



## Study of FEV<sub>1</sub>, VC and PEF<sub>R</sub> in different trimesters of pregnancy

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### Abstract:

**Background & Objectives:** The aim of the present study was to monitor the changes that occur in pulmonary function tests in the different trimesters of pregnancy as compared with that of control group i.e., non pregnant women. **Materials & Methods:** The study consists of recording the Pulmonary Function Tests of four groups of female subjects including pregnant women of various phases of gestational period i.e., 12 weeks, 24 weeks 36 weeks and control group of non pregnant women using computerized Medspiror. The different static lung function parameters measured in this study were vital Capacity (VC), FEV<sub>1</sub> and Peak Expiratory Flow Rate (PEFR). PEFR was measured with Mini Wright's peak flow meter. **Results:** We observed a statistically significant decrease in PEFR ( $p < 0.0001$ ), a significant increase in VC & no significant change FEV<sub>1</sub> in different trimesters of pregnancy. **Conclusion:** Comparative study of pulmonary function tests on different trimesters of pregnancy showed that different respiratory parameters like PEFR were compromised significantly due to mechanical pressure of gravid uterus, diaphragm restricting the movement of lungs. VC showed a significant increase in different trimester's when compared with the non pregnant state whereas FEV<sub>1</sub> showed no significant change. The respiratory system undergoes physiological and anatomical changes during pregnancy and we observed a significant decrease in PEFR and increase in VC among pregnant women.

**Key words:** Forced Expiratory Volume in 1st sec (FEV<sub>1</sub>); Peak Expiratory Flow Rate (PEFR); Pregnancy; Spirometry; Vital Capacity (VC)

### Introduction

Marked local and systemic changes in maternal physiology are initiated by conception and they continue throughout the pregnancy. Pulmonary function is affected by changes of the airway,

thoracic cage and respiratory drive. There is a significant increase in minute ventilation as a result of a direct stimulatory effect of progesterone on central respiratory drive and an enhancement of the hypercapnic ventilatory drive. Lung compliance does

not change, but total respiratory compliance is decreased at term as a result of a reduction in chest wall compliance. Despite the significant increase in intra-abdominal pressure that is due to the enlarging uterus, the maximal inspiratory and expiratory pressures, as well as maximum trans-diaphragmatic pressure do not change significantly. In pregnancy, the hormonal changes along with progressive increase in abdominal volume may have mechanical impact on respiratory function. However, an increased transverse diameter of the chest, resulting from a widened sub costal angle, opposes the effect of the enlarging pregnant uterus and elevated diaphragm, maintaining an altered pulmonary function during pregnancy [1].

There is no significant change in respiratory muscle strength during pregnancy despite the cephalad displacement of the diaphragm and changes in the configuration of the chest wall. Despite the upward displacement of the diaphragm by the gravid uterus, diaphragm excursion actually increases by 2 cm when compared with the non pregnant state [2,3]. Increased diaphragmatic excursion and preserved respiratory muscle strength are important adaptations, given the increase in tidal volume and minute ventilation that accompanies pregnancy. Improved diaphragm mechanics in pregnancy are explained by an increased area of apposition of the diaphragm to the rib cage [3].

Routine spirometric measurements {forced expiratory volume in 1 second - FEV1} and FEV1/FVC ratio are not significantly different compared with non pregnant values. VC has been reported to be either minimally increased, decreased, or unchanged during pregnancy compared with the non pregnant state; on average, there is no significant change in VC in most of the studies [4-7]. The stability of spirometry during pregnancy suggests that there is no significant change in expiratory airflow resistance with pregnancy. Hormone determined changes in smooth muscle tone and elastance of the connective tissue which occur during pregnancy could possibly alter the mechanical properties of the respiratory system. The increase in progesterone and estrogen associated with pregnancy contribute to vascular and central nervous system effects, changes in the balance of bronchoconstrictor and bronchodilator prostanoids and increase in peptide hormones which alter connective tissue characteristics.

The values obtained by forced spirometry, including forced vital capacity (FVC), forced expiratory volume in 1 second (FEV1), and peak

expiratory flow rate (PEFR) have largely been found to remain unchanged during pregnancy [8-10]. In other studies, PEFR is found to decrease with advancing gestational age and to be affected by maternal positioning [11-12] and by living at high altitude [13].

## Materials and Methods

The study consists of recording the Pulmonary Function Tests of four groups of female subjects including both normal and pregnant women of various phases of gestational period i.e., 12 weeks, 24 weeks & 36 weeks and control group of non pregnant women of the child bearing age. The subjects considered for this study are with Hemoglobin more than 10 gm%. The study was approved by the Institutional Ethical Committee.

All the subjects were called for spirometric tracings in the afternoon between 3 to 5pm. (3 to 4 hrs after meal) in the post absorption stage in order to keep uniform conditions for recording the tests. All the subjects are given instructions and demonstration with regard to performance of the tests. The tracings in the spirograph were taken after being fully satisfied that the subject has understood the procedure of the test. Two to three tracings were taken out of which the best is taken as final reading.

The female subjects who are nonsmokers and free from cardiovascular and respiratory ailments were grouped into four groups as: Group 1 - Female normal subjects aged 20-25 years; Group 2 -Pregnant subjects of first trimester gestational period aged 20-25 years ; Group 3- Pregnant subjects of second trimester gestational period aged 20-25 years ; Group 4 - Pregnant subjects of third trimester gestational period of age 20-25 years .The different lung function parameters measured in this study include FEV<sub>1</sub>, VC & PEFR. PEFR was measured with the Mini Wrights peak flow meter. Statistical Analysis was done using Graphpad prism 6 software and unpaired t test was done to estimate the p value between different groups.

## Results

The mean Vital Capacity in I trimester subjects showed a marginal increase of 3.48% when compared with the control non pregnant subjects (p = 0.0116). Similarly , the mean VC in II & III trimester subjects showed an increase of 3.68 % & 6.51% respectively (p value = 0.0155; p <0.0001 respectively) when compared with the control subjects.

**Table 1: Comparison of Mean Values of PEFR, FEV1 & VC along with p value in different trimesters of pregnancy**

	<b>CONTROL MEAN±SD</b>	<b>1<sup>ST</sup> TRIMESTER MEAN±SD</b>	<b>P VALUE</b>
<b>PEFR</b>	5.904 ± 0.02970	5.645 ± 0.02284	P < 0.0001 ****
<b>FEV1</b>	1.582 ± 0.02698	1.598 ± 0.01748	P= 0.6122
<b>VC</b>	3.011 ± 0.02022	3.116 ± 0.03434	P = 0.0116 *

	<b>CONTROL MEAN±SD</b>	<b>2<sup>ND</sup> TRIMESTER MEAN±SD</b>	<b>P VALUE</b>
<b>PEFR</b>	5.904 ± 0.02970	5.430 ± 0.01611	P < 0.0001 ****
<b>FEV1</b>	1.582 ± 0.02698	1.603 ± 0.02839	P= 0.5838
<b>VC</b>	3.011 ± 0.02022	3.122 ± 0.03923	P= 0.0155 *

	<b>CONTROL MEAN±SD</b>	<b>3<sup>RD</sup> TRIMESTER MEAN±SD</b>	<b>P VALUE</b>
<b>PEFR</b>	5.904 ± 0.02970	5.160 ± 0.01633	P < 0.0001 ****
<b>FEV1</b>	1.582 ± 0.02698	1.604 ± 0.03112	P = 0.5897
<b>VC</b>	3.011 ± 0.02022	3.207 ± 0.03687	P < 0.0001 ****

Significant increase in VC seen in III trimester subjects when compared with control subjects is much more when compared with the increase in VC seen in I & II trimester subjects.

The mean PEFR in I trimester subjects showed a significant decrease of 4.39% when compared with the control non pregnant subjects (p value < 0.0001). Similarly, the mean PEFR in II & III trimester subjects also showed a significant decrease of 8.03% & 12.60% respectively when compared with the control subjects (p < 0.0001 & p < 0.0001 respectively). Statistically significant decrease in PEFR was seen in all the trimesters of pregnancy i.e., I, II & III trimester subjects when compared with control subjects and the decrease was much more in III trimester subjects when compared with the decrease in PEFR seen in I & II trimester subjects.

The mean FEV<sub>1</sub> in I trimester subjects showed a non significant increase of 1.01% when compared with the control non pregnant subjects (p value = 0.6122). Similarly, the mean FEV<sub>1</sub> in II & III trimester subjects showed a non significant increase of 1.33% & 1.52 % respectively when compared with the control subjects (p = 0.5838 & p = 0.5897 respectively). No significant change was observed in FEV<sub>1</sub> in different trimesters of pregnancy.

### Discussion

In our study Forced Expiratory Volume in one second (FEV<sub>1</sub>) did not show any significant change in I, II & III trimester's of normal pregnant women as compared to non pregnant women except PEFR which showed a significant decrease in different trimesters of pregnancy. Regarding Vital Capacity, some of the studies [12,14-15] showed a

decrease in FVC and few other studies [10] showed a significant increase in VC. In our study we observed a significant increase in VC.

No change in FEV<sub>1</sub> was observed in few studies [14,16]. The results of our study correlate with the findings of the above studies regarding FEV<sub>1</sub>. Berry M J et al [17] found that during pregnancy, no change in Vital Capacity (VC) was observed. Also concluded that no change occur in FEV<sub>1</sub>/FVC% and Forced Expiratory Volume in 1st second i.e., FEV<sub>1</sub>%. The inspiratory Capacity was increased during pregnancy, due to the altered thoracic configuration and also due to heightened sensitivity to the nervous stimuli required to produce muscular contraction. Also a decline in the Expiratory Reserve Volume and a persistent increase in Tidal Volume (VT) were observed.

Significant decrease in FEV<sub>1</sub>, FVC and FEV<sub>1</sub>/FVC was found in all the three trimesters of pregnancy in a study conducted by Shazia Batool et al [18]. Highly significant decline in FVC, FEV<sub>1</sub>, FEV<sub>3</sub>, MVV and PEFR in all the trimesters of pregnancy as compared to control was observed in a study by Sushma Jadhav et al [19].

No change in Vital Capacity, Peak Expiratory Flow Rate or Forced Expiratory Volume in one second in pregnancy was observed by Nelson Piercy C [20]. Also observed that normal pregnancy is associated with a 20% increase in oxygen consumption and a 15% increase in the maternal metabolic rate and this extra demand is achieved via 5 % increase in Resting Minute Ventilation, resulting from a rise in Tidal Volume rather than respiratory rate leading to hyperventilation. No change in Vital Capacity was observed in other studies conducted [21].

Decrease in FEV<sub>1</sub> during pregnancy was seen in many studies [15,22-24] but we found a non significant increase in FEV<sub>1</sub>, significant increase in FVC. Puranik BM et al [22] conducted a longitudinal study on pulmonary function tests during pregnancy and they concluded that the Vital Capacity remained unchanged throughout pregnancy and no change was observed in FVC, FEV<sub>1</sub> which correlates with the findings of our study. Also decline in PEFR which was observed in this study correlates with our study. Mrunal SP et al [23] studied about the antenatal changes in lung function tests and importance of postpartum exercises in their recovery. They concluded that the change in FVC and FEV<sub>1</sub> % were insignificant, but they observed a significant decline in Expiratory Reserve Volume and PEFR and the increment in Inspiratory Capacity

was also significant. The decline in PEFR observed in our study is in accordance with this study.

Pande Y et al [25] observed that the inconsistent decrease in vital capacity in some of the subjects during pregnancy was in accordance with the observation of earlier investigators who found it to be either unchanged or slightly decreased during pregnancy. The Forced Expiratory Volume in the first second (FEV<sub>1</sub>) and Maximum Mid Expiratory Flow Rate (MMFR) were increased. Kaltreider NL et al [26] observed that during pregnancy, oxygen consumption and Minute Ventilation consistently increased and there was individual variation both in direction and magnitude of the change in Vital Capacity. The FEV<sub>1</sub>/VC remained unaltered. The Expiratory Reserve Volume (ERV) and Functional Residual Capacity (FRC) were decreased progressively. They also found that the Residual Volume (RV) was also progressively decreased. A decrease in FVC, FEV<sub>1</sub> & PEFR in pregnancy was observed by Neeraj Candy S et al [27] and our study do not correlate with this study. A non significant increase in FEV<sub>1</sub> and significant increase in VC and a decrease in PEFR are observed in our study.

Decline in PEFR during the third trimester of pregnancy as observed by Hemant Deshpande et al [28] correlates with our study. In fact, we observed a gradual decline in PEFR in different trimesters of pregnancy. The results of our study were in contradictory to the study of Weerasekara DS et al [29] who observed that there is no significant change in VC & PEFR in pregnancy.

Grindheim G et al [30] conducted a longitudinal study to observe the changes in pregnancy and evaluate the influence of parity, pre-gestational overweight and excessive weight gain on lung function. They observed that the FVC & FVC% increased after first trimester of pregnancy. Also found that VC, FVC%, PEFR and PEF% in early and mid-pregnancy were significantly lower compared with the postpartum value.

Knox AJ et al [31] observed a number of physiological changes which occurred during pregnancy. No significant change in peak flow rates, forced vital capacity or forced expiratory volume in the first second (FEV<sub>1</sub>) is observed. The decline in PEFR in the first trimester can be attributed to morning sickness, lack of nutrition whereas in second and third trimester it may be due to mechanical pressure of enlarging gravid uterus, Elevation of the diaphragm and restrictive movements of lungs. PEFR is more sensitive to muscular element in respiration and as anemia

produces muscle weakness, it reflects in lowering the PEFR (Singhal et al) [32].

Sunyal DK et al [33] studied about the Peak expiratory flow rate in pregnant women in Bangladesh. They observed that the Peak expiratory flow rate and their percentage of predicted values were significantly lower during third trimester of pregnancy compared to controls and it progressively decreased from first to third trimester. PEFR in our study also showed a similar decrease in different trimesters of pregnancy. The decline in PEFR during pregnancy occurs suggestively due to lesser force of contraction of main expiratory muscles like anterior abdominal muscles and internal intercostal muscles [34-35]. The results of most studies done on western populations indicate that vital capacity and peak expiratory flow rate do not change significantly throughout the course of pregnancy.

### Conclusion

Comparative study of pulmonary function tests in different trimesters of pregnancy showed a decrease in PEFR significantly which may be due to mechanical pressure of gravid uterus, diaphragm restricting the movement of lungs especially in third trimester of pregnancy. There was a decrease in respiratory parameters like PEFR from first to third trimesters of pregnancy which may also be due to poor nutrition. To establish the precise cause of decrease in different lung function parameters, further studies are to be undertaken like hormonal assay in different trimesters to study the effect of hormones on different lung function parameters. Continuous Monitoring of lung function in different trimesters provide adequate information regarding maternal healthcare. Obstructive / restrictive lung disorders during pregnancy can be identified early which can be prevented by proper management. Pregnant women need regular monitoring of lung function by spirometry in order to optimize their lung function throughout pregnancy.

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### Conflicts of Interest: Nil

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### References

1. Contreras G, Gutierrez M, Beroiza T, Fantin A, Oddo H, Villarroel L. Ventilatory drive and respiratory muscle function in pregnancy. *Am Rev Respir Dis* 1991; 144: 837–41.
2. Weinberger SE, Weiss ST, Cohen WR. Pregnancy and the lung. *Am Rev Respir Dis* 1980; 121(3):559–81.
3. Gilroy RJ, Mangura BT, Laviertes MH. Rib cage and abdominal volume displacements during breathing in pregnancy. *Am Rev Respir Dis* 1988; 137(3): 668–72.
4. Cugell DW, Frank NR, Gaensler EA. Pulmonary function in pregnancy. I. Serial observations in normal women. *Am Rev Tuberc* 1953; 67(5):568–97.
5. Alaily AB, Carrol KB. Pulmonary ventilation in pregnancy. *Br J Obstet Gynaecol* 1978; 85:518–24.
6. GarciaRio F, PinoGarcia JM, Serrano S. Comparison of helium dilution and plethysmographic lung volumes in pregnant women. *Eur Respir J* 1997; 10 (10):2371–5.
7. Goucher D, Rubin A, Russo N. The effect of pregnancy upon pulmonary function in normal women. *Am J Obstet Gynecol* 1956; 72(5):963–9.
8. Kolarzyk E, Szot WM, Lyszczarz J. Lung function and breathing regulation parameters during pregnancy. *Arch Gynecol Obstet* 2005; 272:53–8.
9. McAuliffe F, Kametas N, Costello J, Rafferty GF, Greenough A, Nicolaides K. Respiratory function in singleton and twin pregnancy. *BJOG* 2002; 109 :765–9.
10. Milne JA. The respiratory response to pregnancy. *Postgrad Med J* 1979;55: 318–24.
11. Harirah HM, Donia SE, Nasrallah FK, Saade GR, Belfort MA. Effect of gestational age and position on peak expiratory flow rate: a longitudinal study. *Obstet Gynecol* 2005; 105:372–6.
12. Puranik BM, Kurhade GA, Kaore SB, Patwardhan SA, Kher JR. PEFR in pregnancy: a longitudinal study. *Indian J Physiol Pharmacol* 1995; 39:135–9.
13. McAuliffe F, Kametas N, Espinoza J, Greenough A, Nicolaides K. Respiratory function in pregnancy at sea level and at high altitude. *BJOG* 2004; 111:311–5.
14. Mokkaipatti R, Prasad EC, Venkatraman, Fatima K. Ventilation functions in pregnancy. *Indian J. Physiol and Pharmacol* 1991; 34(4): 237 - 249.
15. Monga U, Kumar K. Pulmonary functions in Punjabi Pregnant Women. *Indian J Physiol Pharmacol* 2000; 44(1): 115-16.

16. Milne JA, Howie AD and Pack AI. Large airways function during normal pregnancy. *British. J of Obst & Gynec* 1977; 448-51.
17. Berry MJ, Mc Murray RG, Katz VL. Pulmonary & ventilatory response to pregnancy; Immersion and Exercise. *J of Applied physiology* 1989 February (66) Vol 8; 57
18. Shazia Batool, Rabbia Shakoor, Ghulam Mustafa, Ammar Anwer, Shahroona Masood, Zafar H Tanveer. Comparison of lung functions of pregnant women with non pregnant women at sheik zayed hospital, Rahimyarkhan. *JSZMC* 2008, Vol (2) ;155-8.
19. Sushma J, Dudhamal V B , Karadkhedkar S S, Sayeeda Afroz, Razvi N A . Comparative study of pulmonary function tests on different trimesters of pregnancy. *Int J Cur Res Rev*, Jan 2013; Vol 05 (02); 118-22.
20. Nelson Piercy C. Respiratory diseases. *Hand book of Obstetric Medicine*, Oxford 1997; 15-65.
21. Knox AJ, Petkova D, Johnson S .Respiratory disease in pregnancy. *Obstet & Gynaecol* 1999; 9; 69-74.
22. Puranik BM, Kaore SB, Kurhade GA, Agrawal SD, Patwardhan SA, Kher JR. A Longitudinal study of pulmonary function tests during pregnancy. *Indian J Physiol Pharmacol* 1994; 38(2): 129-32.
23. Mrunal SP, Kurhade GA. 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.
24. Pandey MR. Nishith SD, Bhatt RV. Pulmonary function in pregnancy. *J Obst Gynae India* 1972; 22;1-3.
25. Pande Y, Guleria JS, Hingorani V. Pulmonary ventilation & gas exchange in pregnancy. *Indian J. of Obst & Gynae* 1973; 710-5.
26. Kaltreider NL, Rosen M, Cugell DW, Frank NR. Pulmonary function during pregnancy in normal woman and in patients with cardiopulmonary disease. *Thorax* 1970; 25; 445.
27. NeerajCandy S, Pramod J, Singh J, Kaur V. Effect of advanced uncomplicated pregnancy on pulmonary function parameters of north Indian subjects. *Indian J Physiol Pharmacol* 2010: 54(1): 69-72.
28. Deshpande H, Madkar C, Dahiya P. A study of pulmonary function tests in different stages of pregnancy. *Int J Biol Med Res*. 2013; 4(1): 2713-6.
29. Weerasekara DS, Kusuma Ruberu D, Sivayogan S. Pulmonary functions in pregnant srilankan women. *Sabaragamuwa University Journal* 1999; vol 2(1); 57-60.
30. Grindheim G, Toska K, Estensen ME, Rosseland LA. Changes in pulmonary function during pregnancy: a longitudinal cohort study. *BJOG* 2012; 119:94–101.
31. Knox AJ, Petkova D, Johnson S .Respiratory disease in pregnancy. *J Obstet & Gynaecol* 1999; 9; 69-74.
32. Singhal V, Saxena K. Effect of anemia on respiratory and metabolic during 3rd trimester of pregnancy. *Indian J physiol pharmacol* 1987; 3(2); 130-5.
33. Sunyal DK, Amin MR, Ahmed A, Begum S, Begum M, Rahman NJ. Peak expiratory flow rate in pregnant women. *Bangladesh Soc Physiol* 2007;2: 20-23 .
34. Saxena SC, Rao VSC, Mudgal SA. Study of pulmonary function tests during pregnancy. *J Obs Gyn (India)* 1979; 29: 993–5.
35. Ganeriwal SK, Deshpande DR, Reddy BV, Shaikh RM. Effect of pregnancy on pulmonary Ventilation. *J Obs Gyn (India)* 1984; 36: 639–41.