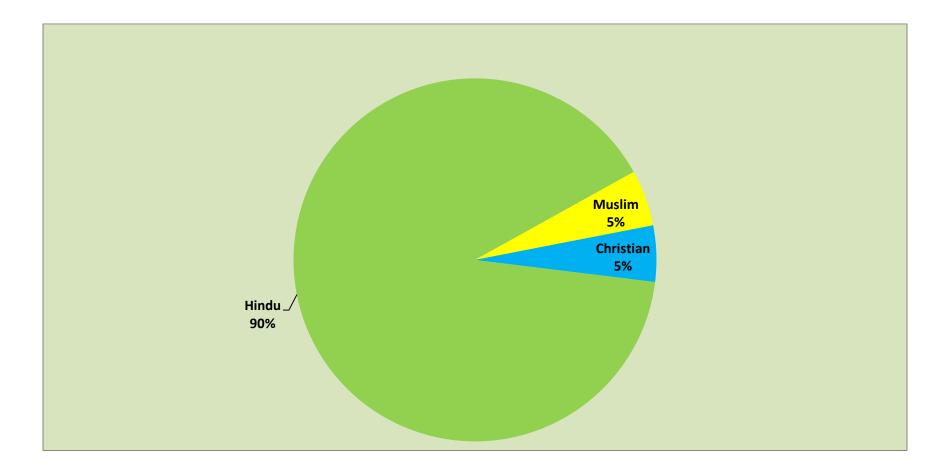


Figure 4.7: Religion wise distribution of antenatal mothers.

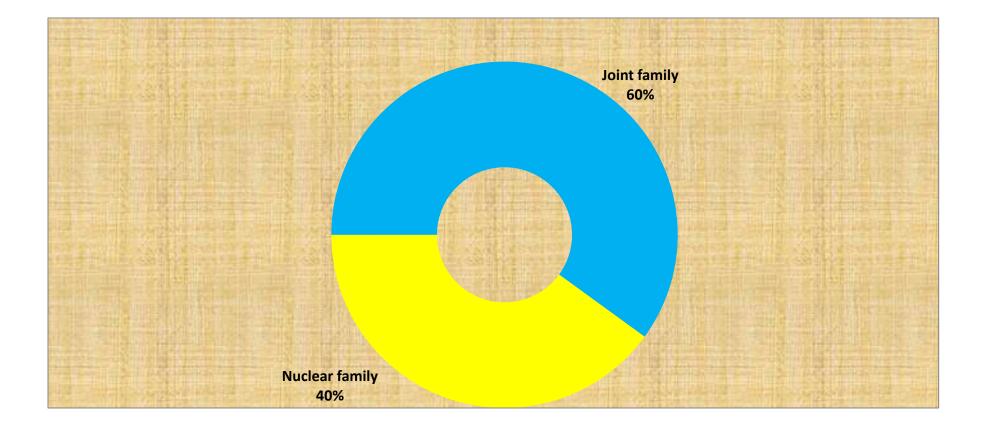


Figure 4.8: Type of family system of antenatal mothers.

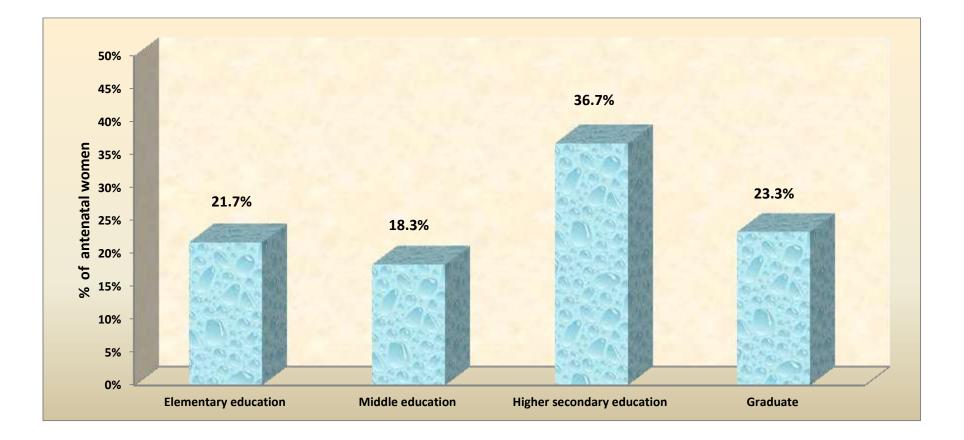


Figure 4.9: Educational status of antenatal mothers.

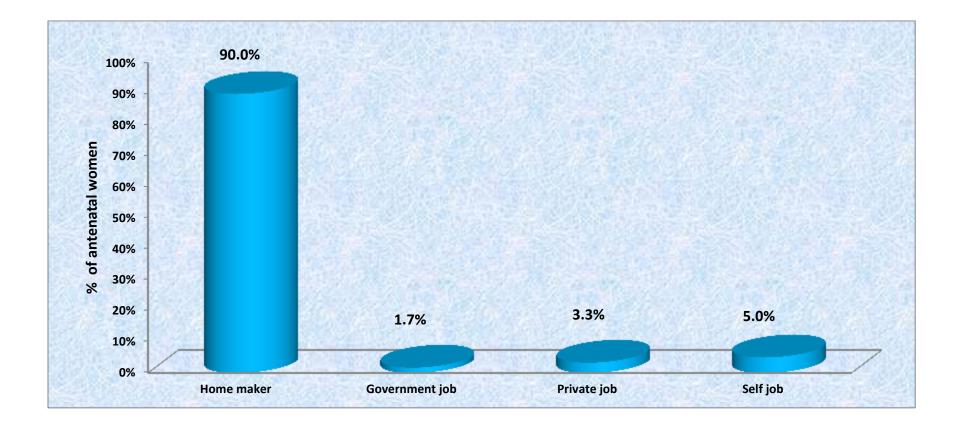


Figure 4.10: Occupational status of the antenatal mothers.

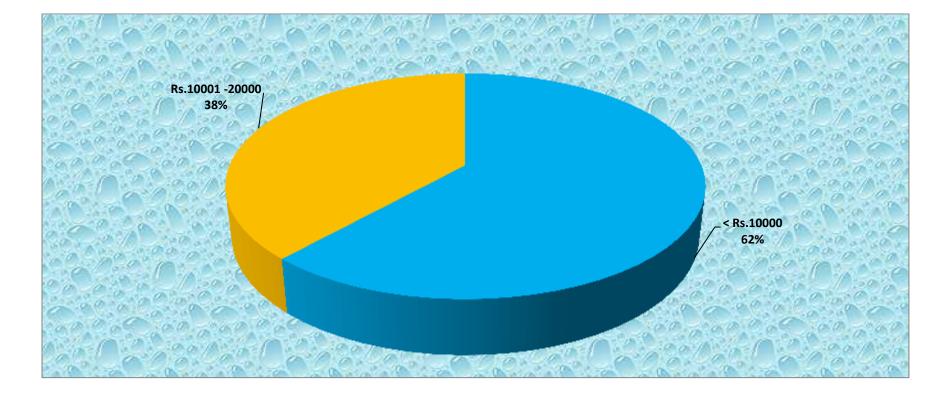


Figure 4.11: Monthly income status of antenatal mothers.

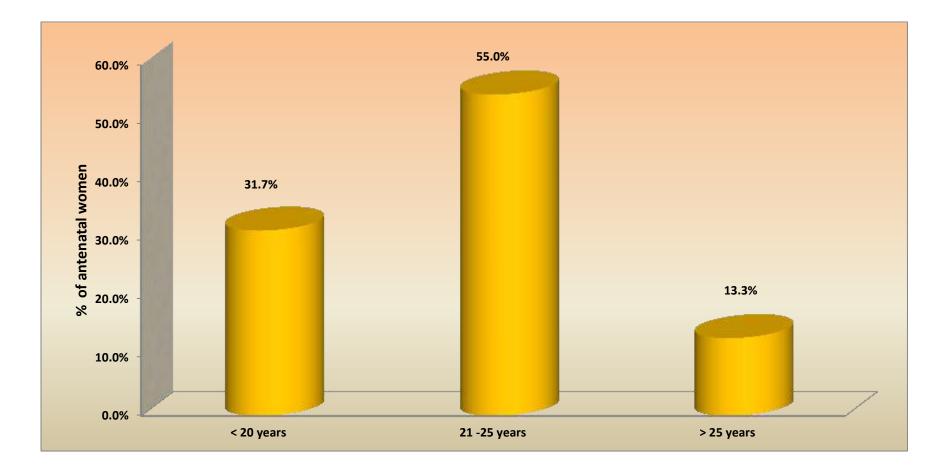


Figure 4.12: Age at marriage of antenatal mother

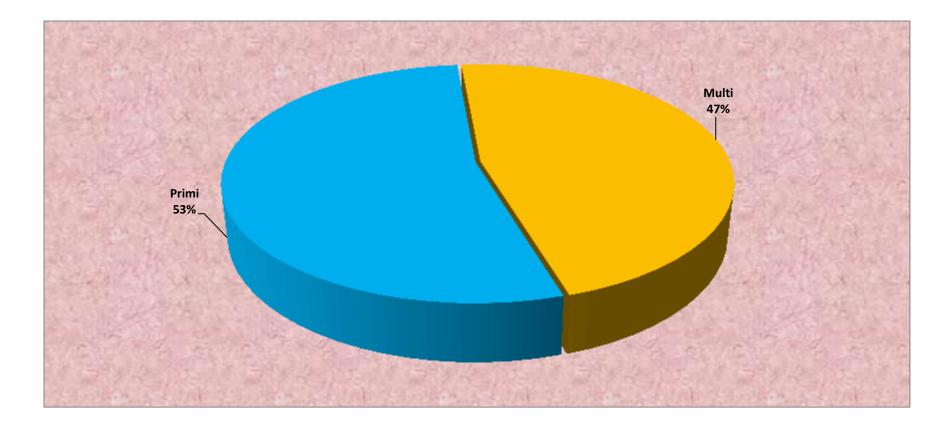


Figure 4.13: Gravida wise distribution of antenatal mothers.

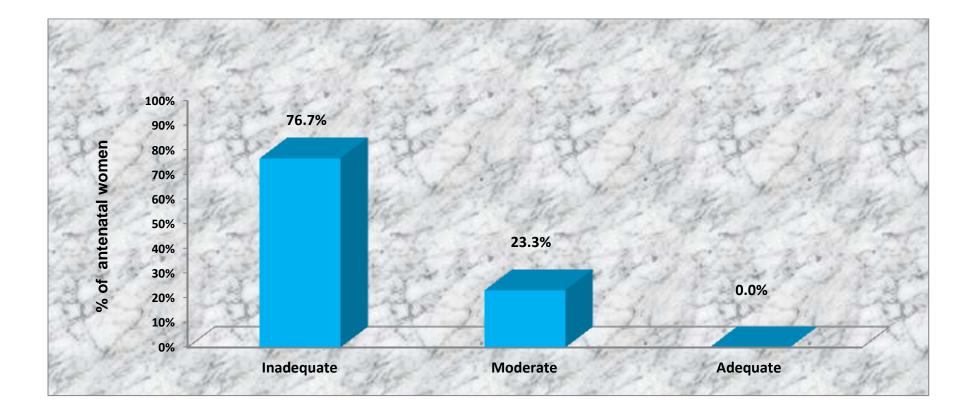


Figure 4.14: Pretest level of knowledge score of antenatal mothers.

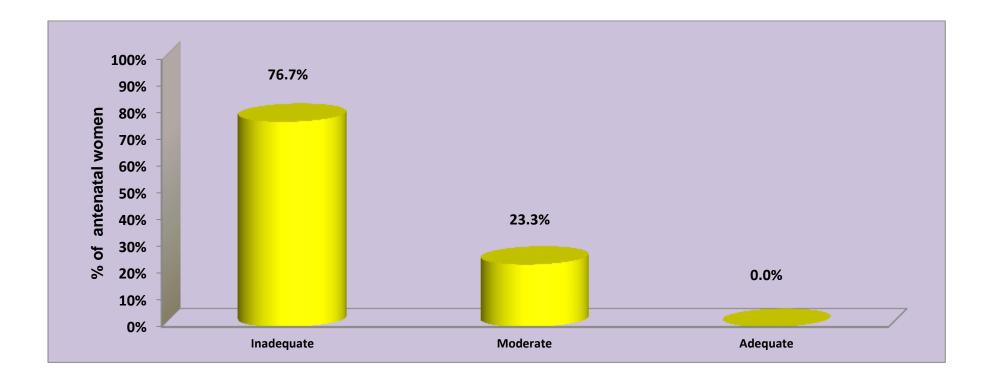


Figure 4.15: Posttest level of knowledge score of antenatal mothers.

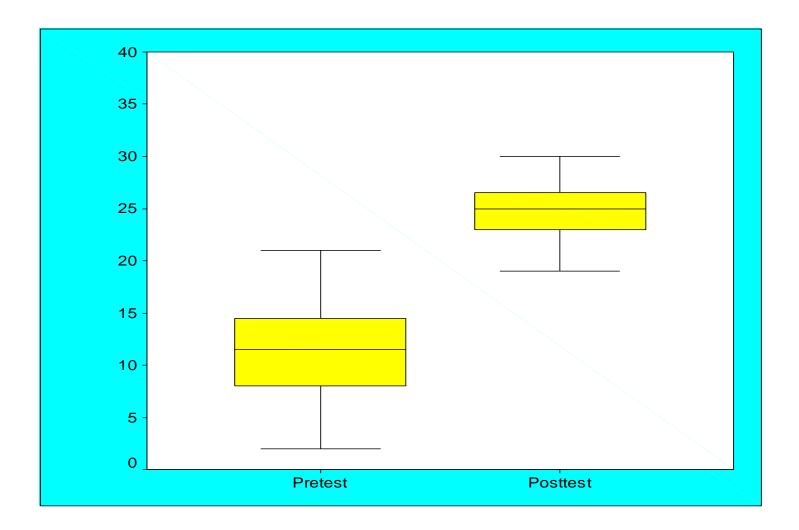


Figure 4.16: Mean pretest and posttest knowledge score of antenatal mothers.

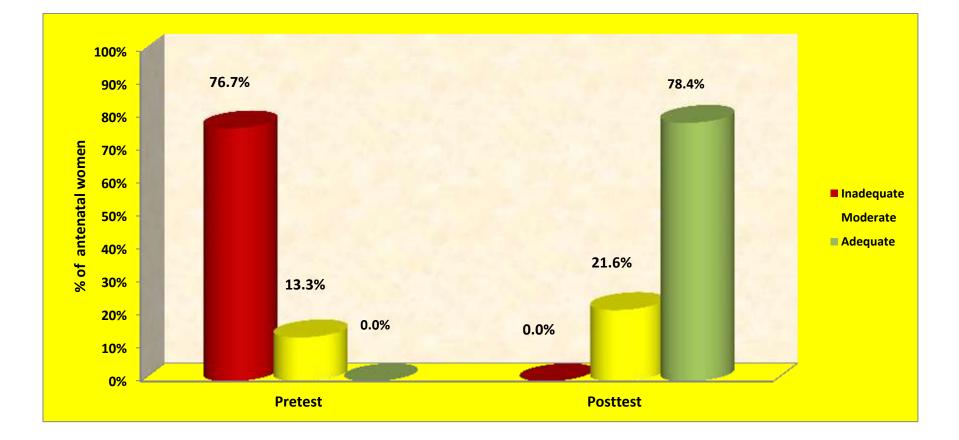


Figure 4.17: Pretest and posttest level of knowledge score.

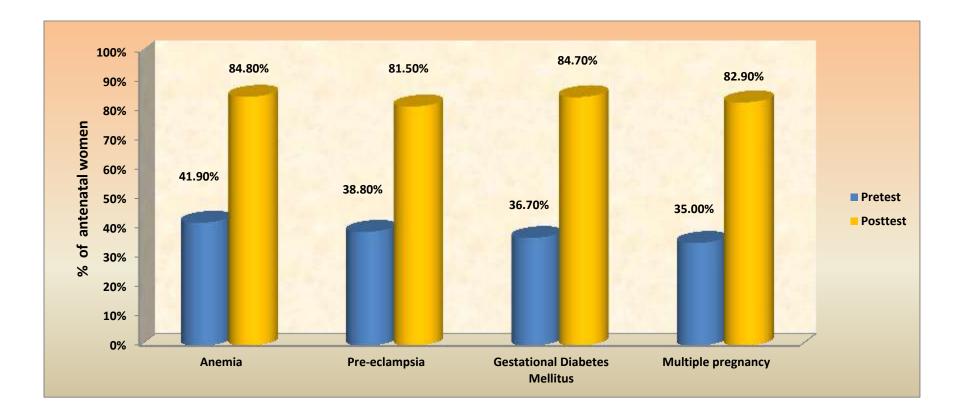


Figure 4.18: Pre and posttest percentage of knowledge score of antenatal mothers.

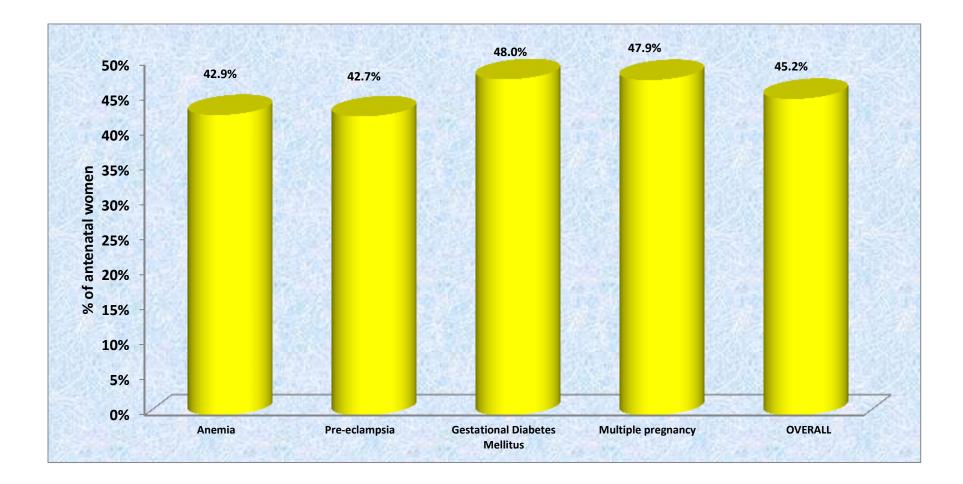


Figure 4.19: Percentage of knowledge gain.

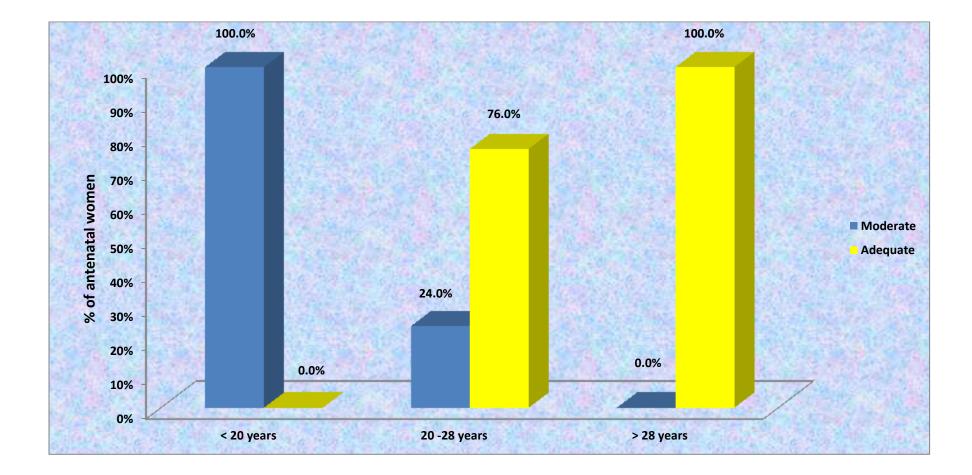


Figure 4.20: Association between posttest level of knowledge and antenatal women age.

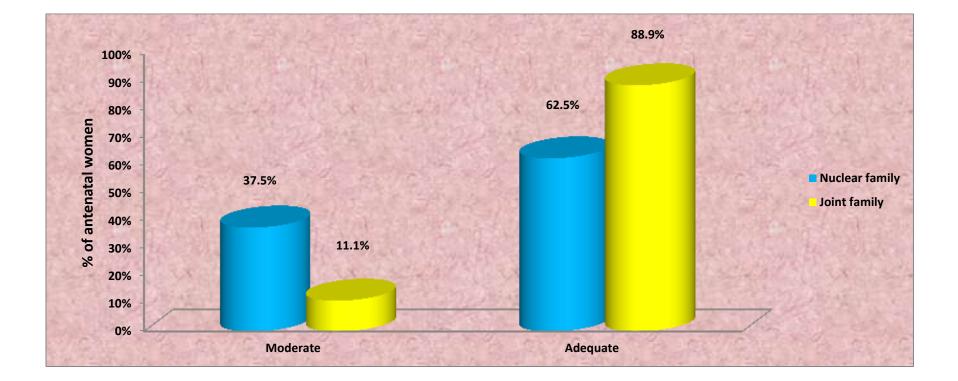


Figure 4.21: Association between posttest level of knowledge and antenatal women type of family system.

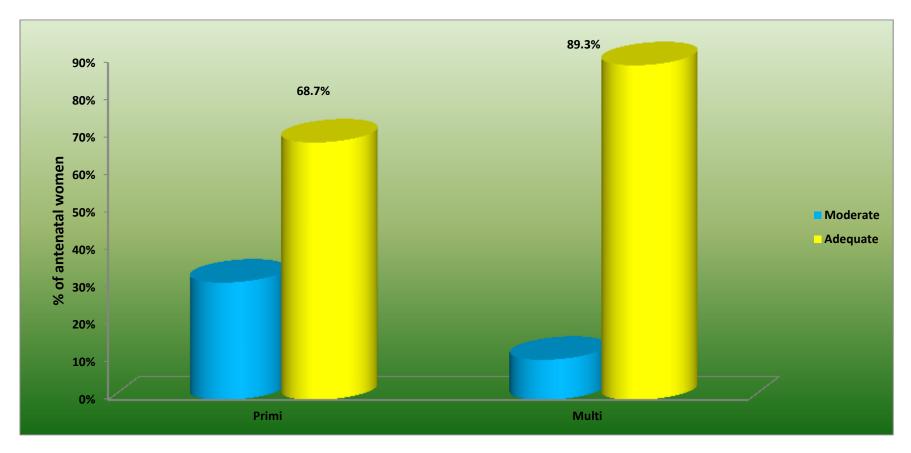


Figure 4.22: Association between posttest level of knowledge score and antenatal women gravida.

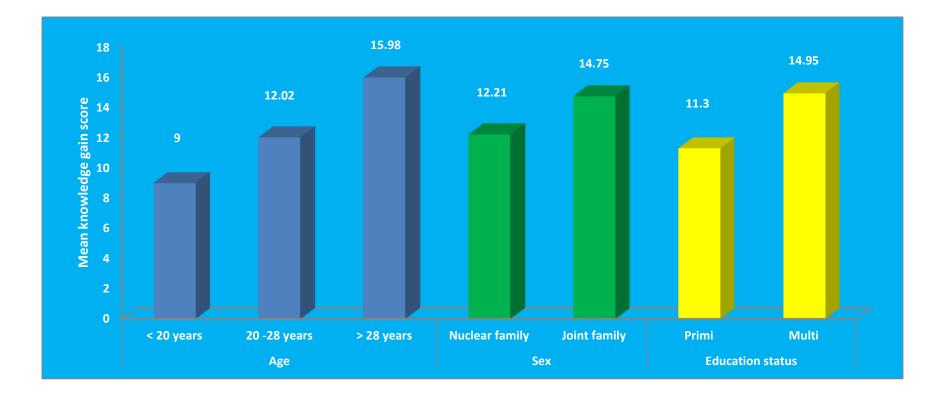
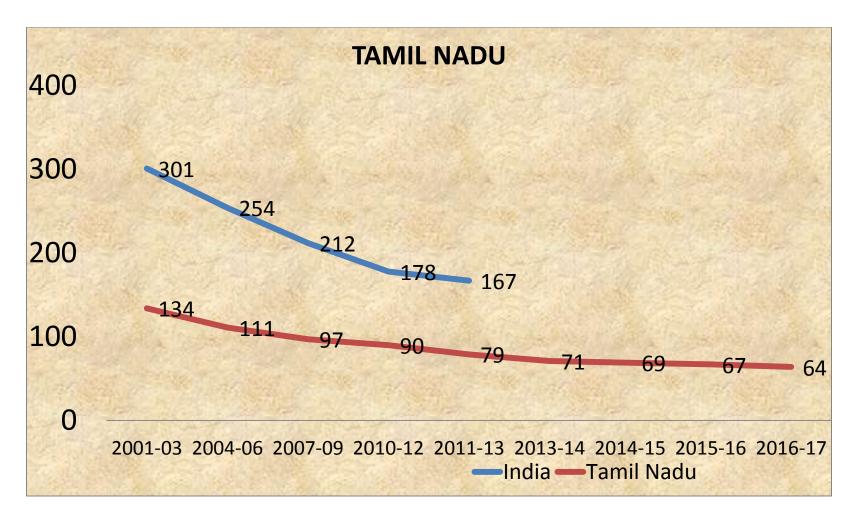
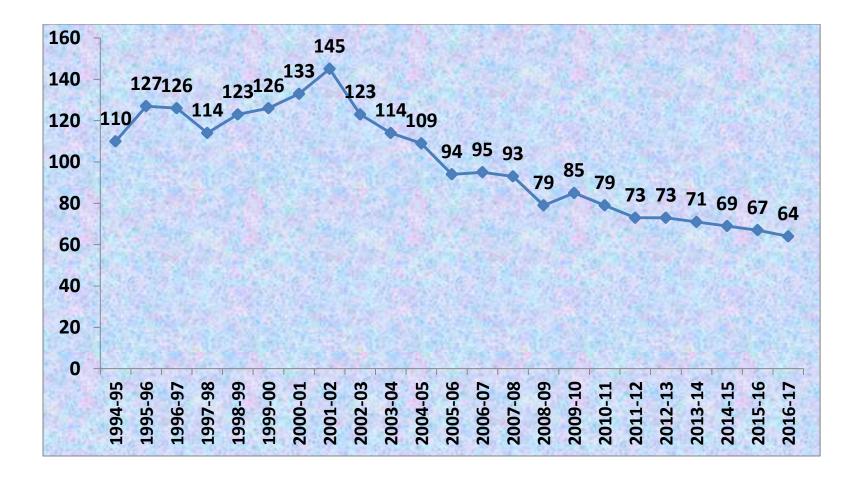


Figure 4.23: Association between knowledge gain score and demographical variables.



Source: SRS (2001-03 to 2011-13) State HMIS (2013-14 to 2016-17) Figure 1.1: Maternal Mortality Ratio



Source: State HMIS Figure1.2: Time trend in Maternal Mortality Ratio-Tamilnadu

DESCRIPTION OF THE TOOL SECTION – A TOOL -1: DEMOGRAPHIC VARIABLES

By interview method the researcher will mark each response of the antenatal women

1	Age (I	n Years)	
	a)	Below 20	
	b)	20-28	
	c)	Above 28	
2	Religio		
	a)	Hindu	
	b)	Muslim	
	c)	Christian	
3	Туре с	of Family	
	a)	Nuclear Family	
	b)	Joint Family	
4	Educ	ational Level	
	a)	Elementary education	
	b)	Middle education	
	c)	Higher secondary education	
	d)	Graduate	
5	Occu	pation	
	a)	Home maker	
	b)	Government job	
	c)	Private job	
	d)	Self job	
6	Family mor	nthly income (in rupees)	
	a)	≤ 10000	
	b)	10001 to 20001	
	c)	>20001	

SECTION-B

$\underline{TOOL-2}$

By the interview method the researcher will mark each response of the antenatal women.

OBSTETRICAL VARIABLES

7.	Age at marriage (in Years)	
8.	Last menstrual period	
9.	Expected date of delivery	
10.	Obstetrical score GPLA	
11.	Age at first child birth (in years)	
12.	Pre pregnant health status	

<u>SECTION – C</u>

<u>TOOL – 3: STRUCTURED QUESTIONARRE REGARDING</u> <u>HIGH RISK PREGNANCY.IT CONSISTS OF PART A,B,C,</u> <u>AND D</u>

By interview method the researcher will mark each response of the antenatal women.

PART A: QUESTION RELATED TO ANEMIA

1.	What is meant by anemia?	
	a) Increased level of hemoglobin in the blood	
	b) Normal level of hemoglobin in the blood	
	c) Decreased level of hemoglobin in the blood	
2.	What is the normal hemoglobin level in a pregnant	
	woman?	
	a) 10-11 gm/dl	
	b) 11-16 gm/dl	
	c) More than 16 gm/dl	
3.	What is the cause of anemia in a pregnant woman?	
	a) Anemia in previous pregnancy	
	b) Family history of anemia	
	c) Increased need of iron during pregnancy.	
4.	Which of the following features are the predictors to	
	identity anemia?	
	a) Pallor, fatigue and difficult breathing	
	b) Headache, vomiting, diarrhea	
	c) Bleeding, abdominal pain and leg cramps	
5.	What ate the food items to be added to prevent	
	anemia? a)Apple, Guava	
	b)Greens, vegetables and dates	
	c) Chicken, Cool drinks	
	-,,,	

6.	cons a) R b) It	y a pregnant women should avoid sumption of tea after meals? educe the absorption of iron. can cause nausea and vomiting. can cause discoloration of teeth.	
7.	Whic	h is the most preferable time to prevent anemia?	
	a)	Preconception period	
	b)	After conception	
	c)	When the antenatal women are diagnosed to have anemia	
8.	What	is the effect to untreated anemia?	
	a)	Increased blood sugar	
	b)	Preterm labour and low birth weight baby	
	c)	Increased blood pressure	
PAR	T B: (QUESTION RELATED TO PRE-ECLAMPSIA	
1.	What	is meant by pre- eclampsia?	
	a)	Presence of normal blood pressure with swelling	
	b)	Presence of low blood pressure with swelling	
	c)	Presence of high blood pressure, and swelling	
2.	What	is normal blood pressure of an antenatal woman?	
		80/50 mm of Hg	
		120/80 mm of Hg	
	c)	140/100 mm of Hg	
3.	Com	nonly at which gestational age the pre- eclampsia will	
	occur		
	a)	At 12 th week	
	,	Between 13-20 week	
	c)	After 20 th week	
4.	What	is relation between bed rest and swelling in pre -	
	eclam	ipsia?	
	a)	Swelling increases with bed rest	
	b)	Swelling decreases with bed rest	
	c)	Swelling persists with bed rest	

5.	Which position a pregnant woman should assume during sleep?				
	a) Right lateral				
	b) Left lateral				
	c) Supine position				
6.	How will you assess the fetal well being?				
	a) By checking blood pressure				
	b) By checking weight				
	c) By monitoring kick count				
7.	What are the warning signs of pre-eclampsia?				
	a) Heart burn, blurred vision, Hands and legs edema and reduced urine output				
	b) Fever				
	c) Back pain				
8.	What is the effect of untreated pre- eclampsia?				
	a) Cancer				
	b) Fits				
	c) Thyroid problems.				

PART C: QUESTIONS RELATED TO GESTATIONAL DIABETES MELLITUS

1.	What is normal blood glucose level of an antenatal woman?	
	a) 135-155 mg/dl	
	b) 80-120 mg/dl	
	c) 50-70 mg/dl	
2.	Which of the following is a risk factor for diabetes in pregnancy?	
2.		
2.	pregnancy?	

at is the cause of diabetes in pregnancy?	
a) Due to Insufficient insulin production	
b) Increased intake of sweet by women	
c) Lack of exercise	
at is symptom of diabetes in pregnancy?	
a) Bleeding per vagina	
b) Frequent urination and thirst	
c) Sudden increase of blood pressure	
ich is the test to be used in diagnosing the diabetes	
ich of the following are treatment measures in a	
_	
-	
ich of the following is the complication of diabetes?	
a) A small sized baby	
b) A baby with high blood pressure	
QUESTION RELATED TO MULTIPLE PREGN	ANCY
	 at is the cause of diabetes in pregnancy? a) Due to Insufficient insulin production b) Increased intake of sweet by women c) Lack of exercise at is symptom of diabetes in pregnancy? a) Bleeding per vagina b) Frequent urination and thirst c) Sudden increase of blood pressure ich is the test to be used in diagnosing the diabetes litus during pregnancy? a) Hemoglobin estimation b) Blood group analysis c) Oral glucose challenge test ich of the following are treatment measures in a gnant woman with diabetes? a) Special diet, exercise and insulin if needed b) Insulin injection only c) Adequate food only ich of the following is the complication of diabetes? a) A small sized baby b) A baby with high blood pressure

1.	What is meant by multiple pregnancy?	
	a) Development of two or more fetus in the uterus	
	b) Development of fetus outside the uterus	
	c) Repeated pregnancies	
2.	What may be the cause of multiple pregnancy?	
	a) Teenage pregnancy	
	b) Heredity	
	c) Obesity	

3.	What	are the signs and symptoms of multiple pregnancy?	
	a)	Fetal movement felt in different parts of abdomen at the	
		same time	
	b)	Increased blood pressure and increased blood sugar	
	c)	Ectopic pregnancy.	
4.	Whic	h method is used to diagnose multiple pregnancy?	
	a)	Blood sugar estimation	
	b)	Through Ultrasonography	
	c)	Urine examination	
5.	What	will be the ideal food recommended for the babies	
	delive	ered by a pregnant woman with multiple pregnancy?	
	a)	Breast milk for all the babies	
	b)	Lactogen	
	c)	One baby with breast milk and others with cow's milk	
6.	What	is the dietary requirement for a woman with multiple	
	pregn	ancy?	
	a)	High calorie, High protein, High iron diet	
	b)	Low fat, Low protein	
	c)	High fat, High Protein	
7.	What	is the complication to fetus due to multiple	
	pregn	ancy?	
	a)	Over weight baby	
	b)	Infectious baby	
	c)	Preterm delivery.	

LESSON PLAN ON

HIGH RISK PREGNANCY

- TOPIC:HIGH RISK PREGNANCY
- DURATION : 30 MINUTES
- **GROUP** : ANTENATAL MOTHER BETWEEN 8-28 WEEKS
- METHOD OF TEACHING : LECTURE CUM DISCUSSION
- AV AIDS : BOOKLET
- VENUE : ANTENATAL OUTPATIENT DEPARTMENT, INSTITUTE OF OBSTETRICS AND GYNAECOLOGY, CHENNAI-08.
- INVESTIGATOR'S NAME : J.DHANALAKSHMI MSC NURSING II YEAR

CENTRAL OBJECTIVE:

At the end of the teaching program the group will acquire knowledge regarding high risk pregnancy and develop desirable attitude towards early identification of high risk pregnancies and prevention of the complications and apply it in practice.

SPECIFIC OBJECTIVES:

The antenatal mothers will be able to,

- state the meaning of each risk pregnancy
- list the causes of each high risk pregnancy
- identify the signs and symptoms of each high risk pregnancy
- describe the diagnosis of each high risk pregnancy
- understand the prevention of each high risk pregnancy
- > explain the management of each high risk pregnancy
- > enumerate the complications of each high risk pregnancy

TIME	SPECIFIC	CONTENT	TEACHER'S		AV AIDS	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY		
1		INTRODUCTION				
min						
	state the meaning of high risk pregnancy	All pregnancies and deliveries are potentially at risk. However, there are certain categories of pregnancies where the mother, the fetus of the neonate is in a state of increased jeopardy. About 20 to 30% pregnancies belong to this category. If we desire to improve obstetric results, this group must be identified and given extra care.	Lecture cum discuss	Discussing	Booklet	What is mean by high risk pregnancy?

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 Min		MEANING OF HIGH RISK PREGNANCY Pregnancy is complicated by variety of disorders and conditions that can profoundly affect the women and her fetus. The pathophysiology of their disorders may adversely affect the pregnancies. These pregnancies are often regarded as high risk. Some of the high risk pregnancies > Anemia > Pre- eclampsia > Gestational diabetes mellitus > Multiple pregnancy				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 Min	state the meaning of anemia	<u>1. ANEMIA</u> <u>MEANING</u> A nemia is a common health problem throughout the world particularly among pregnant women. It is a condition in which there is a low level of hemoglobin in the blood, to a level below 11g/dl. Hemoglobin is red pigment present in the blood cells, and its primary function is to carry oxygen to all parts o the body. The normal hemoglobin level in women is 11-16 g/dl. Approximately 80% of all anemia's in pregnancy result from iron deficiency.	Discussion	Discussing	Booklet	What is normal hemoglobin level of pregnant women?
1 Min	list the causes of anemia	 CAUSES Increased demand for iron Diminished intake Physical stress Pre pregnant health status Blood loss Repeated pregnancies at short interval 	Lecturing	Listening	Booklet	What are all the causes of anemia?

TIME	SPECIFIC	CONTENT	TEACHER'S	LEARNER'S	AV AIDS	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY		
2	identify the signs	SIGNS AND SYMPTOMS	Explaining	Listening	Booklet	How will you assess
Min	and symptoms of					the anaemia?
	anemia	 Paleness which can be seen inside 				
		the lower eyelids, nails, tongue,				
		skin and palms of the hand				
		✤ Fatigue				
		 Labored breathing 				
		 Rapid heart rate 				
		 Giddiness 				
		 Lethargy 				
		 Lack of concentration 				
		 Weigh loss 				
		 Loss of appetite 				
		 Spoon shaped nails in serve 				
		anemia				
		WARNING SIGNS:				
		Severe palmar, conjunctival, and				
		nail pallor.				
		Breathing 30 beats/minute.				
		 Easy fatigability 				
		 Breathlessness even at rest. 				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 Min	describe the diagnosis of anemia	 DIAGNOSIS ➢ Assessment of clinical features ➢ Hemoglobin estimation 	Lecture cum discussion	Discussing	Booklet	What are the methods used to test anaemia?
		Serum binding capacity				
2 Min	understand the prevention of anemia	 PREVENTION OF ANEMIA To prevent anemia it should start from pre conception period itself Early antenatal checkup. 				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
		 Avoidance of frequent child birth. Minimum interval between child birth is at least 2 years Supplementary iron therapy. Even with a well balanced diet supplementary iron to be given. Examination of stool for infestation of worm. In order to prevent worms infestation. 	Discussion	Discussing	Booklet	What are the food items contain more protein and iron?
		Total iron requirement during pregnancy is 1000mg. Intake of diet rich in protein and iron much as groundnuts, beans, dark green leafy vegetables, liver, milk, egg, jaggery and dates.				
		Regular consumption of iron and folio acid as prescribed by the doctor. Every pregnant woman should consume iron and folic acid tablets from the beginning of fourth month of pregnancy.				
		Don't consume tea after breakfast or a meal. Tea contains a substance and when consumed along with food it will prevent iron absorption.				

TIME	SPECIFIC	CONTENT	TEACHER'S	LEARNER'S	AV	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY	AIDS	
		Adequate Treatment to be given in case of				
		malaria, bleeding piles, urinary tract infection				
		 Early detection of falling of hemoglobin level 				
1	Explain the	MANAGEMENT OF ANEMIA				What are the side
Min	management of					effects of iron
	anemia	Diet: A realistic balanced diet, which is rich				therapy?
		in proteins, iron and vitamins.				
		 To improve appetite and facilitate digestion, 				
		preparation containing acid pepsin may be given				
		To eradicate even a minimal septic focus by				
		appropriate antibiotic therapy. Effective therapy				
		to cure the disease contributing to cause anemia				
		Iron therapy: Oral therapy, Parental therapy				
		 Side effects of these therapies are irritation of 				
		stomach, vomiting and black stools. The side				
		effects will be there only for few days and will				
		disappear.				
		The tablets are better tolerated when taken				
		after meals				
		 Blood Transfusion 				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 Min	Enumerate the complication of anemia in mother and fetus	COMPLICATION OF ANEMIA MOTHER	Lecturing	Listing	Booklet	
		 Infection 				
		 Preeclampsia 				
		 Uterine inertia 				
		 Heart failure 				
		Shock				
		 Puerperal sepsis 				
		 Sub involution 				
		 Failing lactation 				
		 Pulmonary embolism 				
		FETUS 🎸 Poor fetal growth				
		 Preterm birth 				
		Low birth weight				

TIME	SPECIFIC	CONTENT	TEACHER'S	LEARNER'S	AV AIDS	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY		
2 Min 2 Min	state the meaning of pre -eclampsia list the risk factors of pre -eclampsia	 PRE-ECLAMPSIA: Blood pressure to the extent of 140/90 mm of Hg or more plus protein in the urine with or without edema occurring after 20th week of pregnancy. <u>RISK FACTORS</u> ✓ Women who have pre -eclampsia in previous pregnancies ✓ Placental factors ✓ Increased placental mass ✓ Abnormal placentation ✓ Pregnancy below 20 years of age ✓ Pregnancy after 35 years of age ✓ Obesity ✓ Family history of high blood pressure ✓ Stress ✓ Past history of diabetes mellitus 	Discussion	Discussing	Booklet	What is pre- eclampsia?
2 Min	identify the signs and symptoms of serve pre -eclampsia	 SIGNS AND SYMPTOMS OF MILD PRE ECLAMPSIA ➢ High blood pressure: It is equal to or more than 140/90 mmHg 				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
		 Swelling Slight swelling on ankles which pits on pressure is found to persist even after 12 hours of bed rest or tightness of the ring on the finger. While pressing the edema the presence of 2mm depth-mild edema. 4 mm-moderate, 6mm severe, 10 mm very severe . Weight gain: The average total weight gain during pregnancy is 10-12 kg.In mild pre eclampsia weight gain is greater than 2 kg per month. SIGNS AND SYMPTOMS OF SEVERE PRE ECLAMPSIA 	Discussion	Discussing	Booklet	How much of weight should be increased during pregnancy?
		 Continuous high blood pressure Blood pressure is equal to or more than 160/110 mm of hg The swelling extends to the face, hands, abdominal wall, vulva, sacral area, ankles, feet and even the whole body Rapid weight gain 				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV SIDS	EVALUATION
1 Min	describe the diagnosis of pre - eclampsia	 Presence of protein per 24 hour urine (30 mg/dl) Head ache, either located over frontal or occipital region Dizziness Disturbed sleep Oliguria Urine output is less than 400 ml in 24 hours Epigastric pain associated with vomiting Blurring vision Decreased fetal movements WARNING SIGNS: Severe head ache Vision defect Reduced urine output Abdominal pain Face and leg edema. DIAGNOSIS Blood pressure monitoring Urine output in 24 hours Weight of the patient Edema testing Urine testing for protein Antenatal fetal monitoring Daily fetal kick count Ultrasonography 	Lecture cum Discussion	Listening	Booklet	What are the methods used to diagnose pre- eclampsia?

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
2 Min	understand the prevention of pre eclampsia	 PREVENTION OF PRE ECLAMPSIA Pre -eclampsia if not totally preventable disease. The following regime should be enforced in risk groups in an attempt to prevent or detect early manifestations of pre -eclampsia. Regular antenatal visits Once a month up to 6 month, once in 15days during seventh and eighth month and thereafter once a week till the expected date of delivery A women at risk should have at least ten antenatal check ups Adequate rest Sleep eight to twelve hours daily with 2 hours of rest in the middle of the day and eighty to ten hours during night from the 20th week of pregnancy onwards. Diet: Well balanced diet rich in protein Screening tests: Doppler assessment 	Lecture cum Discussion	Listening	Booklet	How long the antenatal women have to sleep?

TIME		CONTENT	TEACHER'S	LEARNER'S	AV	EVALUATION
	OBJECTIVES	Roll over test: This is done during 28-32 weeks of pregnancy, blood pressure is measured with women lying on her left side then she is asked to roll on her back to check blood pressure once again. Increases of 20 mm of hg in diastolic pressure from side to back position indicate a positive roll over test, which is the evidence of pre-eclampsia.	ACTIVITY Lecture cum Discussion	ACTIVITY Listening	AIDS Booklet	How many antenatal visits are required for a women with pre eclampsia?
2 min	explain the management of pre - eclampsia	 MANAGEMENT OF PRE ECLAMPSIA Adequate rest. Assume left lateral position during sleep Diet: Well balanced diet rich in protein (about 100 gm). Salt is neither restricted not forced. Total calorie approximately 1600 kcal/day. Monitor blood pressure regularly Monitor weight regularly Undergo all lab tests that are advised during pregnancy 				

TIME	SPECIFIC	CONTENT	TEACHER'S	LEARNER'S	AV	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY	AIDS	
		 ✓ Shock ✓ Dimness of vision even blindness ✓ Hemolytic anemia, elevated liver enzyme, low platelet count ✓ Eclampsia FETAL ✓ intra uterine growth retardation ✓ Pre mature birth ✓ Intrauterine death ✓ Asphyxia 				
1 Min	state the meaning of gestational diabetes mellitus	III. GESTATIONAL DIABETES MELLITUS MEANING Gestational diabetes mellitus is a condition in which the blood glucose level is increased during pregnancy. The normal blood glucose level is 80- 120 mg/dl. It occurs due to insufficient production of insulin by the body.	Lecture cum Discussion	Listening	Booklet	What is mean by gestational diabetes?
1 min	list the risk factors of gestational diabetes mellitus	RISK FACTORS✓Positive family history of diabetes✓History of still birth✓Unexpected perinatal loss				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S	LEARNER'S	AV AIDS	EVALUATION
			ACTIVITY	ACTIVITY		
		✓ Obesity	Discussion	Discussing	Booklet	
		✓ Age over 30 years				
		 Hypertension 				
		 Polyhydramnios . 				
		 Inadequate metabolic control 				
		✓ Five or more previous pregnancies				
1	identify the signs and	SIGNS AND SYMPTOMS				
min	symptoms of					
	gestational diabetes	 Increased thirst 	Lecturing	Listening	Booklet	What are the
	mellitus	 Increased urination 				signs and
		 Weight loss in spite of increased appetite 				symptoms of gestational
		 Fatigue, nausea, vomiting 				diabetes
		 Infections including bladder vagina 				mellitus
		and skin				menieus
		A fetus larger than the normal for				
		the state of pregnancy on prenatal				
		examination.				
		WARNING SIGNS:				
1	describe the diagnosis	✓ Vision defect				
min	of gestational diabetes	✓ Increased tiredness				
	mellitus	 Incresed urine output and thirst 	Lecture	Discussing	Booklet	what are the
		DIAGNOSIS	cum			test used to
			discussion			diagnosis
		 Fasting and post prandial blood 				gestational
		sugar estimation.				diabetes
		Glucose Challenge Test				mellitus?

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1min	understand the prevention of gestational diabetes mellitus	 Confirmation is to be done by testing a second fasting morning specimen of urine acetone . Glucose tolerance test. Ultrasonography PREVENTION: Weight management Diet Exercise 	Lecture cum Discussion	Lecturing	Booklet	
Min	inclinus	MANAGEMENT:				What is the diet to be
	explain the management of Gestational diabetes mellitus	 Antenatal supervision. Peroidic checkup of blood glucose level. Every women should check her blood glucose level. Maintaining the Fasting blood glucose level less than 95 mgs/dl. Special diet: The diet change will involve increased fiber intake ,fat restriction and elimination of concentrated sweets. Regular exercises, 				followed in gestational diabetes mellitus?

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
		Insulin therapy (if needed) only when the blood sugar is not controlled through restricted diet and regular exercises.				
1 min	enumerate the complications of gestational diabetes mellitus	COMPLICATIONS MATERNAL: ✓ Abortion ✓ Preterm labour ✓ Infection ✓ Pre- eclampsia ✓ Maternal distress ✓ Excess amniotic fluid ✓ Prolongation of labour ✓ Failing lactation ✓ Perineal injuries ✓ Risk for mother developing diabetes in future FETAL ✓ Congenital abnormalities like heart diseases ✓ Birth Injuries ✓ Hypoglycemia	Lecture cum Discussion	Discussing	Booklet	What are the complications to mother due to gestational diabetes?
		✓ Respiratory distress				

TIME	SPECIFIC OBJECTIVES	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 min	state the meaning of multiple pregnancy	IV. MULTIPLE PREGNANCY MEANING It means development of two or more fetus in pregnant uterus.	Lecture cum discussion	Discussing	Booklet	What is meant by multiple pregnancy?
1 min	list the risk factors of multiple pregnancy	RISK FACTORS> Heredity> Older age> High parity> Race> Assisted reproductive technology	Discussion	Discussing	Booklet	What are the risk factors of multiple pregnancy?
1 min	identify the signs and symptoms of multiple pregnancy	 SIGNS AND SYMPTOMS Uterus is larger than expected date of pregnancy Increased morning sickness Increased appetite Excessive weight gain Fetal movement felt in different parts of abdomen at the same time 	Explaining	Listening	Booklet	What are the signs and symptoms of multiple pregnancy?

TIME	SPECIFIC OBJECTIVE	CONTENT	TEACHER'S ACTIVITY	LEARNER'S ACTIVITY	AV AIDS	EVALUATION
1 min	describe the diagnosis of multiple pregnancy	 DIAGNOSIS Ultrasonography Physical examination,by inspection and palpation Blood testing Alfa feto protein level 	Lecturing	Listening	Booklet	How will you diagnosis the multiple pregnancy?
2 Min	explain the management of multiple pregnancy	 MANAGEMENT Increased nutrition: Mothers carrying two or more futures need more calories, protein, and other nutrients, including iron. Higher weight gain is also recommended. More frequent parental visits Increased rest Fetal growth assessment Maternal and fetal testing Tocolytics (if preterm labour) Cervical encirclage (for incompetent uterus) Supplementary therapy: Iron therapy to be increased to the extent of 60-80 mg/dl. 	Lecture cum discussion	Discussing	Booklet	What are the management for multiple pregnancy?

TIME	SPECIFIC	CONTENT	TEACHER'S	LEARNER'S	AV	EVALUATION
	OBJECTIVES		ACTIVITY	ACTIVITY	AIDS	
1	enumerate the	<u>COMPLICATIONS</u>	Lecture cum	Discussing	Booklet	What are
min	complications of		discussion			complication of
	multiple pregnancy	MATERNAL				multiple
		 Nausea and vomiting occurs with 				pregnancy?
		increased frequency and severity				
		🗸 Pre -eclampia, Anemia				
		 Abnormal amount of amniotic fluid 				
		✓ Increased operative interference				
		✓ Bleeding				
		✓ Early rupture of membrane.				
		✓ Prolonged labour				
		<u>FETAL</u>				
		✓ Birth defect				
		✓ Premature birth				
		✓ Growth problem				
		 Intra uterine death or still birth 				
		✓ Fetal anomalies				
		✓ Asphyxia				
1		CONCLUSION				
min						
		Despite not every mother pregnant will				
		have complications but knowing				
		complications on risks during pregnant can				
		assist, handles prevent complication occur.				

SCORING KEY

QUESTION NUMBER	ANSWER	SCORE
I.ANEMIA		
1	c	1
2	b	1
3	c	1
4	a	1
5	b	1
6	a	1
7	a	1
8	b	1
II.PRE-ECLAMPSIA		
1	с	1
2	b	1
3	С	1
4	С	1
5	b	1
6	с	1
7	a	1
8	b	1

III.GESTATIONAL	ANSWER	SCORE
DIABETES MELLITUS		
1	b	1
2	с	1
3	а	1
4	b	1
5	с	1
6	а	1
7	с	1
IV.MULTIPLE		
PREGNANCY		
1	a	1
2	b	1
3	а	1
4	b	1
5	a	1
6	a	1
7	с	1

<u>விபரங்களை தொகுப்பதற்குரிய கருவி</u>

பகுதி-அ கருவி-1: தனிநபர் அழப்படை தகவல்கள்

மாதிரி எண்:

1)	ഖലള	து (ஆண்டுகளில்)	
	එ	20க்கும் குறைவு	
	ஆ)	20-28 ഖயது	
	B	28க்கும் அதிகம்	
2)	மதம்		
	එ	இந்து	
	ஆ)	முஸ்லிம்	
	Ð	கிறிஸ்தவா	
3)	குடுப	ப்ப வகைபாடு	
	அ	தனிக்குடும்பம்	
	ஆ)	கூட்டுக்குடும்பம்	
4)	கல்வ	ித்தகுதி	
	එ	ஆரம்பக் கல்வி	
	ஆ)	இடைநிலைக் கல்வி	
	B	உயா்நிலைக்கல்வி	
	(म	பட்டதாரி	
5)	தொ	ழில்	
	එ	குடும்பத் தலைவி	
	ஆ)	அரசுப்பணி	
	Ð	தனியாா் பணி	
	म)	சொந்த வேலை	
6)	கடுப	ப் மாத வருமானம்	
	එ)	ரூ.10000க்கும் குறைவு	
	ஆ)	ரூ.10001 முதல் 20000/–	
	B	ரூ.20001க்கு மேல்	

பகுதி-ஆ கருவி-2: பேறுத்தன்மை பற்றிய விபரங்கள்

7)	திருமணத்தின்போது வயது (ஆண்டுகளில்)	
8)	கடைசி மாதவிடாய் காலம்	
9)	குழந்தை பிறப்பினை எதிர்பார்க்கும் நாள்	
10)	பேறுத்தன்மை பற்றிய குறியீடு ஜி.பி.எல்.ஏ	
11)	முதல் குழந்தை பிறப்பின்போது வயது	
12)	முன் காப்பகால உடல்நிலை	

பகுத்-இ

கருவி-3: தீட்டமிட்ட வினாத்தாள் அதிக பாதிப்பிற்குரிய கர்ப்பங்கள் பற்றிய வினாக்கள் கொண்டது. இது பிரிவு அ, ஆ, இ மற்றும் ஈ கொண்டது

<u>பகுத்-அ: இரத்தசோகை குறித்த வினாக்கள்</u>

1)	இரத்	த சோகை என்றால் என்ன?	
	அ)	இரத்தத்தில் ஹீமோகுளோபின் அளவு அதிகமாதல்	
	ஆ)	இரத்தத்தில் ஹீமோகுளோபின் அளவு சராசரியாயிருத்தல்	
)	இரத்தத்தில் ஹீமோகுளோபின் அளவு குறைதல்	
2)	പെൽ	ரகளில் சரியான ஹீமோகுளோபின் அளவு என்ன?	
	එ)	1O–11 கி/டெசி.லி	
	ஆ)	11–16 ക്/ டെசி.லி	
)	16 கீ/ டெசி.லி க்கும் அதிகம்	
3)	கர்ப்ப	ிணி பெண்ணில் இரத்த சோகை ஏற்படக் காரணம் என்ன?	
	அ)	முந்தைய காப்பகாலத்தில் இரத்த சோகை	
	ஆ)	இரத்த சோகை கொண்ட வரலாறு	
)	கா்ப்பகாலத்தில் அதிகமான இரும்புச்சத்து தேவை	
4)		வருவனவற்றில் எந்த அறிகுறி இரத்த சோகை இரு பிடுகின்றது?	ப்பதை
	එ)	வெளிரி காணப்படுதல், களைப்பு, மூச்சுவிடுதலில் சிரமம்	
	ஆ)	தலைவலி, வாந்தி பேதி	
)	இரத்தக் கசிவு, வயிற்று வலி, கால் தசை இறுக்கம்	
5)	• •	த சோகையை தடுக்க எந்த உணவுப்பொருட்களை	
	சோ்த்	துக்கொள்ள வேண்டும்?	
	එ)	ஆப்பிள், கொய்யாப்பழம்	
	ஆ)	பச்சைக் கீரை, காய்கறிகள், போீச்சம்பழம்	
	S	கோழிக்கறி, குளிா்பானங்கள்	

6)		கா்ப்பினி பெண் உணவு உண்டபின் தேநீா் அருந்துவதை த எடும்?	விர்க்க
	அ	இரும்புச்சத்து உறிஞ்சும் தன்மையை குறைக்கும்	
	ஆ)	இது குமட்டல் மற்றும் வாந்தியை ஏற்படுத்தும்	
)	இது பற்களில் கரையை ஏற்படுத்தும்	
7)	இரத்	த சோகையை தடுக்க எது சரியான நேரம்?	
	அ	காப்பமடையும் முன்னா்	
	ஆ)	காப்பமடைந்த பின்னா்	
	B	கா்ப்பினி பெண்ணில் இரத்த சோகை இருப்பதை	
		கண்டறிந்தபின்	
8)	சிகிச்ச	சை அளிக்கப்படாத இரத்த சோகையின் விளைவு என்ன?	
	அ	அதிக இரத்த சாக்கரை	
	ஆ)	குறைமாத மற்றும் எடை குறைவான குழந்தை	
	B	அதிக இரத்த அழுத்தம்	

<u>பிரிவு-அ: கர்ப்பகாலத்திற்கு முன் ஏற்படும் வலிப்பு நோய்</u> <u>பற்றிய வினாக்கள்</u>

1) கா்ப்பகால வலிப்ப	நோய் என்றால் என்ன?	
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ക)	சராசரி இரத்த அழுத்தத்துடன் கூடிய வீக்கம் காணப்படுதல் 🗌	
<u> </u>		

- ஆ) குறைவான இரத்த அழுத்தத்துடன் கூடிய வீக்கம் காணப்படுதல்
- இ உயர் இரத்த அழுத்தத்துடன் கூடிய வீக்கம் காணப்படுதல்
- 2) ஒரு காப்பிணிப் பெண்ணின் சராசரியான இரத்த அழுத்தத்தின் அளவு என்ன?
 - அ) 80–50 மி.மீ. பாதரச அழுத்தம்
 - ஆ) 120–80 மி.மீ. பாதரச அழுத்தம்
 - இ 140–100 மி.மீ. பாதரச அழுத்தம்

3)	பொ	துவாக எந்த கா்ப்ப காலத்தில் கா்ப்ப கால வலிப்பு நோய் ஏற்ப(6ம்?
	ළ)	12 வாரத்தில்	
	ஆ)	13–20 வது வாரத்தில்	
	മ	20வது வாரக்கிற்கு மேல்	

5)

6)

7)

- ஆ) 13–20 வது வாரத்தில்
- 2 20வது வாரத்திற்கு மேல்
- காப்பகால வலிப்பு நோயில் படுக்கை ஓய்விற்கும் வீக்கத்திற்கும் உள்ள 4) சம்பந்தம் என்ன?

அ)	படுக்கை ஓய்வினால் வீக்கம் அதிகரிக்கிறது	\bigcirc
ஆ)	படுக்கை ஓய்வினால் வீக்கம் குறைகிறது	
B	படுக்கை ஓய்வினால் வீக்கம் அப்படியே இருக்கிறது	
எந்த	நிலையில் ஒரு கா்ப்பிணி பெண் உறங்க வேண்டும்?	
அ	வலதுபுறமாக சரிந்த நிலையில்	
ஆ)	இடதுபுறமாக சரிந்த நிலையில்	
B	மல்லாா்ந்த நிலையில்	
ദ ിക്ക്	ின் ஆரோக்கிய நிலையினை எவ்வாறு அறியலாம்?	
அ	இரத்த அழுத்தத்தை பரிசோதிப்பதன் மூலம்	
ஆ)	எடையை அளப்பதன் மூலம்	
B	சிசுவின் அசைவினை வைத்து	
கர்ப்ப	பகால வலிப்பு நோயின் எச்சரிக்கை அறிகுறிகள் என்ன?	
ച	ைக்க எரிச்சல் மங்கலான பார்வை கைகால்களில் வீச்சும்	

ආ)	நெஞ்சு எரிச்சல், மங்கலான பார்வை, கைகால்களில் வீக்கட	Ď
	குறைவான சிறுநீா் வெளியேற்றம்	
ஆ)	காய்ச்சல்	
Ð	ധ്രച്ചക്രഖலി	

- 8) சிக்ச்சை அளிக்கப்படாத கா்ப்பகால வலிப்பு நோயின் விளைவு என்ன?
 - அ புற்றுநோய்
 - ഖலിப்பு ஆ)
 - தைராய்டு 2

<u>பிரிவு-ஆ: கர்ப்பகால சர்க்கரை நோய் பற்றிய வினாக்கள்</u>

1	கர்ப்ப	ப காலத்தின் சரியான இரத்த சா்க்கரையின் அளவு என்ன?
	එ)	135–155 ഥി.ക്/ പെഴി.லി
	ஆ)	80–120 ഥി.ക്/ പെഴി.லി
	B	50–70 மி.கீ/ டெசி.லி
2)		வருவனவற்றில் எது கா்ப்பகாலத்தில் சா்க்கரை நோய்க்கான க்கும் காரணி?
	එ)	இரத்த சம்பந்தப்பட்ட உறவுகளில் திருமணம்
	ஆ)	இரத்தத்தில் குறைந்த அளவு ஹீமோகுளோபின்
	Ð	சா்க்கரை நோய் கொண்ட குடும்ப வரலாறு
3)	கர்ப்ப	ப காலத்தில் சா்க்கரை நோய் ஏற்படக் காரணம் என்ன?
	එ)	இன்சுலின் உற்பத்தியின் அளவு குறைவாயிருத்தல்
	ஆ)	பெண்கள் அதிக அளவு இனிப்பை உட்கொள்ளுவதால் 📃
	Ð	உடற்பயிற்சியின்மை
4)	கர்ப்ப	பகாலத்தில் சா்க்கரை நோய்க்கான அறிகுறி என்ன?
	එ	கா்ப்பப்பை வாய்க்குழியில் இரத்தம் கசிதல்
	ஆ)	அதிக தாகம் மற்றும் சிறுநீா் கழிப்பு
	D	திடீர் இரத்த அழுத்த உயர்வு
5)		ப காலத்தில் சா்க்கரை நோயை எந்த பாிசோதனையை பயன்படுத்தி டுபிடிக்கலாம்?
	එ)	ஹீமோகுளோபின் அளவை அளப்பதன் மூலம்
	ஆ)	இரத்த பிரிவினை பரிசோதிப்பதன் மூலம்
	Ð	வாய்க்குழி குளுகோஸ் தாங்குதிறன் ஆய்வின் மூலம் 📃
6)		வருவனவற்றில் எது கா்ப்பகாலத்தில் சா்க்கரை நோயினால் டக்கூடிய விளைவு?
	එ)	அளவில் சிறிய குழந்தை
	ஆ)	அதிக இரத்த சா்க்கரையுடன் கூடிய குழந்தை
	Ð	அளவில் பெரிய குழந்தை

7)		வருவனவற்றில் எது கா்ப்பகாலத்தில் சா்க்கரை நே னணிற்கு அளிக்க வேண்டிய சிகிச்சை முறைகள்?	ாயுள்ள
	அ	சிறந்த உணவு, உடற்பயிற்சி, இன்சுலின் ஊசி	
	ஆ)	இன்சுலின் ஊசி மட்டும்	
	Ŋ	போதுமான உணவு மட்டும்	
	<u>ப</u> ி	<u>ரிவு-®: பல கரு கர்ப்பத்தை பற்றிய வினாக்கள்</u>	
1	പல ര	கருவுற்ற கா்ப்பம் என்றால் என்ன?	
	ළ)	2 அல்லது அதற்கு மேற்பட்ட சிசு கருப்பையில் வளா்தல்	
	ஆ)	கருப்பையின் வெளியே சிசு வளா்தல்	
	<u>s</u>	அடுத்தடுத்த காப்பங்கள்	
2)	പരം പ	கருவுற்ற காப்பத்தின் காரணம் என்ன?	
	அ	இளம் வயதில் கா்ப்பமாதல்	
	ஆ)	பாரம்பரியம்	
	®)	உடல் பருமன்	
3)		கருவுற்ற கா்ப்பத்திற்கான அறிகுறி என்ன? சிசனின் அசையை வயிற்றின் வெல்லோட பசரிசனில் வி	-
	එ)	சிசுவின் அசைவை வயிற்றின் வெவ்வேறு பகுதிகளில் ஒழே கடியர் சில் உணகால்	
	ر س	சமயத்தில் உணருதல் உயர் செர்ச ஆயர்சால் உயர் செர்சாசர்சாசா	
	ஆ) ி	உயா் இரத்த அழுத்த மற்றும் உயா் இரத்த சா்க்கரை 	
	Ð	வெளிப்புற காப்பம்	
4)	പல ര	கருவுற்ற கா்ப்பத்தை எம்முறையை பயன்படுத்தி கண்டறியல	ாம்?
	එ)	இரத்த சா்க்கரையின் அளவினை பாிசோதிப்பதன் மூலம்	
	ஆ)	ஸ்கேன் பரிசோதனை மூலம்	
	®)	சிறுநீா் பாிசோதனையின் மூலம்	
5)		கருவுற்ற கா்ப்பத்தில் குழந்தை பிறப்பிற்கு பின்பு குழந்தை டுக்க வேண்டிய உணவு என்ன?	களுக்கு
	ආ	எல்லா குழந்தைகளுக்கும் தாய்ப்பால்	
	ஆ)	லேக்டோஜன்	
	B	ஒரு குழந்தைக்கு தாய்ப்பாலும், மற்ற குழந்தைகளுக்கு	_
		பசும்பாலும்	

6)	பல என்க	கருவுற்ற கா்ப்ப ன?	பத்தில் கா்ப்பிண பக்குக்குக்குக்குக்கு	<u>നി</u>	உணவு	ട്ടേഞഖ
	එ)	அதிக கலோரி, எ	அதிக புரதம், அ	திக இரும்புச் சத	ந்து	
	ஆ)	குறைந்த கொயு	ழப்பு, குறைந்த ப	ரதம்		
	8)	அதிக கொழுப்பு	l, அதிக புரதம்			
7)	പல ര	கருவுற்ற தன்மை	யினால் சிசுவிற்	க்கு உண்டாகும்	பாதிப்பு எ	ன்ன?
	එ)	அதிக எடையுள்	ள குழந்தை			
	ஆ)	நோய் தொற்று	ள்ள குழந்தை			
	R	ക്രണ്ഡാന് വിന്നം	வம்			

குறைமாத பிரசவம் **)**

அத்க பாதிப்பிற்குரிய கர்ப்பம் பற்றிய பாடத்திட்டம்

பாடத்தீட்டம்

தலைப்பு	:	அதிக பாதிப்பிற்குரிய காப்பம்
காலம்	:	30 நிமிடங்கள்
ወ ው	:	8–28 வாரங்களில் உள்ள கா்ப்பிணி பெண்கள்
இடம்	:	கா்ப்பிணி பெண்கள் புறநோயாளிகள் பிாிவு, மகப்பேறு மருத்துவமனை, எழும்பூா்
கற்பிக்கும் முறை	:	விரிவுடையாற்றலும், கலந்துரையாடலும்
கற்பிக்கும் மொழி	:	தமிழ்
துணைக் கருவி	:	புத்தகக் குறிப்பேடு
ஆயவாளர் பெயர்	:	ஜெ.தனலெட்சுமி, முதுநிலை இரண்டாமாண்டு மாணவி

பொதுவான நோக்கம்

கற்பித்தலின் முடிவில் குழு அதீக பாதீப்பிற்குாிய கா்ப்பத்தை பற்றிய அறிவினையும், அதன் அறிகுறிகளை எவ்வாறு ஆரம்பத்திலேயே கண்டறிவது என்பதையும், அதன் விளைவுகளை எவ்வாறு தடுக்க வேண்டும் என்பதையும் மற்றும் நீஜ வாழ்க்கையில் எவ்வாறு உபயோகப்படுத்த வேண்டும் என்பதையும் அறிய முடியும்.

சிறப்பு நோக்கங்கள்

காப்பிணி பெண்கள் கீழ்க்கண்ட திறன்களைப் பெறுவா

- 1. அதிக பாதிப்பிற்குரிய காப்பம் என்பதன் பொருள் யாது என்பதைக் கண்டறிவா
- 2. அதிக பாதிப்பிற்குரிய காப்பத்திற்கான காரணங்களை பட்டியலிடுவா
- 3. அதிக பாதிப்பிற்குரிய காப்பத்திற்கான அடையாளங்கள் அறிகுறிகள் இவற்றைக் கண்டறிவர்
- 4. அதிக பாதிப்பிற்குரிய காப்பத்திற்கான கண்டறியும் முறையை வரையறுப்பா
- 5. அதிக பாதிப்பிற்குரிய காப்பத்தினை தடுக்கும் முறையை புரிந்துகொள்வா
- 6. அதிக பாதிப்பிற்குரிய காப்பத்திற்கான சிகிச்சை முறையை விவரிப்பா
- 7. அதிக பாதிப்பிற்குரிய காப்பத்தினுடைய விளைவுகளை பட்டியலிடுவா்

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	ஆராய்ச்சியாளர் காரியத்திட்டத்தின் தலைப்பினை அறிமுகப்படுத்துதல்	முன்னுரை : அனைத்து கர்ப்பங்களும், பிரசவசங்களும் பாதிப்பிற்குரிய விஷயங்கள். இருப்பினும் ஒரு சில வகைகளில், தாய் மற்றும் சேய்யிற்கு அதிக விளைவுகள் ஏற்பட வாய்ப்புள்ளது. ஏறத்தாழ 20⊷30% கர்ப்பங்கள் இந்த வகையை சார்ந்தது. மகப்பேற்று நிலையை உயர்த்த வேண்டுவெனில், மேற்கண்ட குழுமம் முதலில் கண்டறியப்பட்டு சிகிச்சை அளிக்கப்பட வேண்டும்.	அறிமுகப்படுத்துதல்	கவனித்தல்	புத்தகக் குறிப்பேடு	
1 நிமிடம்	கற்றுக்கொள்பவர்கள் அதிக பாதிப்பிற்குரிய கர்ப்பம் என்பதன் வொருள் யாது என்பதைக் கண்டறிவர்	 அதிக பாதிப்பிற்குரிய கர்ப்பத்தின் அர்த்தம் : கர்ப்ப காலத்தில் பல வகையான நோய்கள் மற்றும் காரணங்கள் தாய் மற்றும் சேயிற்கு பாதிப்பினை ஏற்ப்படுத்துகிறது. இவ்வகை கர்ப்பங்கள் அதிக பாதிப்பிற்குரியவை என கருதப்படுகிறது. சில அதிக பிரச்சனைக்குரிய கர்ப்பங்கள் இரத்த சோகை கர்ப்பகால வலிப்பு நோய் கர்ப்பகால சர்க்கரை நோய் பல கருவுற்ற தன்றை 	அதன் அர்த்தத்தை விவரித்தல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	அதிக பாதிப்பிற்குரிய கர்ப்பம் என்றால் என்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் இரத்த சோகை என்பதன் பொருள் யாது என்பதைக் கண்டறிவர்	 இரத்த சோகை இரத்த சோகை உலகத்தில் அதிலும் குறிப்பாக கர்ப்பிணிப் வபண்களில் என்பது வபாதுவான பிரச்சனை. இந்நிலையில் ஹீமோகுளோபின் அளவு குறைவாக காணப்படும். அதாவது அதன் அளவு 11கி/வடசி.லி-க்கும் குறைவு. ஹீமோகுளோபின் என்பது இரத்தத்தில் உள்ள ஒரு சிவப்பனு ஆகும். இதன் முதன்மை வேலை உடலில் அனைத்து உறுப்புகளுக்கும் ஆக்ஸிஜனை எடுத்து வுசல்வதாகும். சராசரியான ஹீமோகுளோபின் அளவு 11–16 கி/வடசி லி. வாதுவாக 80% இருப்புச்சத்து குறைவினால் ஏற்படுகிறது. 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	கர்ப்பிணி வெண்ணின் ஹீமோகுளோபின் அளவு என்ன?
1 நிமிடம்	கற்றுக்கொள்பவர் இரத்த சோகை காரணங்களை பட்டியலிடுவர்	 அதிகப்படியான இரும்பு சத்து தேவை குறைவாக உணவு உட்கொள்ளுதல் உடல் அலைச்சல் கர்ப்பத்திற்கு முன்புள்ள உடல்நிலை இரத்த இழப்பு குறைந்த இடைவெளியில் அடிக்கடி கர்ப்பமாதல் 	விவரித்தல்	கவனித்தல்	புத்தகக் குறிப்பேடு	ூரத்த சோகைக்கான காரணங்களை குறிப்பிடுக?

நோம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளாின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	றதிப்பிடுதல்
2 நிமிடம்	கற்றுக்கொள்பவர்கள் இரத்த சோகை அடையாளங்கள் அறிகுறிகள் இவற்றைக் கண்டறிவர்	 அறிதறிகள் வெளிரிக் காணப்படுதல் குறிப்பாக கீழ் இமையின் உட்புறம், நகங்கள், நாக்கு, தோல் மற்றும் உள்ளங்கை. களைப்பு மூச்சுவிடுதலில் சிரமம் அதிகப் படியான இதய துடிப்பு தலைச்சுற்றல் சோம்பல் கவனம் செலுத்தலில் சிரமம் எடை இழப்பு திவிர இரத்த சோகையின் கரண்டி வடிவில் நகங்கள். எச்சரிக்கை அறிதறிகள் உள்ளங்கை மற்றும் கீழ் இமையின் உட்புறம் வெளிரி காணப்படுதல். சுவாம் ஒரு நிமிடத்திற்கு 30க்கும் அதிகமாக இருத்தல். அதிகமான களைப்பு ஒய்வு நேரத்திலும் மூச்சுவிடுவதில் சிரமம் 	விவரித்தல் விவரித்தல்	கவனித்தல் கவனித்தல்	புத்தகக் குறிப்பேடு	ூரத்த சோகையை எவ்வாறு கண்டறியலாம?
2 நிமிடம்	கற்றுக்கொள்பவர்கள் கண்டறியும் முறையை வரையறுப்பர்	 கண்டறியும் முறைகள் நோய்கள் உடல் அறிகுறிகள் ஆராய்வதன் மூலம் அளவை மதிப்பிடுதலின் மூலம் இரத்தப் வொருள் இணையும் திறன் மூலம் 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	ஞரத்தசோகை கண்டறியும் முறைகளை குறிப்பிடுக?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
2 நிமிடம்	கற்றுக் கொள்பவர்கள் இரத்த சோகை தடுக்கும் முறையை புரிந்துகொள்வர்	 தடுக்கும் முறைகள் இரத்த சோகையை தடுக்க கர்ப்பகாலத்திற்கு முன்பே ஆரம்பிக்க வேண்டும். ஆரம்ப கர்ப்பகால பரிசோதனை ஹீமோகுளோபிளின் அளவு குறைவினை ஆரம்பத்திலேயே கண்டறிதல் அடிக்கடி குழந்தைப்பேற்றினை தவிர்ப்பதன் மூலம் (குறைந்த யட்ச இடைவெளி 2 வருடங்கள்) இணை இரும்பு சத்து சிகிச்சை சத்தான உணவு வகைகளுடன் இணை இரும்பு சத்தும் தேவையானது. நாடாப்புளு தாக்குதலை தவிர்க்க பாதங்களை வசுரூப்புகள் பயன்படுத்தி பாதுகாக்க வேண்டும். கர்ப்பகாலத்தில் தேவையான வெரத்த இரும்பு சத்து 1000 மி.கி. ஆகும். புரதச் சத்து மற்றும் இருமபுசத்து அதிகமுள்ள உணவு வகைகளை உட்கொள்ளுதல் அவையாவன நிலக்கடலை, பீன்ஸ், அடர் பச்சை காய்கறிகள், கல்லீரல், பால், முட்டை, வெங்காயம் மற்றும் பேரீட்சைப் பழம். 	கலந்துரையடால்	கலந்துரையாடல்	புத்தக குறிப்பேடு	புரதம் மற்றும் இரும்புச் சத்து அதிகம் கொண்ட உணவு வகைகள் யாவை?

நோம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
2	கற்றுக் கொள்பவர்கள்	 மருத்துவரின் அறிவுப்படி தினசரி இரும்புச்சத்து மற்றும் போலிக் அளவு மாத்திரையை உட்கொள்ளுதல் அனைத்து கர்ப்பிணிப் வெண்களும் இவ்வகை மாத்திரைகளை 4வது மாதம் முதற்கொண்டு உட்கொள்ள வேண்டும். உணவு உண்டப்பின் தேநீர் அருந்தக்கூடாது. ஏவெனில் உணவுடன் தேநீர் அருந்தும் போது தேநீரில் உள்ள ஒரு வகை வொருள் இரும்புச்சத்து உறிஞ்சும் தன்மையை தடுக்கிறது. மலேரியா, இரத்தம் மற்றும் சிறுநீர்குழாய் சம்மந்தப்பட்ட நோய்களுக்கான சிகிச்சை 	கலந்துரையாடல் சிகிச்சை	புத்தகக் குறிப்பு கவனித்தல்	புத்தகக்	இரும்புச்சத்து
நிமிடம்	சிகிச்சை முறைகளை விவரிப்பர்	 உணவு அதிக புரதம், இரும்புச்சத்து மற்றும் வைட்டமின் கொண்ட சமச்சீர் உணவு பசியினை தூண்டவும், செறித்தலை அதிகரிக்கவும் வெய்சின் அதிகம் கொண்டு தயாரிக்கப்பட்ட வொருட்கள் உயர் எதிர் வொருளை பயன்படுத்தி குறைந்தபட்ச கிருமி தாக்குதலையும் இல்லாமல் செய்தல். இரத்தசோகை ஏற்படுத்தல் காரணிகளுக்கு சிகிச்சை இரும்புச் சத்துள்ள மருந்துகள்: வாய் வழி இரும்புச்சத்து மாத்திரை மற்றும் ஊசி மூலம் இரும்புச்சத்து ஏற்றுதல். உட்கொள்ளும்போது வயிற்று எரிச்சல், வாந்தி, கருப்பு நிற மலம் போன்ற பின்விளைவுகள் ஏற்படலாம். இவை சில நாட்களில் சரியாகிவிடும். இரத்தம் செலுத்துதல் 	முறைகளை விவரித்தல்		குறிப்பேடு	மாத்திரையின் பக்க விளைவுகள் என்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவாின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக் கொள்பவர்கள் இரத்த சோகை விளைவுகளை பட்டியலிடுவர்	 விளைவுகள் தாயிற்கு: நோய்தொற்று கர்ப்ப கால இரத்த அழுத்த நோய் கர்ப்பப்பை சுருங்காமை இதயக்கோளாறு அதிக இரத்தப்போக்கு மகப்பேறுக்குப்பின் நோய்தொற்று கர்ப்பயை சரிவர சுருங்காமை தாய்ப்பால் சுரத்தலில் குறைபாடு நுரையீரல் இரத்தக்குழாயில் அடைப்பு சிசுவின் வளர்ச்சி குறைபாடு குறைவான பிறப்பு எடை 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	ூரத்த சோகையினால் சிசுவிற்கு உண்டாகும் யாதிப்பு என்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக் கொள்பவர்கள் கர்ப்பகால வலிப்பு பொருள் யாது என்பதைக் கண்டறிவர்	ூய்த்தம்: கர்ப்பிணிப்வெண்ணில் சரியான இரத்த அழுத்தத்தின் அளவு 120/80 மிமீ பாதரச அழுத்தம். கர்ப்பகால வலிப்பு நோய் என்பது உயர் இரத்த அழுத்தம் அதாவது 140/90 மிமீ. பாதரச அழுத்தம் அல்லது அதற்கு அதிகமாகவும், சிறுநீரில் புரதம் உடன் வீக்கம் மற்றும் வீக்கமற்ற நிலை ஆகும். இது குறிப்பாக கர்ப்பகாலத்தில் 20வது வாரத்திற்கு மேல் ஏற்படும்.	விவரித்தல்	கவனித்தல்	புத்தகக் குறிப்பேடு	கர்ப்பகால வலிப்பு நோய் என்றால் என்ன?
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோய் பாதிக்கும் காரணிகளை பட்டியலிடுவர்	 பாதிக்கும் காரணிகள்: கர்ப்பகால வலிப்பு நோய் கொண்ட சரித்திரம் ஒருக்கும் நஞ்சுக்கொடி காரணிகள் அதிகமான நஞ்சுக்கொடியின் அளவு நஞ்சுக்கொடியின் இடமாற்றம் சிலுநீரகத்தில் ஏதேனும் நோயிந்தால் 20 வயதிற்கு முன் கர்ப்பமாதல் 30 வயதிக்குபின் கர்ப்பமாதல் உடல் பருமன் உயர் இரத்த அழுத்தம் கொண்ட குடும்ப சரித்திரம் மன அழுத்தம் முன்பே உள்ள சர்க்கரை நோய் 	விவரித்தல்	கவனித்தல்	புத்தக குறிப்பேடு	

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோயிற்கான ஆரம்ப அறிகுறிகளை கண்டறிவர்	 ஆரம்ப அறிகுறிகள் : உயர் இரத்த அழுத்தம் 140/90 மீ.மீ. பாதரச அழுத்தம் அல்லது அதற்கு மேல் வீக்கம் 12 மணி நேர ஒய்விற்கு பின்பும் முட்டியில் வீக்கம் வதாடர்ந்து காணப்படுதல் விரல் மோதிரம் இறுகி காணப்படுதல். வீக்கத்தை அழுத்தும் போது 2 மி.மீ. க்கு உள்ளே வுசன்றால் அது மிதமான வீக்கம், 4 மிமீ க்கு உள்ளே வுசன்றால் அது நடுத்தர வீக்கம் 6 மி.மீக்கு உள்ளே வுசன்றால் அது நடுத்தர வீக்கம் 6 மி.மீக்கு உள்ளே வுசன்றால் அது மாசமான நிலை 10 மி.மீ க்கு உள்ளே வுகன்றால் அது மிக மோசமான நிலை. எடை அதிகரிப்பு கர்ப்பகாலத்தில் சராசரி எடை அதிகரிப்பு 10 முதல் 12 கிலோ ஆகும். ஆனால் கர்ப்பகால வலிப்பு நோயில் எடை அதிகரிப்பு மாதத்திற்கு 2 கிலோவிற்கு மேல் இருக்கும். 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	கர்ப்பகாலத்தில் தாய்க்கு சராசரியாக எவ்வளவு எடை அதிகரிக்க வேண்டும்?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் சையல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைய்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோயிற்கான தீவிர நிலையில் உள்ள அறிகுறிகளை கண்டறிவர்	 தீவிர நிலையில் அறிதறிகள் தீவிர தொடர் உயர் இரத்த அழுத்தம் இரத்த அழுத்தத்தின் அளவு 160/110 மி.மீ யாதரச அழுத்தம் அல்லது அதற்கு அதிகமாக இருக்கும். வீக்கம், முகம், கைகள், வயிற்றுப் பகுதி, வென்னினப்வழுக்க உறுப்பின் வெளிப்பகுதி, முதுகுப்புறப்பின் பகுதி, யாதம் மற்றும் மற்ற உடல் பகுதிகளில் காணப்படும். விரைவான எடை அதிகரிப்பு 24 மணிநேர சிறுநீரில் புரதம் காணப்படுதல். அதாவது 30 கிராம்/வடசி.லி. தலை வலி முன்புற அல்லது பின்புறம் நிம்மதியற்ற உறக்கம் குறைவான சிறுநீர் வெளியேற்றம் 24 மணிநேரத்தில் சிறுநீர் வெறியேற்றம் 400 மி.லி க்கும் குறைவாக இருக்கும் வயிற்றுவலியுடன் கூடிய வாந்தி மங்கலான யார்வை குறைவான சிசு அரைவு 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	தீவிர கர்ப்ப கால வலிப்பு நோயின் அறிகுறிகளை கூறுக?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
		<u>எச்சரிக்கை அறிகுறிகள்</u> • தாங்கமுடியாத தலைவலி • யார்வை குறையாடு • குறைவான சிறுநீர் வெளியேற்றம் • மேல் வயிற்று வலி • கால் மற்றும் முகத்தில் வீக்கம்	விவரித்தல்	கவனித்தல்		
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோயினை கண்டறியும் முறையை வரையறுப்பர்	 இரத்த அழுத்தத்தை அளப்பதன் மூலம் 24 மணி நேர சிறுநீர் வெளியேற்றம் தாயின் எடையை அளப்பதன் மூலம் புரதத்திற்கான சிறுநீர் ஆய்வு சிசுவின் ஆரோக்கிய நிலையை கண்காணிப்பதன் மூலம் தினசரி சிசு உதைக்கும் எண்ணிக்கையை கணக்கிடுதல். மீவயாலி கதிர் சிதறல் (ஸ்கேன்) 	விவரித்தல்	கவனித்தல்	புத்தக குறிப்பேடு	கர்ப்பகால வலிப்பு நோய் கண்டறியும் முறைகளை கூறுக?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோயினை தடுக்கும முறையை பற்றி புரிந்துகொள்வர்	 தடுக்கும் முறைகள்: கர்ப்பகால வலிப்பு நோயை முழுமையாக தடுக்க முடியாது. பின்வருவனவற்றை பயன்படுத்த பாதிப்புக்குள்ளானவர்களை பாதிப்பிலிருந்து தடுக்கவோ அல்லது ஆரம்ப நிலையிலே கண்டுபிடிக்கவோ முடியும். தொடர் கர்ப்பகால பரிசோதனை முதல் 6 மாதங்களுக்கு மாதம் ஒரு முறையும், 7வது 8வது மாதங்களில் 15 நாட்களுக்கு ஒருமுறையும், அதற்கு பின் ஒவ்வொரு வாரமும் பரிசோதனைக்கு செல்ல வேண்டும். பாதிப்புக்குள்ளவர்கள் குறைந்தது 10 பரிசோதனையாவது செய்ய வேண்டும். போதுமான ஒய்வு 20 வது வாரத்தின் தொடக்கத்திலிருந்த 8–12 மணி நேர உறக்கம். இதில் 2 மணி நேரம் பிற்பகலிலும், 8–10 மணி நேரம் இரவிலும் உறங்க வேண்டும். உணவு வகைகள், புரதச்சத்து நிறைந்த உணவுப் வாருட்கள் கண்டறியும் ஆய்வு டாப்ளர் கொண்டு மதிப்பிடுதல் 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	ஒரு கர்ப்பிணி வெண் எவ்வளவு நேரம் உறங்கலாம்.

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	றதிப்பிடுதல்
		 ரோல் ஒவர் ஆய்வு– இதனை கர்ப்பகாலத்தில் 28–32 வாரங்களில் செய்ய வேண்டும். முதலில் தாயினை இடது புறமாக சாய்நத நிலையில் படுக்குமாறு செய்து இரத்த அழுத்தத்தினை பரிசோதிக்க வேண்டும். பின்பு திரும்ப செய்து மீண்டும் பரிசோதிக்க வேண்டும். அப்போது டையஸ்டோலிக் அழுத்தம் 20 மி.மீ. பாதரச அழுத்தத்தற்கு அதிகமாக இருந்தால் அது கர்ப்பகால வலிப்பு நோய் இருப்பதற்கு சாதகமான நிலையாகும். 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	கர்ப்பகால வலிப்பு நோய் உள்ள வெண் குறைந்தது எத்தனை முறை பரிசோதனைக்கு வர வேண்டும்?
2	கற்றுக்கொள்பவர்கள்	சிகிச்சையின் முறைகள்	சிகிச்சை	கவனித்தல்	புத்தக	
நிமிடம்	கர்ப்பகால வலிப்பு நோயிற்கான சிகிச்சை முறையை விவரிப்பர்	 யோதுமான ஒய்வு : உறங்கும்போது இடதுபுறமாக சாய்ந்த நிலையில் படுக்க வேண்டும். உணவுகள் பூரதம் நிறைந்த உணவு (வொதுவாக 100 கி/நாள்) உய்புசத்தினை அதிகமாகவோ அல்லது தவிர்க்கவோ கூடாது. வமாத்த கலோரி சராசரியாக 1600 கலோரி/நாள். வநாத்த அழுத்த பரிசோதனை வதாடர் எடை அளப்பு ஆய்வக பரிசோதனை குறிப்பாக, சிறுநீரில் சர்க்கரை மற்றும் புரதத்திற்கான ஆய்வு மற்றும் இரத்த பரிசோதனை கர்ப்பக்கால வலிப்பு நோய்க்கான அறிகுறிகளை கண்காணித்தல் அதிகப்படியான உடலியக்கத்தை தவிர்த்தல் 	முறைகளை விவரித்தல்		குறிப்பேடு	

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
		 மன அழுத்தம் மற்றும் பயத்தை குறைத்தல்– புத்தகத்தை படிப்பதன் முலம் தொலைக்காட்சியை பார்ப்பதன் மூலமும் மன அழுத்தத்தை குறைக்கலாம். தினமும் குழந்தை உதைக்கும் எண்ணிக்கையை கணக்கிடுதல்– காலை 10 மணியிலிருந்து கணக்கெடுப்பு ஆரம்பிக்க வேண்டும். 10 அசைவாவது 1 நாளில் தெரிய வேண்டும். வதாடர் 2 நாட்ளில் 12 நேரத்தில் 10 அசைவிற்கும் குறைவாகவோ அல்லத அசைவற்ற நிலையையோ உணர்ந்தால் மருத்துவரிடம் வதரிவிக்க வேண்டும். உயர் குரத்த அழுத்தத்திற்கான மருந்துகள் 	விவரித்தல்	கவளித்தல்		

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவரின் சையல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால வலிப்பு நோயின் விளைவுகளை பட்டியலிடுவர்	 விளைவுகள் தாயிற்கு குறைவான சிறுநீர் வெளியேற்றம் அல்லது சீறுநீர் வெளியேற்றமின்மை எதிர்பராத இரத்தக்கசிவு நோய்தொற்று பார்வை குறைவு அல்லது பார்வை இழப்பு இரத்தசோகை கல்லீரல் நாளச்சுரப்பி அளவு அதிகரித்தல் தட்டணுக்களின் அளவு குறைதல் வலிப்பு. கேரயக்கு: கருப்பையில் சிசுவின் வளர்ச்சி குறையாடு குறை பிரசவம் கர்பய்யையிலே சிசுவின் இறப்பு யுச்சு தினறல் 	விவரித்தல்	கவனித்தல்	புத்த குறிப்பேடு	கர்ப்பகால வலிப்பு நோயின் விளைவுகளை குறிப்பிடுக?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால சர்க்கரை நோய் என்பதன் வொருள் யாது என்பதைக் கண்டறிவர்	ூர்த்தம் : கர்ப்பகால சர்க்கரை நோய் என்பது கர்ப்பகாலத்தின் போது ஞரத்தச்சர்க்கரையின் அளவு அதிகமாக காணப்படுதல் ஆகும். சரியான ஞரத்த சர்க்கரையின் அளவு 80–120 மிகி/வெசி.லி. ஆகும். ஞது உடலில் ஞன்சுலின் குறைபாட்டினால் ஏற்படுகிறது.	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	கர்ப்பகால சர்க்கரை நோய் என்றால் என்ன?
1 நிமிடம்	கற்றுக்கொ்பவர்கள் கர்ப்பகால சர்க்கரை நோய் பதிக்கும காரணிகளை பட்டியலிடுவர்	 பாதிக்கும் காரணிகள் இரத்த சர்க்கரை கொண்ட குடும்ப சரித்திரம் பிறப்பன் போது குழந்தை இறப்பு சரித்திரம் காரணமில்லாத சிசு இழப்பு உடல் பருமன் 30 வயதிற்கு மேல் கர்ப்பமாதல் உயர் இரத்த அழுத்தம் அதிகப்படியான கருப்பை திரவத்தின் அளவு பற்றாகுறையான வளர்சிதை மாற்றம் 5 அல்லது அதற்கு மேற்பட்ட கர்ப்பங்கள் 	விவரித்தல்	கவனித்தல்	புத்தக குறிப்பேடு	கர்ப்பகால சர்க்கரை நோயின் அறிகுறிகளை கூறுக?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால சர்க்கரை நோயிற்கான அறிகுறிகளை கண்டறிவர்	 அறிதறிகள் அதிகப்படியான தாகம் அதிகப்படியான சிறுநீர் வெளியேற்றம் அதிகப்படியான பசி ஒருப்பினும் எடை இழப்பு களைப்பு, குமட்டல், வாந்தி சிறுநீர்ப்பை, பெண்ணினப்பெருக்க உறுப்பின் வெளிப்பகுதி மற்றும் தோலில் நோய்தொற்று யரிசோதனையின் சிசுவின் அளவு சராசரி அளவினை அதிகமாக இருக்கும். <u>எச்சரிக்கை அறிகுறிகள்</u> பார்வை கோளாறு அதிகமான களைப்பு அதிகமான சிறுநீர் வெளியேற்றம் மற்றும் தாகம் 	விவரித்தல்	கவனித்தல்	புத்தக குறிப்பேடு	கர்ப்பகால சர்க்கரை நோயின் அறிகுறிகளை கூறுக?
1 நிமிடம்	கற்றுக் கொள்பவர்கள் கர்ப்பகால சர்க்கரை நோயினை கண்டறியும் முறையை வரையறுப்பர்	 கண்டறியும் முறைகள்: யாஸ்டிங் மற்றும் யோஸ்பிராண்டியல் இரத்த சர்க்கரை அளவு பரிசோதித்தல். குளுகோஸ் தாங்கு திறன் ஆய்வு அசிட்டோன் இருப்பதற்கான சிறுநீர் பரிசோதனை மீவயாலி வரைவு (ஸ்கேன்) 	கலந்துரையாடல்	கலந்துரையடால்	புத்தகக் குறியீட்டு	கர்ப்கால சர்க்கரை நோயினை கண்டறியும் முறைகள் என்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவாரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடூதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால சர்க்கரை நோயை தடுக்கும் முறையை புரிந்துகொள்வர்	தடுப்பு முறைகள் எடை குறைப்பு உணவு முறை மாற்றம் உடற்பயிற்சி 	விவரித்தல்	கவனித்தல்	புத்தகக் குறியீட்டு	
2 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால சர்க்கரை சிகிச்சசை முறையை விவரிப்பர்	 சிகிச்சை முறைகள் : கர்ப்பகால மேற்பார்வை குறிப்பிட்ட கால ஒடைவெளியில் ஒரத்த சர்க்கரையின் அளவை யரிசோதிப்பது அனைத்து கர்ப்பிணி வபண்களும் இரத்த சர்க்கரையின் அளவை யரிசோதிக்க வேண்டும். ஃயாஸ்டிங்க் இரத்த சர்க்கரையின் அளவை 95மிகி/வடசிலி அளவுக்கு குறைவாக வைத்துக்கொள்ள வேண்டும். சிறந்த உணவு வகைகள் அதிகப்படியான நார்சத்தினை உட்கொள்ள வேண்டும். கொழுப்பு மற்றும் அடர் சுவைமிக்க இனிப்பு பண்டங்களை தவிர்க்க வேண்டும். தினசரி உடற்பயிற்சி இன்சுலின் (தேவைப்பட்டால்) 	விவரித்தல்	கவனித்தல்	புத்தகக் குறியீட்டு	கர்ப்பக்கால சர்க்கரை நோயில் கடைபிழக்க வேண்டிய உணவு வகைகள் என்வென்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் கர்ப்பகால சர்க்கரை நோயின் விளைவுகளை பட்டியலிடுவர்	 விளைவுகள் தாயிற்கு கருச்சிதைவு குறைமாத பிரசவம் நோய் தொற்று கர்ப்பகால வலிப்பு நோய் அதிகப்படியான கருப்பை திரவம் நீண்ட நேர பிரசவம் நீண்ட நேர பிரசவம் தாய்ப்பால் சுரத்தலின் அளவு குறைவாக காணப்படுதல் வபண்ணினப்பெருக்க உறுப்பில் காயங்கள் எதிர் காலத்தில் தாயிற்கு சர்க்கரை நோய் ஏற்பட வாய்ப்பு சோய்பிற்கு : இதய நோய் போன்ற பிறவிக் கோளாறு பிறப்புக் காயங்கள் இரத்தச் சர்க்கரையின் அளவு குறைந்து காணப்படுதல் வுரச்சுத்தின்றல் 	விவரித்தல்	கவனித்தல்	புத்தகக் குறிப்பேடு	கர்ப்பகால சர்க்கரை நோயினால் தாயிற்கு ஏற்படும விளைவுகள் என்ன?

நோம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்றத் தன்மை என்பதன் வொருளை அறிவர்	ூர்த்தம் : பல கருவுற்ற தன்மை என்றால் ஞரண்டு அல்லது அதற்கு மேற்யட்ட சிசு கருப்பையில் வளர்வது ஆகும்.	கலந்துரையாடல்	கலந்துரையாடல்	புத்தகக் குறிப்பேடு	பல கருவுற்ற தன்மை என்றால் என்ன?
1 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்ற தன்மைக்கான பாதிக்கும் காரணிகளை பட்டியலிடுவர்	யாதிக்கும் காரணிகள்: • பாரம்பரியம் • முதிர்ச்சிப் பருவம் • அதிக குழந்தைப்பிறப்பு • ூனம் • செயற்கைமுறை கருத்தரிப்பு	விவரித்தல்	கவனித்தல்	புத்தக குறிப்பேடு	பல கருவுற்ற தன்மையை உண்டாக்கும் காரணிகள் என்ன?
1 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்ற தன்மைக்கான அறிகுறிகளை கண்டறிவர்	 அறிகுறிகள்: கர்ப்பப்பை அளவு தேவையான கர்ப்பங்காலத்தை விட வெரியதாக இருத்தல். அதிகப்படியான குமட்டல் மற்றும் வாந்தி அதிகப்படியான பசி அதிகப்படியான எடை உயர்வு வயிற்றுப்பகுதியில் ஒரே சமயத்தில் வெவ்வேறு பகுதியில் குழுந்தை அசைவினை உணர்தல் 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	பல கருவுற்ற தன்மையின் அறிகுறிகள் என்ன?

நேரம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாளரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்ற தன்மையினை கண்டறியும் முறையை வரையறுப்பர்	கடண்டறியும் முறைகள்: • மீவயாலி வரைவு • உடல் யரிசோதனை • இரத்த யரிசோதனை • ஆல்பா பீட்டா புரதத்தின் அளவு	விவரித்தல்	கவளித்தல்	புத்தக பதிவேடு	பல கருவுற்ற தன்மையை கண்டறியும் முறைகளை கூறுக?
2 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்ற தன்மைக்கான சிகிச்சை முறையை விவரிப்பர்	 சிகிச்சை முறைகள் : அதிகப்படியான சத்துணவு அதிக கலோரி, பூதம் மற்றும் ஒரும்புச் சத்து முக்கியம் அடிக்கடி முன்பேறுக்கால பரிசோதனை அதிகப்படியான ஒய்வு சிசுவின் வளர்ச்சியை கண்காணித்தல் தாய் மற்றும் சிசு நலப்பரிசோதனை டோகோலைட்டிக்ஸ் (குறைமாத பிரசவமாயிருப்பின்) சர்வீகல் என்சர்க்கலேஜ் (தாங்குதிறனற்ற கர்ப்பப்பையிருந்தால்) இணை ஒரும்புச்சத்து சிகிச்சை 60–80 மி.கி/வெசி.லி அளவுக்கு ஒரும்புச்சத்து உயர்த்த வேண்டும். 	கலந்துரையாடல்	கலந்துரையாடல்	புத்தக குறிப்பேடு	கல கருவுற்ற தன்மையின் சிகிச்சை முறைகளை குறிப்பிடுக?

நோம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவரின் செயல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
1 நிமிடம்	கற்றுக்கொள்பவர்கள் பல கருவுற்று தன்மையின் விளைவுகளை பட்டியலிடுவர்	 விளைவுகள் தாயிற்க : குமட்டல் மற்றும் வாந்தி அதிக முறையும், தீவிரமாகவும் குருக்கும். கர்ப்பகால வலிப்பு நோய் மற்றும் இரத்தசோகை இயற்கைககு புறம்பான கருப்பைத் திரவம் அறுவை சிசிச்சைக்கு அதிகப்படியான வாய்ப்பு இரத்தப்போக்கு பனிக்குடம் முன்னமே உடைதல் நீண்ட நேர பிரசவம் சேய்யிற்க : பிறவிக் குறைபாடு குறை பிரசவம் வளர்ச்சிக் குறைபாடு 	விவரித்தல்	கவளித்தல்	புத்தக குறிப்பேடு	பல கருவுற்ற தன்மையின் விளைவுகளை குறிப்பிடுக?

நோம்	குறிக்கோள்	வொருளடக்கம்	ஆராய்ச்சியாவரின் சையல்	கற்றுக் கொள்பவரின் செயல்	ஒலி, ஒளி அமைப்பு	மதிப்பிடுதல்
		 கருப்பையிலேயே சிசுவின் இறப்பு மூச்சுத்திணறல் குழந்தைப் பிறப்பின் போது இறத்தல் 	விவரித்தல்	கவனித்தல்		
1 நிமிடம்		முடிவுறை அனைத்து கர்ப்பிணிப் வபண்களும் பிரச்சனைகளை சந்திப்பதில்லை. ஆனால் பிரச்சனைகளை பற்றி அறிவதன் முலம் அதன் விளைவுகளை தடுக்க முடியும்.				

வினாவிற்கான விடைகள்

கேள்வி எண்	விடைகள்	மதிப்பெண்
I.இரத்த சோகை		
1	A	1
2	ஆ	1
3	@	1
4	୬	1
5	அ	1
6	୬	1
7	୬	1
8	ஆ	1
II.கர்ப்பகால		
வலிப்பு நோய்		
1	A	1
2	அ	1
3	g	1
4	A	1
5	ஆ	1
6	8	1
7	அ	1
8	ஆ	1

III.கர்ப்பகால	விடைகள்	மதிப்பெண்
சர்க்கரை நோய்		
1	ஆ	1
2	8	1
3	அ	1
4	ஆ	1
5	g	1
6	9	1
7	இ	1
IV.பல கருவுற்ற		
தன்மை		
1	9	1
2	அ	1
3	9	1
4	अ	1
5	9	1
6	9	1
7	9	1

PATIENT CONSENT FORM

TITLE: A Pre-Experimental study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology,Chennai-08

Name of the Participant	:			
Date	:			
Age/sex	:			
Name of the Principal Ir	vestigator:	J. Dhanalak	shmi	
Name of the Institution	:	Institute of O	bstetrics and Gyr	necology
Enrollment No	:			
Documentation of the	Informed C	onsent: (lega	1 representative	can sion i

Documentation of the Informed Consent: (legal representative can sign if the participant is minor or incompetent)

- I have read the information in this form (or it has been read to me). I was free to ask any questions and they have been answered. I am over 18 years of age and exercising my free power of choice, hereby give my consent to be included as a participant in the study.
- I have read and understood this consent form and the information provided to me.
- I had the consent document explained in detail to me.
- I have been explained about the nature of my study.
- My rights and responsibilities have been explained to me by the investigator.
- I am aware of the fact that I can option out of the study at any time without having to give any reason and this will not affect my future treatment in this hospital.
- I hereby give permission to the investigator to release the information obtained from me as result of participation in this study to the sponsors, regulatory

authorities, Government agencies and IECI, understand that they are publicly presented.

- I have had my questions answered to my satisfaction.
- I have decided to be in the research study.
- I am aware that if I have any question during this study, I should contact the investigator. By signing this consent form I attest that that the information given in this document has been clearly explained to me and understood by me. I will be given a copy of this consent document.

1. Name and Signature/thumb impression of the participant (or legal representative if participant incompetent)

Name	 	 	
Signature_	 	 	

Date_____

Name and Signature of the Investigator or her representative obtaining consent:

Name			 	

Signature_____

Date_____

INFORMATION TO PARTICIPANTS

TITLE: A Pre-Experimental study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology,Chennai-08

Name of the Participant	:	
Date	:	
Age/sex	:	
Name of the Principal In	vestigator:	J. Dhanalakshmi
Name of the Institution	:	Institute of Obstetrics and Gynecology
Enrollment No	:	

You are invited to take part in this research/study/procedures. The information in this document is meant to help you decide whether or not to take part. Please feel free to ask if you have any queries or concerns.

You are being asked to participate in this study being conducted at Institute of Obstetrics and Gynecology, Chennai-08.

What is the purpose of the Research (explain briefly)

This research is conducted to assess the effectiveness of structured teaching programme on knowledge regarding early identification of high risk pregnancy among antenatal mothers attending at institute of obstetrics and gynaecology,chennai-08

We have obtained permission from the Institutional Ethical Committee.

Study Design

Pre experimental research design (One group pre-test post-test design)

Study procedure

1. The study will be undertaken after approval from Institutional Ethics Committee.

- 2. Those who are willing to participate will be enrolled and informed consent will be obtained.
- 3. Women who fulfill the inclusion criteria and exclusion criteria are selected in the group.
- The level of awareness about early identification of high risk pregnancy is assessed with structured questionnaire of antenatal mothers with gestational age 8-28 weeks.
- 5. STP for 30 minutes regarding early identification of high risk pregnancy which includes aspects like meaning, causes, signs and symptoms, diagnosis, prevention, management and complications of high risk pregnancy.
- 6. After one week period, post-test will be conducted with the same structured pretest questionnaire for analyzing the effectiveness of STP.
- 7. The results of the study will be analyzed by using descriptive and inferential statistics.

Possible risks to you-Briefly Mention.

No risks involved.

Possible benefits to you

After finishing the study, investigator will provide information about

- meaning of high risk pregnancies like anemia, pre-eclampsia, gestational diabetes mellitus and multiple pregnancy.
- causes and risks of high risk pregnancy.
- signs and symptoms of high risk pregnancy.
- investigations of high risk pregnancy
- prevention and management of high risk pregnancy.
- complications of high risk pregnancy.

Possible benefits to other people

The result of the research may provide benefit to the society in terms of advancement in medical knowledge and/or early identification of high risk pregnancy in future.

Confidentiality of the information obtained from you

You have the right to confidentiality regarding the privacy of your medical information (personal details, results of physical examinations, investigations, and your medical history). The information from this study, if published in scientific journals or presented at scientific meetings, will not reveal your identity.

Your privacy in the research will be maintained throughout the study in the event of any publication or presentation resulting from research, no personal identity information will be shared.

How will your decision not to participate in the study affect you?

Your decision not to participate in this research study will not affect your daily living activities, medical care or your relationship with investigator or the institution.

Can you decide to stop participating in the study once you start?

The participation in this research is purely voluntary and you have the right to withdraw from this study at any time during the course of the study without giving any reasons. The result of the study will be informed to you at the end of the study.

Signature of the Investigator Date:

Signature of the Participant Date:

<u> ஆராய்ச்சி தகவல் தாள்</u>

<u>ஆராய்ச்சி தலைப்பு:</u> அதிக பாதிப்பிற்குரிய கர்ப்பம் பற்றிய பாடத்திட்டத்தினை கற்பித்தலின் மூலம், அதனால் தாய்மார்களுக்கு ஏற்பட்ட அறிவினை கண்டறிதல்.

ஆய்வாளர் பெயர் : ஜெ.தனலெட்சுமி

பங்கேற்பாளா் பெயா் :

வயது:

ஆராய்ச்சி சேர்க்கை எண்:

தேதி:

நான் அரசு தாய் சேய் நல மருத்துவமனையின் கா்ப்பிணிகள், வெளிநோயாளிகள் பகுதியில் பாிசோதனைக்காக வரும் தாய்மாா்களை திறனாய்வு மேற்கொள்கிறேன்.

அதிக பாதிப்பிற்குரிய கா்ப்பம் பற்றிய பாடத்திட்டத்தினை கா்ப்பிணிகளுக்கு கற்பித்து, அதற்கு முன் உள்ள அறிவினையும், பாடம் கேட்டு தெரிந்து கொண்ட பின் உள்ள அறிவினையும், அறிய போகிறேன்.

இந்த செயல்முறையின் மூலம், கா்ப்பிணி தாய்மாா்களுக்கு, அதிக பாதிப்பிற்குாிய கா்ப்பத்தினை, முன்கூட்டியே அறியும் வாய்ப்பு அதிகம் உள்ளது. இம்முறையை தாய்மாா்கள் நன்றாக பயன்படுத்திக் கொள்ளலாம்.

தாய்மாா்கள் தங்கள் சொந்த விருப்பத்தின் போில் ஆராய்ச்சியில் இணைக்கப்படுவா். விருப்பமில்லை என்றால் எந்நேரமும் விலகிக் கொள்ளலாம். இதனால் ஆராய்ச்சிக்கு எந்தவித பாதிப்பும் ஏற்படாது.

முடிவுகளை அல்லது கருத்துக்களை வெளியிடும் போது தங்களின் பெயரையோ அல்லது அடையாளங்களையோ வெளியிட மாட்டேன் என்பதை தெரிவித்துக் கொள்கீறேன்.

ஆராய்ச்சியாளா் கையொப்பம்

பங்கேற்பாளா் கையொப்பம்

தேதி:

தேதி :

<u> ஆராய்ச்சி ஒப்புதல் பழவம்</u>

ஆராய்ச்சி தலைப்பு	:	அதிக பாதிப்பிற்குரிய கர்ப்பம் பற்றிய பாடத்திட்டத்தினை கற்பித்தலின் மூலம், அதனால் தாய்மார்களுக்கு ஏற்பட்ட அறிவினைக் கண்டறிதல்
ஆய்வு நடத்தப்படும் இடம்	:	கா்ப்பிணிகள் புறநோயாளிகள் பிரிவு
ஆராய்ச்சியாளர் பெயர்	:	ஜெ.தனலெட்சுமி
பங்குபெறுபவரின் பெயர்	:	வயது:

பங்குபெறுபவரின் புறநோயாளி எண். :

இந்த ஆராய்ச்சியின் விவரங்களும் அதன் நோக்கங்களும் முழுமையாக எனக்கு விளக்கப்பட்டது.

எனக்கு விளக்கப்பட்ட விஷயங்களை நான் புரிந்து கொண்டு நான் எனது சம்மதத்தை தெரிவிக்கீறேன்.

இந்த ஆராய்ச்சியில் பிறரின் நீர்பந்தமின்றி நான் எனது சொந்த விருப்பத்தின் பேரில் பங்கு கொள்கீறேன். மற்றும் நான் இந்த ஆராய்ச்சியிலிருந்து எந்த நேரமும் பின்வாங்கலாம் என்பதையும் அதனால் எவ்வித பாதிப்பும் ஏற்படாது என்பதையும் புரிந்து கொண்டேன்.

இந்த ஆராய்சியின் தகவல்களை வெளியிட சம்மதிக்கிறேன். அப்படி வெளியிடும் போது என் அடையாளம் வெளிவராது என்பதை அறிவேன்.

நான் இந்த ஆராய்ச்சிக்கு என்னுடைய முழு ஒப்புதலை அளிக்கீறேன். எனக்கு இந்த ஒப்புதல் கடிதத்தின் நகல் கொடுக்கப்பட்டது.

ஆராய்ச்சியாளர் கையொப்பம்

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3	1	2	4	1	1	25	G3P1L1A1	28
2	1	2	4	1	1	19	PRIMI	
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PRE-TEST SCORE																													
		i	ane	mia		- 1		pre-eclampsia GDM											r	multiple gestation									
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		i	ane	mia	1				pre-eclampsia									Ģ	SDN	1			multiple gestation							
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F anemia pre eclampsia											051	-1ES	TEST GDM multiple gestation																
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			ane	mia						pre	e ecl	amp	osia					(GDN	Λ			multiple gestation						
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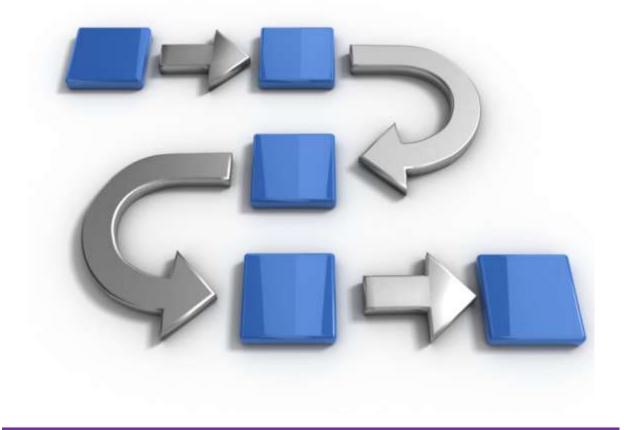
Chapter-I Introduction



Chapter-II Review of Literature



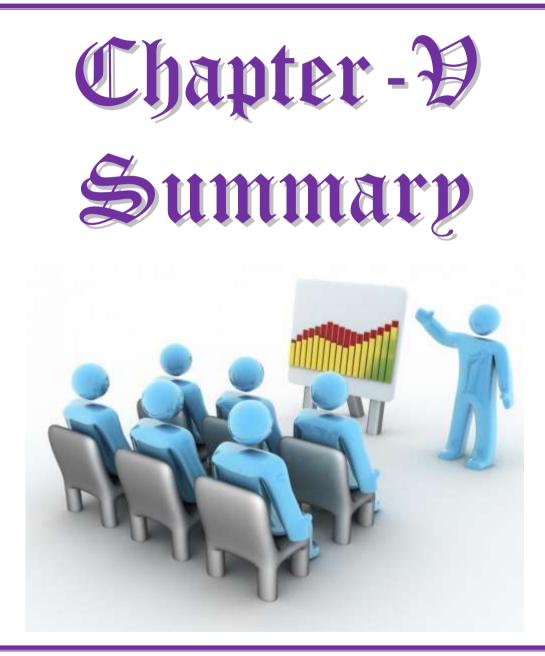
Chapter-III Research Methodology





Interpretation





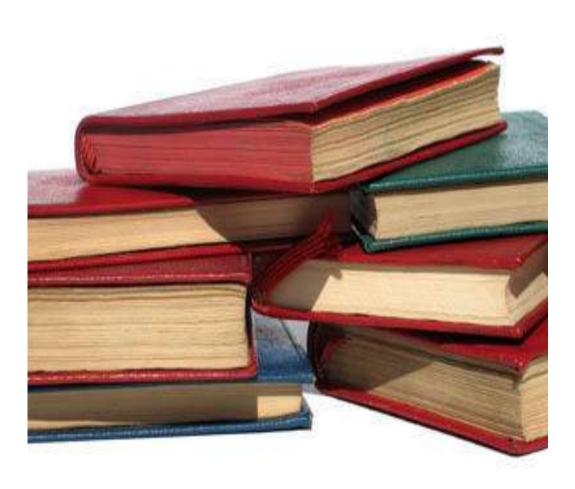


No.

Chapter - HII Conclusion & Recommendation











10G & Government hospital for

Women & children . Egmore.

Chennai. 08 dated: 01.06.2017.

Sub: Training MSc nursing II year- obstetrics & Gynecological nursing – clinical practice Dissertation, Practical examination and Lecture training in IOG & Government hospital for women & children. Egmore. Chennai, 08 dated: 31.03.2017 for the period from 20/11/2016 to 17/12/2016 Permission. Orders issued.

Ref: letter dated 20/11/2016 of the Head of Department. O & G Nursing, college of nursing. Madras Medical college, Chennai - 3.

As per the letter reference cited, the following the M.Sc II years students of Madras Medical College, chennai -3 are permitted to undergo the clinical experience, lectures classes. University practical examination and also to carry out dissertation work in IOG & Government hospital for women & children. Egmore. Chennai- 8 for the period from 20/11/2016 to 17/12/2016 under the guidance of the Assistant Professor of O & G mentioned against their names.

S.No	Name of the students	Name of the Assistant professor of & G of this hospital.							
1	Mrs. Alagirisamy anitha	Dr. kavitha							
2	Mrs. R. Bama	Dr. sumathi							
3	Mrs. J. Dhanalakshmi	Dr. A. vijayalakshmi							
4	Ms. B. Hemalatha	Dr. P. Priyadharshini.							
5	Ms. R. Revathi	Dr. R.Sridevi							
6	Mrs. P. Savitha	Dr. sadhana.							
7	Mrs. Shanthi Grace	Dr.chandrakala							
8	Mrs. Subbulakshmi	Dr. K. Priyadharshini.							

Director and superintendent elrector and Superintendes Institute of Obstetrics and wasscology and Govt. Hosphus for Women and Children. BMORE, MADRAS-R

To

The individuals concerned

Copy to

Dr. A. vijayalakshmi Assistant Professor of O & G, IOG and Government Hospital for women and Children, Egmore. Chennai- 8.

CERTIFICATE OF CONTENT VALIDITY

This is to certify that the tool constructed by Ms. J.Dhanalakshmi, M.Sc., (Nursing) II year, College of Nursing, Madras Medical College which is to be used in her study titled, "A Study to Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High risk Pregnancy Among Antenatal Mothers Attending At Institute of Obstetrics and Gynecology, Egmore, Chennai-08" has been validated by the undersigned. The suggestions and modifications given by me will be incorporated by the investigator in concern with their respective guide. Then she can proceed to do the research.

Signature with seal

PROF. Dr. ROSALINE RACHEL, M.S.C., (M), Ph.D.(N) PRINCIPAL MMM COLLEGE OF NURSING No.131, SAKTHI NAGAR, NO.131

Place: Chennai. Date: 7-11-06.

INSTITUTIONAL ETHICS COMMITTEE MADRAS MEDICAL COLLEGE, CHENNAI 600 003

EC Reg.No.ECR/270/Inst./TN/2013 Telephone No.044 25305301 Fax: 011 25363970

CERTIFICATE OF APPROVAL

To J.Dhanalakshmi I Year M.Sc.(Nursing) Student College of Nursing Madras Medical College Chennai 600 003

Dear J.Dhanalakshmi,

The Institutional Ethics Committee has considered your request and approved your study titled "A PRE-EXPERIMENTAL STUDY IN EVALUATE THE EFFECTIVENESS OF STRUCTURED TEACHING PROGRAMME ON KNOWLEDGE REGARDING EARLY IDENTIFICATION OF HIGH RISK PREGNANCY AMONG ANTENATAL MOTHERS ATTENDING AT INSTITUTE OF OBSTETRICS AND GYNAECOLOGY, CHENNAI 600 008 " NO. 19072016.

The following members of Ethics Committee were present in the meeting hold on **12.07.2016** conducted at Madras Medical College, Chennai 3

1. Prof. C. Rajendran, MD.	Chairperson
2. Prof. Isaac Christian Moses, MD., Dean(FAC)MMC, Ch-3 Dep	uty Chairperson
3. Prof. Sudha Seshayyan, MD., Vice Principal, MMC. Ch- 3. Me	mber Secretary
4. Prof. B. Vasanthi, MD., Prof of Pharmacology, MMC.	Member
5. Prof. P.Raghumani.MS., Professor of Surgery, Inst. of surgery	Member
6. Prof. Md Ali, MD., DM., Prof & HOD of MGE, MMC, Ch-3.	Member
7. Prof. Baby Vasumathi., MD, Director. Inst. of O&G.	Member
8. Prof. K.Ramadevi., MD, Director, Inst of Bio-Chemistry, MMC.	Member
9. Prof. R.Padmavathy, MD., Professor, Inst. of Pathology, MMC. Ch	Member
10. Prof.S. Tito, MD, Director, Inst. of Inter Med, Ch-3.	Member
11.Tmt.J.Rajalakshmi, Junior Administrative Officer, MMC.Ch	Layperson
12. Thiru.S. Govindasamy., B.A.B.L., High Court, Chennai-1	Lawyer
13.Tmt.ArnoldSaulina, MA., MSW.,	Social Scientist

We approve the proposal to be conducted in its presented form.

The Institutional Ethics Committee expects to be informed about the progress of the study and SAE occurring in the course of the study, any changes in the protocol and patients information/informed consent and asks to be provided a copy of the final report.

Member Secretary MEMBER SECRETARY MADRAS MEDICAL COLLEGE CHENNAI-500 003

Lr. No. 40 (S) (CON. mme, dt. 19.11.16. DRMD BRMD 11.16

From

J. Dhanalakshmi,

M.Sc., (N) II year,

College of Nursing,

Madras Medical College,

Chennai -03.

To

The Director,

Institute of Obstetrics and Gynecology Hospital for Women and Children,

Egmore,

Chennai -08.

Through

Principal,

College of Nursing, Madras Medical College, Chennai -03.

Respected Sir/Madam,

Sub: Requesting permission to conduct research at Institute of Obstetrics and

Gynecology Hospital for Women and Children. Chennai -08.

I, M.Sc., Nursing II year student have to conduct the research study for the fulfillment of M.Sc.,(N) Programme. My topic is "A Pre-Experimental study to assess the effectiveness of Structured teaching programme on early identification of high risk pregnancy among antenatal mothers attending at Institute of Obstetrics and Gynecology" from 20-11-2016 to 18-12-2016. I assure that I will not disturb the routine activities of the OPD.

With due respect, I request your good self to kindly permit me to conduct this research study.

Forwarded

Thanking you,

(9/11/1b

DR. V. KUMARI, M.Sc(N)., Ph.D., PRINCIPAL COLLEGE OF NURSING MADRAS MEDICAL COLLEGE CHENNAI - 600 003.

Yours sincerely,

analaki

(J. Dhanalakshmi)

CERTIFICATE OF CONTENT VALIDITY

This is to certify that the tool constructed by Ms. J.Dhanalakshmi, M.Sc., (Nursing) II year, College of Nursing, Madras Medical College which is to be used in her study titled, "A **Pre-Experimental study To Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High Risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08**" has been validated by the undersigned. The suggestion and modifications given by me will be incorporated by the investigator in concern with their respective guide. Then she can proceed to do the research.

Name: Dr. A.VIJAYALAKSHMI Designation: Asst professor Signature with Seal 27[7][17 Assistant Surgeon I.O.G. & Government Hospital For Women and Children Egmore, Chennai-8.

CERTIFICATE OF CONTENT VALIDITY

This is to certify that the tool constructed by Ms. J.Dhanalakshmi, M.Sc., (Nursing) II year, College of Nursing, Madras Medical College which is to be used in her study titled, "A Study to Evaluate The Effectiveness Of Structured Teaching Programme on Knowledge Regarding Early Identification Of High risk Pregnancy Among Antenatal Mothers Attending At Institute of Obstetrics and Gynecology, Egmore, Chennai-08" has been validated by the undersigned. The suggestions and modifications given by me will be incorporated by the investigator in concern with their respective guide. Then she can proceed to do the research.

Signature with seal

Name: DHANALANJHME.V Designation: READER College: APROLLO COLUZUE OF NURLING



Place: CHENNAL Date: 9/11/16

CERTIFICATE FOR ENGLISH EDITING

TO WHOM SO EVER IT MAY CONCERN

This is to certify that the dissertation topic "A Pre-Experimental study to Evaluate the Effectiveness of Structured Teaching Programme on Knowledge Regarding Early Identification of High risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08" done by Mrs.J.Dhanalakshmi, M.Sc Nursing II year, college of Nursing, Madras Medical College, Chennai-03, has been edited for English language appropriateness.

Place: Chennai Date: 31.07.2017

M. Dhan lat-L Signature

M. DHANALAKSHMI, M.A., M Phil Asst. Professor Dept. of English Madras Medical College Chennal - 600 003

CERTIFICATE OF TAMIL EDITING

TO WHOM SO EVER IT MAY CONCERN

This is to certify that the dissertation topic "A Pre-Experimental study to Evaluate the Effectiveness of Structured Teaching Programme on Knowledge Regarding Early Identification of High risk Pregnancy Among Antenatal mothers attending at Institute of Obstetrics and Gynaecology, Chennai-08" done by Ms.J.Dhanalakshmi, M.Sc Nursing II year, college of Nursing, Madras Medical College, Chennai-03, has been edited for Tamil language appropriateness.

Place: Malaireddiyur Date ! - 7.7.2017

Date :

Signature Signature A. BABOND, M.A., B.Ed UPLANS (SIDIS)

அரசு உயர்நிலைப்பள்ளி மலைரெட்டியுர்-635 701, வே. மா







