

Original Research Article

# Cardiopulmonary and hematological parameters in pregnancy


Nagadeepa Wulsa<sup>1\*</sup>, Geeta Soren<sup>2</sup>, Rama Mohan Pathapati<sup>3</sup>,  
Madhavulu Buchineni<sup>3</sup>

<sup>1</sup>Assistant Professor, Department of Physiology, MNR Medical College, Sangareddy, Telangana, India

<sup>2</sup>Associate Professor, Department of Pharmacology, Maheshwara Medical College, Patancheru, Telangana, India

<sup>3</sup>Associate Professor, Department of Pharmacology, Narayana Medical College and Hospital, Nellore, Andhra Pradesh, India

\*Corresponding author email: [drdeepamesh6@gmail.com](mailto:drdeepamesh6@gmail.com)

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## Abstract

**Introduction:** We have studied cardiopulmonary and hematological parameters in pregnancy, such as electrocardiographic parameters, peak expiratory flow rates and packed cell volumes in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters of pregnancy.

**Material and methods:** 50 consecutive normal healthy pregnant women in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester each, between 20- 30 years and 50 normal non-pregnant women of the same age group were enrolled. All subjects underwent onetime ECG, pulmonary function tests and complete hemogram.

**Results:** Heart rate statistically significant increased in pregnant women in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester when compared to controls ( $p < 0.01$ ). PR interval showed statistically significant decrease in pregnant women in 2<sup>nd</sup> trimester and 3<sup>rd</sup> trimester when compared to controls ( $p < 0.01$ ). QTc showed statistically significant increase in pregnant women all the trimesters when compared to controls with ( $p < 0.01$ ). QRS frontal axis showed statistically significant decrease in pregnant women of 1<sup>st</sup> trimester, 2<sup>nd</sup> trimester ( $p < 0.01$ ) and 3<sup>rd</sup> trimester ( $p < 0.01$ ) when compared to controls. PEFR values showed statistically highly significant decrease in pregnant women of 1<sup>st</sup> trimester, 2<sup>nd</sup> trimester and 3<sup>rd</sup> trimester when compared to controls with ( $p < 0.001$ ). PCV values showed statistically significant decrease in pregnant women of 1<sup>st</sup> Trimester, 2<sup>nd</sup> trimester and 3<sup>rd</sup> trimester when compared to controls with ( $p < 0.01$ ).

**Conclusion:** We found a statistically significantly increase in the HR QTc, PEFr, PCV and a decrease in PR interval in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters of pregnancy as compared to controls. Additionally, QRS frontal axis was also deviated to left side with advancing pregnancy.

## Key words

Electrocardiography, Pulmonary function, Hematology, Trimesters, Pregnancy.

## Introduction

Normal pregnancy is characterized by profound changes in almost every organ system to ensure the fetus grows properly and receives adequate nutrition. Altered physiological findings and such findings can be misinterpreted as pathological changes in reference to pre-pregnancy standards [1, 2]. Additionally, assessment of these adaptations in normal women during pregnancy is also necessary for better antenatal care, in the assessment of fitness for surgery and to know the progress of pre-existing disease [3-7]. To this purpose we studied cardiopulmonary and hematological parameters in pregnancy, such as electrocardiographic parameters, peak expiratory flow rates and packed cell volumes in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters of pregnancy.

## Materials and methods

The study was conducted in the Physiology department, Kakatiya Medical College, Warangal. Ethical committee has approved the study protocol. 50 consecutive normal healthy pregnant women in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester each, between 20- 30 years and 50 normal non-pregnant women of the same age group were enrolled. Following an explanation about the nature and purpose of the study, those subjects who gave written informed were enrolled. Normal healthy pregnant women in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester between 20-30 years of age were included. Subjects were excluded if they were less than 20 and more than 30 years and / or with any organic cardiac disease, renal disease, severe anemia, thyroid disease, diabetes and hypertension. Relevant information which includes demographic and clinical data from each individual was recorded in a pretested structured proforma. Then these subjects

underwent a detailed physical and systemic examination. Physical examination includes measuring height in centimeters, weight in kilograms, and recording of resting pulse rate by palpating the radial artery and blood pressure recording with a mercury sphygmomanometer using the appropriate sized cuff. Clinical examination of the cardiovascular system and respiratory system was done in detail. Data acquisition was performed in the morning. ECG was recorded using "Digital Physiograph the Lab Tutor". It is an HTML-based software package, designed specifically for laboratory teaching and used in conjunction with ADInstruments' Power Lab. The recorded ECG was evaluated for different parameters such as heart rate, PR interval, QT, QTc interval and QRS axis deviation. Peak expiratory flow was monitored using small hand held apparatus with a mouthpiece at one end and a scale with a moveable indicator. This test was performed three times with Wright peak flow meter (Clement Clarke) and the highest value of PEFr from three accurately performed blows was measured. The PCV can be measured by centrifuging heparinized blood in a capillary tube at 10,000 rpm for five minutes. This separates the blood into layers. The volume of packed red blood cells divided by the total volume of the blood sample gives the PCV. Since a tube is used, this can be calculated by the lengths of the layers.

## Statistical analysis

Results are analyzed with the help of Graph pad calculator and expressed as Mean  $\pm$  SD for continuous data and number and percentages for categorical data. One way ANOVA was used for multiple group comparisons followed by 'Post – hoc – Tukey' test for group – wise comparisons.

Categorical data was analyzed by Chi – square test. A two tailed p value less than 0.05 was considered statistically significant.

## Results

The results were as per **Table - 1**. Heart rate showed statistically significant increase in pregnant women in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester when compared to controls (p <0.01). There was also statistically significant increase in heart rate in pregnant women in third trimester when compared to pregnant women in second trimester and first trimester (p<0.01). PR interval showed statistically significant decrease in pregnant women in 2<sup>nd</sup> trimester and 3<sup>rd</sup> trimester when compared to controls (p < 0.01). There was also statistically significant decrease in PR interval in pregnant women in 3<sup>rd</sup> trimester when compared to pregnant women in 2<sup>nd</sup> trimester and 3<sup>rd</sup> Trimester (p <0.01). There was slight increase in QT interval in pregnant women in third trimester, there was no statistical significance when values of controls, pregnant women in 1st trimester, 2nd trimester and pregnant women in 3rd trimester were compared (p > 0.05). QTc showed statistically significant increase in pregnant women Of 1<sup>st</sup> trimester, 2nd trimester and 3rd

trimester when compared to controls with (p <0.01). There was also statistically significant increase in QTc interval in pregnant women in 3rd trimester when compared to pregnant women in 2nd trimester and 1st trimester. (p < 0.05). QRS frontal axis showed statistically significant decrease in pregnant women of 1<sup>st</sup> trimester, 2<sup>nd</sup> trimester ( p < 0.01 ) and 3<sup>rd</sup> trimester ( p < 0.01 ) when compared to controls. There was no statistically significant difference between pregnant women in 1st trimester, 2nd trimester and pregnant women in 3<sup>rd</sup> trimester (p > 0.05). PEFR values showed statistically highly significant decrease in pregnant women of 1st Trimester, 2nd trimester and 3rd trimester when compared to controls with (p <0.001). There was also statistically significant decrease in PEFR values of pregnant women in 3rd trimester when compared to pregnant women in 2nd trimester and 1st trimester. (p < 0.01). PCV values showed statistically significant decrease in pregnant women Of 1<sup>st</sup> Trimester, 2<sup>nd</sup> trimester and 3<sup>rd</sup> trimester when compared to controls with (p <0.01). There was also statistically significant decrease in PCV values of pregnant women in 3rd trimester when compared to pregnant women in 2<sup>nd</sup> trimester and 1<sup>st</sup> trimester. (p < 0.01).

	Controls [A]	1 <sup>st</sup> TM [B]	2 <sup>nd</sup> TM [C]	3 <sup>rd</sup> TM [D]	Differences between groups				
					A-B	A-C	A-D	B-C	B-D
<b>Age (years)</b>	24.7 ± 2.2	24.1 ± 2.4	24.3 ± 2.6	25.5 ± 2.8	n	n	n	n	n
<b>HR (b/m)</b>	79.0 ± 6.60	84.2 ± 6.40	87.2 ± 6.16	94.2 ± 8.5	**	**	**	**	**
<b>PR (sec)</b>	0.15 ± 0.01	0.14 ± 0.02	0.13 ± 0.02	0.12 ± 0.02	**	**	**	**	**
<b>QT (sec)</b>	0.35 ± 0.02	0.35 ± 0.01	0.35 ± 0.01	0.36 ± 0.01	n	n	n	n	n
<b>QT<sub>c</sub></b>	0.38 ± 0.01	0.39 ± 0.01	0.40 ± 0.01	0.41 ± 0.01	**	**	**	**	**
<b>QRS (degrees)</b>	64.10 ± 6.8	57.0 ± 12.5	53.0 ± 16.7	45.9 ± 22.1	**	**	**	N	N
<b>PEFR (Lit/min)</b>	331.46 ± 40.84	297.52 ± 32.81	234.77 ± 34.44	183.81 ± 33.90	**	**	**	**	**
<b>PCV %</b>	38 ± 2	33 ± 2	31 ± 2	29 ± 3	*	*	*	*	*

\*=p<0.05, \*\*= p<0.01, n=P>0.05

## Discussion

Pregnancy is associated with remarkable changes in the cardiovascular system. The ECG during normal pregnancy may show wide variation from the normal limits even in the absence of demonstrable heart disease. This can be due to the altered spatial arrangement of the chest organs, and electrical properties of the myocardium are due to change in sympathetic and hormonal modulation of the electrical activity of the heart during pregnancy. It is also interesting to note that pregnancy may be associated with a concentric enlargement of the left ventricle in response to the hemodynamic requirements, which can explain these ECG changes [8-10].

In this study, we found linear increase in HR across 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimester pregnant women who were statistically significant as compared to controls as well as between trimesters. Pregnancy is associated with dramatic cardiovascular adaptations like increase in HR. This increase in HR is linked to autonomic nervous system changes that produce alterations in cardiac autonomic modulation such an association was seen in early pregnancy where the initial change in HR may be linked to the production of chorionic gonadotropin, with the later gradual increase being related to the vascular changes which accompany placental and fetal growth [11]. This increased in heart rate during third trimester compensates for the fall in stroke volume resulting from caval compression [12]. Additionally, the cardiac output increases as early as 5 weeks and rises to 45% above the baseline at 24 weeks of gestation.

We also observed that PR interval significantly decreased in pregnant women of 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters when compared to controls. Furthermore, significant decrease in PR interval was also observed in pregnant women in 3<sup>rd</sup> trimester when compared to pregnant women in 2<sup>nd</sup> trimester and 1<sup>st</sup> trimesters. The decrease in PR interval during pregnancy could be due to shortening of AV conductance with respect to increase in heart rate that accompany during

pregnancy [13-14]. QT Interval a marker for ventricular depolarization and repolarization was in pregnant women during 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters when compared to controls. However when corrected to heart rate QTc showed a significant increase in the pregnant women of 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters when compared to controls. It is possible that the increase in QTC may be due to increase in HR. This could be linked to changes in ventricular depolarization and repolarization patterns during pregnancy and they must be considered as a complex consequence with changes in regulatory mechanisms during normal pregnancy [15-17]. It can be concluded that prolonged QTc is simply just as a non specific sign of changed course of repolarization [8, 18]. In pregnancy, the change in the electrical axis can be raising diaphragm as pregnancy advances, left shift of apical impulse of axis due to changes in the left ventricular size and mass with associated increased volume, elevation and rotation of the heart, resulting from the enlarging uterus [8-10, 13-14], Similarly we also observed, a significant decrease in QRS frontal axis in pregnant women of 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters when compared to controls.

There was a decline in mean PEFR as the pregnancy advances from 1<sup>st</sup> to 3<sup>rd</sup> trimester and this decline was significant when compared to controls as well as between trimesters. Similar observation was noted in the earlier studies on PEFR in pregnancy [19, 20, 21-31]. The decrease in PEFR may be attributed to upward displacement of the diaphragm, increased metabolic rate, changes in the mechanics of breathing, and increases in progesterone level [32, 33].

This study has shown that in pregnancy hemorrheologic activity differs significantly due to increased values of PCV as compared to non-pregnant subjects. Likewise, there was also statistically significant increase in PCV in pregnant women in 3<sup>rd</sup> trimester when compared to pregnant women in 2<sup>nd</sup> trimester and 1<sup>st</sup> trimester. This increase in PCV could be due to marked increase in plasma volume associated

with normal pregnancy causing dilution of many circulating factors and cells resulting in physiological anemia [34-37].

## Conclusion

We found a statistically significantly increase in the HR QTc, PEFr, PCV and a decrease in PR interval in 1<sup>st</sup>, 2<sup>nd</sup> and 3<sup>rd</sup> trimesters of pregnancy as compared to controls. Additionally, QRS frontal axis was also deviated to left side with advancing pregnancy. Our study provides a quick look into the variety of adaptations in cardiopulmonary and hematological systems during normal pregnancy which brings about changes in even in the absence of any disease.

## References

1. Oakley C, Warnes CA. Heart disease in pregnancy: Wiley Online Library; 2007.
2. Hall JE. Guyton and Hall textbook of medical physiology: Elsevier Health Sciences; 2010.
3. Moore LE, Pereira N. Physiological changes of pregnancy. Maternal Critical Care: A Multidisciplinary Approach, 2013; 107.
4. Campbell DC, editor Physiological changes of pregnancy. Seminars in Anesthesia, Perioperative Medicine and Pain; Elsevier, 2000.
5. Carlin A, Alfirevic Z. Physiological changes of pregnancy and monitoring. Best Practice & Research Clinical Obstetrics & Gynaecology, 2008; 22(5): 801-23.
6. Horan C. Physiological changes in pregnancy. Obstetric Evidence Based Guidelines, 2007; 17.
7. May WY. Physiological changes in pregnancy. Essential of Obstetrics, 2004.
8. Lechmanova M, Kittnar O, Mlcek M, Slavicek J, Dohnalova A, Havranek S, et al. QT dispersion and T-loop morphology in late pregnancy and after delivery. Physiological research, 2002; 51(2): 121-30.
9. Sunitha M, Chandrasekharappa S, Brid S. Electrocardiographic Changes in Normal Pregnancy. International Journal of Physiology, 2014; 2(2): 47-52.
10. Hoogsteder PH, KRÜSE AJ, Sep SJ, Dassen WR, Gorgels AP, Peeters LL. Electrocardiographic findings in women with a recent history of pre-eclampsia. Acta obstetrica et gynecologica Scandinavica, 2012; 91(3): 372-8.
11. Clapp JF. Maternal heart rate in pregnancy. American journal of obstetrics and gynecology, 1985; 152(6): 659-60.
12. Julian D, Wenger N. Heart disease and heart surgery during pregnancy. women and heart disease United Kingdom: Martin Dunitz, 2000.
13. Sunitha M, Chandrasekharappa S. Electrocardiographic QRS Axis, Q Wave and T-wave Changes in 2<sup>nd</sup> and 3<sup>rd</sup> trimester of Normal Pregnancy. Journal of clinical and diagnostic research JCDR, 2014; 8(9): BC17.
14. Madras V, Challa N. Electrocardiographic variations during three trimesters of normal pregnancy. International Journal of Research in Medical Sciences, 2015; 3(9): 2218-22.
15. Carruth JE, Mirvis SB, Brogan DR, Wenger NK. The electrocardiogram in normal pregnancy. American heart journal, 1981; 102(6): 1075-8.
16. Ozmen N, Cebeci B, Yiginer O, Muhcu M, Kardesoglu E, Dincturk M. P-wave dispersion is increased in pregnancy due to shortening of minimum duration of P: does this have clinical significance? Journal of international medical research, 2006; 34(5): 468-74.
17. Ozmen N, Cebeci BS, Kardesoglu E, Celik T, Dincturk M, Demiralp E. P wave dispersion is increased in pulmonary stenosis. Indian pacing and electrophysiology journal, 2006; 6(1): 25.
18. Chia P, Chia H, Subramaniam R. A clinical approach to heart disease in pregnancy. The Obstetrician &



- Gynaecologist., 2002; 4(3): 162-8.
19. Phatak MS, Kurhade G. Short communication a longitudinal study of antenatal changes in lung function tests and importance of postpartum exercises in their recovery. *Indian J Physiol Pharmacol.*, 2003; 47(3): 352-6.
  20. Mokkapatti R, Prasad E, Fatima K. Ventilatory functions in pregnancy. *Indian journal of physiology and pharmacology*, 1991; 35(4): 237-40.
  21. Bansal M, Goyal M, Dhillon JK, Kaur P. Longitudinal study of peak expiratory flow rate in pregnant women. *NJIRM*, 2012; 3(1): 34-8.
  22. Priyadarshini G, Mishra A. Effect of gestational age on pulmonary functions in pregnant Odia women. *International Journal of Medical Science and Public Health*, 2014; 3(11): 1364-7.
  23. Monga U, Kumari K. Pulmonary functions in Punjabi pregnant women. *Indian J Physiol Pharmacol.*, 2000; 44(1): 115-6.
  24. Puranik BM, Kurhade GA, Kaore SB, Patwardhan SA, Kher JR. PEFr in pregnancy: a longitudinal study. *Indian J Physiol Pharmacol.*, 1995; 39(2): 135-9.
  25. Harirah HM. Effect of gestational age and position on peak expiratory flow rate: a longitudinal study. *Obstetrics & Gynecology*, 2005; 106(1): 199-200.
  26. Grindheim G, Toska K, Estensen ME, Rosseland L. Changes in pulmonary function during pregnancy: a longitudinal cohort study. *BJOG: An International Journal of Obstetrics & Gynaecology*, 2012; 119(1): 94-101.
  27. Yerneni S, Sajja S. Study of FEV1, VC and PEFr in different trimesters of pregnancy. *Int J Res Health Sci.*, 2014; 2(1): 41-6.
  28. Rasheed B, Hussain K, Hussain S. PEFr in relation to phases of pregnancy. *Indian journal of physiology and pharmacology*, 2000; 44(4): 511-2.
  29. Rasheed B, Mansoor A, Hussain S. Incentive spirometry and PEFr in different phases of pregnancy. *Indian journal of physiology and pharmacology*, 2002; 46(1): 126.
  30. Sroczyński T, editor [Evaluation of respiratory tract function in healthy women in the last month of uncomplicated pregnancy]. *Annales Academiae Medicae Stetinensis*; 2001.
  31. Sodhi C, Pramod J, Singh J, Kaur V. Effect of advanced uncomplicated pregnancy on pulmonary function parameters of north Indian subjects. *Indian J of Physiol Pharmacol*, 2010; 54(1): 69-72.
  32. Bobrowski RA. Pulmonary physiology in pregnancy. *Clinical obstetrics and gynecology*, 2010; 53(2): 285-300.
  33. Chesnutt AN. Physiology of normal pregnancy. *Critical care clinics.*, 2004; 20(4): 609-15.
  34. Imoru M, Emeribe A. Haemorrhologic profiles in apparently healthy pregnant women in Calabar, Nigeria. *African Journal of Biotechnology*, 2008; 7(24).
  35. Imoru M, Emeribe A. Haemorrhologic and fibrinolytic activities in pregnant women: Influence of gestational age and parity. *African Journal of Biotechnology*, 2009; 8(23).
  36. Oke OT, Awofadeju SO, Oyedeji SO. Haemorrhological profiles in different trimesters among pregnant women in South West Nigeria. *Pak J Physiol.*, 2011; 7(2): 17-19.
  37. Salawu L, Durosinmi M. Erythrocyte rate and plasma viscosity in health and disease. *Nigerian journal of medicine: journal of the National Association of Resident Doctors of Nigeria*, 2000; 10(1): 11-3.