A STUDY OF FVC, PEFR AND MEP IN DIFFERENT TRIMESTERS OF PREGNANCY

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Abstract

Objective: Pregnancy is characterized by profound changes in the function of virtually every regulatory system in the human body. The events in pregnancy elicit one of the best examples of selective anatomical, physiological & biochemical adaptation that occurs during pregnancy & profound changes in respiratory physiology is a part of the same process. Thus this study was designed to evaluate the pulmonary function tests in 1^{st} , 2^{nd} and 3^{rd} trimesters of pregnancy & compare them with non-pregnant control group.

Method: A cross-sectional study was carried in 200 healthy women in the age range of 19-35 years .The subjects were distributed in four groups, i.e control (non-pregnant) group and 1^{st} , 2^{nd} & 3^{rd} trimester pregnant groups. Number of subjects in each group is 50. We recorded respiratory parameters in control and study groups. Statistical analysis done by 'Z' test.

Result: There was significant decrease in FVC, PEFR & MEP in all trimesters of pregnancy with maximum decrease of FVC in 1st trimester & PEFR, MEP in 2nd trimester.

Conclusion: The changes in pulmonary function are attributed to major adaptations in the maternal respiratory system & are also be influenced by the mechanical pressure of enlarging gravid uterus, elevating the diaphragm & restricting the movements of lungs thus hampering the forceful expiration & decrease in 1^{st} trimester might be due to decline in alveolar Pco₂ caused by hyperventilation which acts as bronchoconstrictor & due to sensitization of respiratory centre due to progesterone.

Keywords: Pregnancy, FVC, PEFR, MEP.

1. Introduction

Pregnancy is characterized by profound changes in the function of virtually every regulatory system in the human body. The events in pregnancy elicit one of the best examples of selective anatomical. physiological & biochemical adaptation that occur during pregnancy & profound changes in respiratory physiology is a part of the same process¹. The changes in the respiratory physiology² are due to increasing size of the fetus with advancing gestation which constitutes a mechanical impediment to normal process of ventilation³& due to hormone Progesterone which increases ventilation by increasing respiratory centre sensitivity to carbondioxide as a result the tidal volume and minute ventilation is increased ⁴⁻⁶.

The physiological adaptation of the pregnant woman involves the circulatory, respiratory, digestive, renal, endocrine & metabolic systems. Their precise knowledge allows the clinician to verify the extent of the adaptation in pregnant women & helps to avoid unnecessary treatment of physiology changes misinterpreted as pathological changes in reference to pre pregnancy standards⁷.

The knowledge of the expected or desired changes in pulmonary parameters is fundamental in understanding of how the disease states affect pregnancy & vice versa⁸. Also, information regarding status of pulmonary function is essential for assessment of fitness for anaesthesia⁹.

The serial testing initiated early in pregnancy permits valid interpretation of pulmonary function changes with advance in gestation. The respiratory changes are adaptive in nature. In order to evaluate any respiratory ailment during pregnancy, an accurate knowledge of the physiological changes in pulmonary function during normal pregnancy is necessary. The changes in maternal pulmonary function tests during pregnancy have been reported.¹⁰

Aim of this study was to evaluate the changes in the pulmonary functions of women in the age group of 19-35 yrs at 1st, 2nd& 3rd trimesters of pregnancy & compare them with that of healthy non-pregnant age matched controls.

A cross sectional study was conducted in the Department of Physiology, Shri B.M.Patil Medical College, Hospital and Research Centre, Bijapur.

The study was undertaken to determine the pulmonary function changes in 1^{st} , 2^{nd} & 3^{rd} trimesters of pregnancy. The observations were compared with age matched healthy non pregnant women

2.1 Method of Collection of data:

Study Group: 150 pregnant women in the age group of 19-35yrs who were attending the OPD of OBGy of Shri B.M. Patil Medical College were included in the study group. The study group was in turn divided into 3 subgroups. Each sub group was comprised of 50 women in first, second and third trimesters of pregnancy.

Control Group: It comprised of apparently healthy age matched 50 non pregnant women.

The nature and purpose of the study were explained to the subjects who had volunteered for the study. From each participant an informed consent was obtained. A proforma was used to record the relevant information from each selected individual who had fulfilled inclusion criteria. The subjects who had exclusion criteria were dropped from the study. A thorough physical & systemic examination of each subject was done (in particular, cardiovascular and respiratory system). Recordings were taken during morning hours between 9 am to 12 Noon. **Inclusion Criteria:** Apparently healthy subjects of Indian origin were included in the study. The apparent health status of the subject was determined through thorough clinical examination and history taking.

Exclusion Criteria : Subjects with acute respiratory infection in the previous three months, chronic respiratory infection including asthma, history or clinical signs of cardiovascular diseases, diabetes mellitus, hypertension, tobacco consumption in any form , alcohol intake, endocrine disorders, obesity & moderate to severe anemia were excluded from the studies.

Following parameters were recorded in each subject:

A .Record of physical Anthropometry of subjects: Height (in centimeters), Weight (in kilograms), Body surface area & Body Mass Index were measured.

B. Record of physiological parameters: Pulse Rate (beats per minute), Blood Pressure (SBP and DBP in mmhg), Respiratory Rate (cycles /minute) were recorded.

C. Record of pulmonary function parameters: The subject was informed about the procedure. Consent was taken from each subject before recording .For each test, three readings were taken. The highest reading of the three was taken for calculation. All tests were recorded in a sitting posture at room temperature, in morning hours.

The following pulmonary parameters are recorded by Computerized Spiropac (Medicad)¹¹ I) 1. FVC (Forced Vital Capacity in L)

2. PEFR (Peak Expiratory Flow Rate in L/sec)

II). MEP (Maximum Expiratory Pressure in mmHg) - Recorded by using modified Black's apparatus.

2.2 Statistical analysis: The results were expressed as Mean \pm SD. Z test was used for comparison between control and study groups in consultation with statistician. A 'p' value of 0.05 or less was considered as statistically significant.

3. Results

3.1 Forced Vital Capacity (FVC) in Litres: There was statistically very highly significant decrease in FVC in 1^{st} (p<0.001), 2^{nd} (p<0.001) & 3^{rd} (p<0.001) trimesters of pregnancy when compared to control group. However, there was statistically significant increase in FVC of 3^{rd} trimester (p<0.01) as compared to 2^{nd} trimester of pregnancy.

3.2 Peak Expiratory Flow Rate (PEFR) in Liters/sec: There was statistically very highly significant decrease in PEFR in 1^{st} (p<0.001), 2^{nd} (p<0.001) & 3^{rd} (p<0.001) trimesters of pregnancy when compared to non pregnant women. Maximum decrease was in 1^{st} trimester. **3.3 Maximum Expiratory Pressure in mmHg:** There was statistically significant decrease in MEP in 1^{st} (p<0.041), 2^{nd} (p<0.004) & 3^{rd} trimesters when compared to control group. Maximum decrease in MEP was seen in 2^{nd} trimester of pregnancy.

| subjects in Control and Study Groups | | | | | | | |
|--------------------------------------|--------------------|---------------------------|----------|---------------------------|----------|---------------------------|----------|
| Parameters | Control | 1 ST Trimester | | 2 ND Trimester | | 3 RD Trimester | |
| | Mean <u>+</u> SD | Mean + SD | P value | Mean <u>+</u> SD | P value | Mean <u>+</u> SD | P value |
| Age (yrs) | 24 <u>+</u> 5.8 | 25 <u>+</u> 4.4 | 0.183 | 25 <u>+</u> 0.36 | 0.244 | 26 <u>+</u> 3.4 | 0.032* |
| Height (cms) | 159 <u>+</u> 5.06 | 151 <u>+</u> 16 | 0.001*** | 155 <u>+</u> 5.2 | 0.001*** | 155 <u>+</u> 5.3 | 0.001*** |
| Weight (kg) | 59 <u>+</u> 9.6 | 52 <u>+</u> 7.1 | 0.001*** | 55 <u>+</u> 7.1 | 0.007** | 60 <u>+</u> 9.2 | 0.342 |
| BMI (kg/m^2) | 23.19 <u>+</u> 3.5 | 21.93 <u>+</u> 3.86 | 0.045* | 22.4 <u>+</u> 3.79 | 0.139 | 24.73 <u>+</u> 3.08 | 0.010** |
| BSA (Sq m) | 1.61 <u>+</u> 0.14 | 1.49 <u>+</u> 0.13 | 0.001*** | 1.5 <u>+</u> 0.1 | 0.001*** | 1.58 <u>+</u> 0.13 | 0.091 |

Table 1: Mean + SD and level of significance of Age and Anthropometric Parameters of subjects in Control and Study Groups

p>0.05: Not Significant, *p: <0.05: Significant, ** p: <0.01: Highly significant, *** p: <0.001: Very highly significant

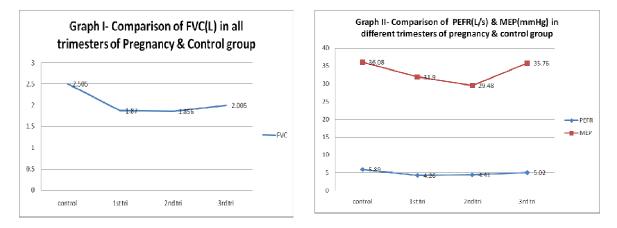
| Table 2: Mean + SD and Range of Physiological Parameters of subjects in Control and Study |
|---|
| groups |

| groups | | | | | | |
|---------------|---------------------|-------------------------------------|--------------------------------------|----------------------------------|--|--|
| Parameters | Control | 1 ST Trimester | 2 nd Trimester | 3 rd Trimester | | |
| | Mean <u>+</u> SD | Mean <u>+</u> SD | Mean <u>+</u> SD | Mean <u>+</u> SD | | |
| PR(beats/min) | 76 <u>+</u> 9.4 | 89 <u>+</u> 10 (p=0.054*) | 77 <u>+</u> 5.5 (p=0.167) | 78 <u>+</u> 7.7 (p=0.053*) | | |
| SBP (mm Hg) | 118.66 <u>+</u> 4.4 | 115.60 <u>+</u> 7.67 (p=0.0146*) | 109.56 <u>+</u> 5.68 (p=0.001***) | 117.56 <u>+</u> 7.94 (p=0.17) | | |
| DBP (mm Hg) | 74.32 <u>+</u> 4.75 | 73.36 <u>+</u> 5.97 (p=0.378) | 66.52 <u>+</u> 5.68 (p=0.001***) | 75.60 <u>+</u> 7.41 (p=0.441) | | |
| PP (mm Hg) | 44.34 <u>+</u> 5.47 | 42.44 <u>+</u> 7.81 | 43.20 <u>+</u> 5.33 | 42.44 <u>+</u> 8.33 | | |
| MAP (mm Hg) | 88.94 <u>+</u> 5.72 | 88.32 <u>+</u> 5.80 | 80.90 <u>+</u> 4.19 (p=0.001) | 89.45 <u>+</u> 7.26 | | |
| RR(cpm) | 16 <u>+</u> 3 | 22 <u>+</u> 3 | 22 <u>+</u> 4.3 | 26 <u>+</u> 3.7 | | |

p>0.05: Not Significant, *p: <0.05: Significant, ** p: <0.01: Highly significant, *** p: <0.001: Very highly significant

| Denometers | Control | 1 ST Trimester | 2 ND Trimester | 3 RD Trimester | |
|------------|---------------------|---------------------------|---------------------------|---------------------------|--|
| Parameters | Mean <u>+</u> SD | Mean <u>+</u> SD | Mean <u>+</u> SD | Mean <u>+</u> SD | |
| FVC(L) | 2.50 ± 0.20 | 1.87 <u>+</u> 0.37 | 1.85 <u>+</u> 0.39 | 2.19 <u>+</u> | |
| | 2.50 <u>+</u> 0.39 | (p=0.001***) | (p=0.001***) | 2.0(p=0.001***) | |
| PEFR(L/s) | 5 90 1 1 16 | 4.26 <u>+</u> 1.28 | 4.41 <u>+</u> 1.35 | 5.02 <u>+</u> | |
| | 5.89 <u>+</u> 1.16 | (p=0.001***) | (p=0.001***) | 1.26(p=0.001***) | |
| MEP(mmHg) | 36.08+14.13 | 31.9 <u>+</u> 9.35 | 29.48 <u>+</u> 10.33 | 35.76 <u>+</u> | |
| | <u>50.06+</u> 14.15 | (p=0.041*) | (p=0.004**) | 12.6(p=0.458) | |

p>0.05: Not Significant, *p: <0.05: Significant, ** p: <0.01: Highly significant, *** p: <0.001: Very highly significant



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4. Discussion

A cross-sectional study was carried in 200 healthy women in the age range of 19-35 years. The subjects were distributed in four groups, i.e control (non-pregnant) group and 1st, 2nd &3rd trimester pregnant groups. Number of subjects in each group is 50. We have recorded various physical, physiological & respiratory parameters in control and study groups.

4.1 Forced vital capacity: Our study showed significant decrease in FVC from 1st trimester to 3rd trimester as compared to control which is in agreement with other workers. The decrease in FVC was maximum in first trimester which may be attributable to hormonal changes which requires further studies.

A study of forced vital capacity in pregnant women by Dipok Kumar Sunyal and workers showed reduced FVC in all trimesters as compared to control & maximum decrease in third trimester. The decrease in FVC is attributable to the mechanical pressure of enlarging gravid uterus, elevating the diaphragm & restricting the movements of lungs thus hampering the forceful expiration¹².

A study by Deepal & workers showed no significant changes in FVC during all trimesters of pregnancy Hormonal alteration in pregnancy causes a reduction in the tracheo-bronchial smooth muscle tone & the increasing thoracic width may be due to enlarging uterus as a result there is no impairment in large airway function throughout pregnancy¹³.

4.2 PEFR: Our study showed significant decrease in PEFR from 1^{st} trimester to 3^{rd} trimester as compared to control & maximum decrease was seen in 1^{st} trimester. The maximum decrease in 1^{st} trimester may be attributable to the hormonal levels.

As per studies on PEFR by Neeraj & workers, there was decrease in PEFR in third trimester & decrease was attributable to the decline in alveolar Pco_2 which acts as bronchoconstrictor. Also the decrease in PEFR could be due to lesser force of contraction of main expiratory muscles like the anterior abdominal wall muscles & internal intercostals muscles¹⁴.

A work on PEFR by Sunyal DK et al there was decrease in PEFR in all trimesters of pregnancy which is significant in 2nd & 3rd trimesters of pregnancy. The cause for the decrease was more likely due to lesser force of contraction of the expiratory muscles like anterior abdominal muscles & internal intercostals muscles in this state ^{15,16}. Moreover, progressively reduced value of PEFR in three trimesters of pregnancy

may be attributed to the mechanical effects of enlarged gravid uterus reducing vertical dimension by limiting movement of diaphragm¹⁷. In addition some degree of obstruction to the expiratory flow, especially late in pregnancy also contributes¹⁸. Some studies suggest, inadequate nutrition due to morning sickness, altered eating habits associated with advancing gestation that resulted in muscular weakness & the lesser force of contraction of main expiratory muscles¹⁹.

A study by Leo R. Brancazio & workers showed that PEFR does not change with pregnancy. Not only are the absolute mean peak expiratory flow rates similar at all four times, but the mean normalized peak expiratory flow rates (calculated by using formula PEFR (L/min) = 198.07 + 3.07 age - 0.0477 age² + 3.6 height) in all groups are close to unity ¹⁹.

4.3 MEP: Our study showed significant decrease in MEP from 1^{st} trimester to 3^{rd} trimester as compared to control with maximum decrease in 2^{nd} trimester.

Conclusion

The significant decrease in FVC & PEFR might be due to the mechanical pressure of enlarging gravid uterus, elevating the diaphragm & restricting the movements of lungs thus hampering the forceful expiration and maximum decrease in 1^{st} trimester due to decline in alveolar Pco₂ caused by hyperventilation which acts as bronchoconstrictor.

The statistical significant decrease in PEFR & MEP in all trimesters might be due to decrease in expiratory muscle power.

To establish the cause of decrease in respiratory parameters more in first trimester of pregnancy than in 2^{nd} & 3^{rd} , further studies are to be undertaken by hormonal assay in different trimesters to know the relation between hormone and respiratory parameter

References

- Chhabra S, Nangia V, Ingley KN. Changes in respiratory function tests during pregnancy. *Ind J Physiol Pharmacol* 1988; 32: 56-60
- 2. Pandya KD, Chandwani S, Desai CA, Dadlani AG. Study of vital capacity and timed vital capacity in normal non-pregnant and pregnant women. *J Obst Gynecol Ind*, 1984: 36 : 1053-1057
- Saxena SC, Rao VSC, Mudgal SA. Study of pulmonary function tests during pregnancy. *J Obst Gynecol* 1979; 29 : 993-995

- 4. Bayliss DA, Millhorn DE. Central neural mechanics of progesterone action: application to the respiratory system. *J Appl Physiol* 1992; 73(2):393-404
- 5. Skatrud JB, Dempsey A, Kaiser DG. Ventilatory response to medroxyprogesterone acetate in normal subjects: time course and mechanism. *J Appl Physiol* 1978; 44(6): 939-944
- 6. Lysons HA, Antonio R. The sensitivity of the respiratory centre in pregnancy and after the administration of progesterone. *Trans Assoc Am Phys* 1959; 72: 173-180
- Foidart M. Physiology of the pregnant woman and risk factors. *Contracept fertile* sex 1993; 21: 811-815
- 8. Elkus R, Popovich J Jr. Respiratory physiology in pregnancy. *Clin Chest Med* 1992; 13: 555-565
- Mokkapati R, Prasad EC, Venkatraman, Fatima K. Ventilatory functions in pregnancy. *Indian J Physiol Pharmacol* 1991; 35: 237-240
- Kolarzyk E, Szot WM, Lyszczarz J. Lung function and breathing regulation parameters during pregnancy. *Arch Gynecol Obstet* 2005 Jun; 272(1): 53-8.
- Pal G.K. Textbook of practical physiology 2nd edition Orient Longmann publications Chennai 2005;p:154-161
- Sunyal DK, Md Ruhul Amin, MH Molla, Abida Ahmed, Shameena Begum. Forced vital capacity in normal pregnancy. *J med. Sci. Res* 2007 July; 09: 01: 21-25
- 13. Weerasekara Deepal S., Ruberu D.Kusua and S.Sivayogan Pulmonary Functions in Pregnant Sri Lankan Women 1999 *sabaragauwa university journal*, vol.,2, no., 1, pp. 57-60
- 14. Neeraj , Sodhi Candy , John Pramod, Singh Joydeep & Kaur Vaneet, *Indian J Physiol Pharmacol* 2010 : 54 (1) : 69-72.
- 15. Phatak MS, Kurhade GA. Longitudinal study of antenatal changes in lung function tests and importance of postpartum exercises in their recovery. *Indian journal of physiology pharmacology* 2003;47(3): 352-356
- Spiropoulos K, Prodromaki E, Tsapanos V. Effect of body position on PaO2 and PaCO2 during pregnancy. *Gynecol Obstet Invest* 2004; 58: 22 -25.
- 17. Rasheed MA, Mansoor A, Hussian S. Incentive spirometry and PEFR in different phases of pregnancy. *Indian J physiol pharmacol* 2002; 46 (1):126-128.

- Norregaard O, Schultz P, Ostergaard A and Dahl R. Lung function & Postural changes during pregnancy. *Respir Med.* 1989; 83: 467-70
- 19. Brancazio LR, Laifer SA, Schwartz T. Peak Expiratory Flow Rate in normal pregnancy. *Obstet Gynecol.* 1997; 89 (3): 383-386.