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Pulmonary Function Tests in Patients with Chronic Rhinosinusitis: A Comparative Study

By Prof Dr. Ramesh Azad, Dr. Siddharth Sharma & Dr. Sunil Sharma

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Methods: The study was conducted in the Department of ENT and Pulmonary Medicine, IGMC Shimla on 27 adult patients of chronic rhinosinusitis and 26 age matched healthy control subjects between November 2018 and November 2019. The pulmonary function tests were performed using a spirometry and included Vital Capacity (VC), Forced vital capacity (FVC), Forced expiratory volume in one second (FEV1) and Forced expiratory volume in one second by Forced vital capacity ratio (FEV1/FVC).

Results: Following observations were made from the results of the study, among controls majority of the subjects i.e 23 (92%) had a normal PFT with only 1 patient (4%) had mild and another 1 patient (4%) had moderate restriction on PFT. Whereas among the patients group 10 patients (37%) had a normal pre-operative baseline PFT. Another 10 (37%) and 7 patients (25.9%) had mild and moderate obstruction respectively.

Keywords: chronic rhinosinusitis, unified airway, PFT, lung function.

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PULMONARYFUNCTIONTESTSINPATIENTSWITHCHRONICRHINDSINUSITISACOMPARATIVESTUDY

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Pulmonary Function Tests in Patients with Chronic Rhinosinusitis: A Comparative Study

Prof Dr. Ramesh Azad ^a, Dr. Siddharth Sharma ^e & Dr. Sunil Sharma ^e

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Conclusion: Findings of the current study suggest that their are latent derangements in the pulmonary functions of patients with upper airway disease ie chronic rhinosinusitis and the difference between the pulmonary functions of patients and controls is significant. Hence our study validates the Unified airway disease model and suggests that lower airway evaluation shall be done in patients with chronic upper airway disease and a timely diagnosis and intervention could be beneficial for the pulmonary diseases.

Keywords: chronic rhinosinusitis, unified airway, PFT, lung function.

I. INTRODUCTION

hronic rhinosinusitis (CRS) is an inflammatory disease of the mucosa of the nasal cavity and paranasal sinuses, with symptoms lasting 12 weeks or more. It is a common disease and represents a public health problem resulting in socioeconomic burden throughout the world[1]. The exact pathogenesis of CRS is still not clearly understood; it significantly affects quality-of-life measures with decrements in general health perception and social functioning. It has been long recognized that diseases coexist in the upper and lower airways. The "United-Airways" concept implies that there is a link between upper and lower airway inflammation[2].

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Galenus was the first to note an association of nasal symptoms and asthma in the 2nd century he defined the nose as a respiratory instrument in his work "De usupartium". From the literature, it appears that Kariya et al[3] were the first to report the latent obstruction in low diameter airways associated with CRS. Their suggestion was that obstructive pulmonary alterations were present in CRS cases, even where they caused no symptoms and had not led to a diagnosis of obstructive airway disease. Chien et al[4] reported that chronic obstructive pulmonary disease was associated with an increased risk of Chronic rhinosinusitis without nasal polyps, independent of a number of potential confounding factors. Both the upper and lower respiratory tracts are continually exposed to gases and airborne particles. The importance of host defence mechanisms is vital. Physical barriers, mucociliary clearance systems with immune processes combine to protect the lung from damage. When abnormalities of these mechanisms occur, the patient may first present to the otolaryngologist before serious bronchopulmonary involvement occurs. The mechanisms that connect upper and lower airway dysfunction are under investigation. They include a nasal-bronchial reflex, mouth breathing caused by nasal obstruction, and pulmonary aspiration of nasal contents and a direct evidence of an association could be obtained by examining the objective results of pulmonary function tests of patients with CRS and comparing them with results of healthy controls.

II. METHODS AND MATERIALS

The study was conducted in the Departments of Otorhinolaryngology and Pulmonary medicine, Indra Gandhi Medical College, Shimla, Himachal Pradesh between November 2018 and November 2019.

a) Study type

A hospital based study.

b) Study Design

A comparative observational study.

c) Subjects and Sample size

The study included 52 participants with 27 patients fulfilling the clinical criteria for CRS and 25 age matched healthy controls. The subjects in both groups belonged to the age group 18–60 years.

d) Inclusion Criteria

Patients suffering from CRS as documented by examination and history, clinical radiological investigations were included in the patients group. Symptoms and findings of CRS were based on the CRS criteria and divided into major and minor factors as American established by the Academy of Otolaryngology and Head and Neck Surgery in 1996. These parameters have since become widely adopted researchers and by health maintenance bv organizations CRS diagnosis requires presence of at least 2 major factors or one major factor with 2 or more minor factors, or nasal purulence on examination.

e) Exclusion Criteria

- Pregnant women

Coexistent systemic diseases like diabetes, hypertension, neoplasia.

- Patients with known psychiatric illness.
- Prior paranasal sinus, nose or throat surgery.

f) Controls

The study participants belonging to the same age group as patients group who presented to ENT OPD after ruling out CRS and the exclusion criteria by means of clinical examination and were taken as controls.

g) Pulmonary Function Tests

PFT were assessed using Spirometer, Vitalograph model number 6800. Spirometry is a powerful tool that can be used to detect, follow, and manage patients with lung disorders. It is reliable and relatively simple to incorporate into a routine office visit. To determine the validity of spirometric results, a Minimum of 3 acceptable maneuversviz: FEV1 and FVC maneuver, VC and IC maneuver and Peak exploratory flow were done as per criteria laid down by latest ATS (American thoracic society) guidelines. In each test,

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patients exhaled for at least six seconds and stopped when there was no volume change for one second. The test session was finished when the difference between the two largest FVC measurements and between the two largest FEV1 measurements was found to be within 0.2 L.

Results of spirometery were graded as per Gold 2017 standards[5]. The PFT values were considered to indicate significant airway obstruction when FEV1/FVC < 0.7 and FEV1 < 80 % of the predicted value for a patient's age, height and weight.

h) Statistical analysis

Data is presented as frequency, percentages, and maximum minimum and mean \pm SD wherever applicable. Data is also depicted in tables and bar graphs wherever possible. Proportions are compared by Chi square as appropriate to the data set. Means are compared by student t test and a P value of < 0.05 is considered to be statistically significant.

III. Results

A total of 52 adult subjects including 27 CRS patients and 25 healthy controls who presented to ENT OPD after thorough clinical history and physical examination were enrolled for the study after matching them with our inclusion and exclusion criteria.

a) Age distribution of participants

The age of participants included in patient group ranged from 18 to 60 years with a mean of 40.59 \pm 13.49 years. The maximum number of patients (15) belonged to 41-60 age group representing 55.6% of total, in control group age ranged from 30 to 60 years with a mean of 48.68 \pm 11.32 years with maximum participants in 41- 60 years representing 72% of the total. The data is depicted in Table 1.

Table 1:	Age wise distribution of	f study group and	control group

Age (in years)	Study	/ group	Co	ntrols	Chi-square value	P value
	Frequency	Percentages	Frequency	Percentages		
≤20	2	7.4	0	0.0	3.201	0.362 NS
21-30	6	22.2	3	12.0		
31-40	4	14.8	4	16.0		
41-60	15	55.6	18	72.0		
Total	27	100.0	25	100.0		

Statistical Analysis: Chi-square test. P≤0.05: Statistically significant (S)

b) Demographic profile of participants

The weight of patients in the study group ranged from 35kg to 82 kg with a mean of 61.56 ± 13.45 kgs. The height of the patients in study group ranged from 145cm to 185cm with a mean of 165.07 \pm 10.38cm. The BMI ranged from 15 to 28.50 with a mean

of 22.69 \pm 3.53. In the control group weight ranged from 35kg to 79 kg with a mean of 60.12 \pm 11.37kg. Height ranged from 105cm to 180cm with a mean of 160.16 \pm 15.35. Mean BMI among control group was 22.60 \pm 3.50. The demographic data is represented in Table 2.

Statistics	Age (control)	Age (Patients)	Weight (Controls)	Weight (Patients)	Height (Controls)	Height (Patients)	BMI (Controls)	BMI (Patients)
Minimum	30	18.00	35.00	35.00	105.00	145.00	14.50	15.00
Maximum	60	60.00	79.00	82.00	180.00	185.00	29.30	28.50
Mean	48.68	40.59	60.12	61.56	160.16	165.07	22.60	22.69
Median	50.00	41.00	60.00	60.00	160.00	165.00	22.80	24.00
Std. deviation	11.32	13.41	11.37	13.45	15.35	10.38	3.50	3.53
Std. Error of mean	2.26	2.58	2.27	2.59	3.07	2.00	0.70	0.68

Table 2: Demography of participants

c) PFT Values in controls

Majority of patients (23) in control group had a normal pulmonary function accounting for 92% of total whereas one patient each had mild and moderate restriction in pulmonary function test. Mean Vital capacity among controls was 82.80% of predicted. The mean FVC was 90.08% of predicted value. FEV₁ mean was 72.89% of the predicted value. FEV₁/FVC ratio had a mean 97.12% of the predicted value. The data is represented in Table 3.

Table 3: Pulmonary	function in	Control group
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Statistics	VC/ Vital capacity	VC/ Vital capacity %*	FVC(L)	FVC%	FEV1(L)	FEV1%	FEV1/FVC	FEV1/FVC%
Minimum	1.85	60.00	1.99	72.00	1.58	50.00	1.07	70.00
Maximum	3.82	114.00	4.19	119.00	3.48	117.00	1.43	112.00
Mean	3.07	82.80	3.23	90.08	2.63	84.80	1.28	97.12
Median	3.33	83.00	3.36	88.00	2.68	84.00	1.29	100.00
Std. deviation	0.57	13.15	0.61	11.65	0.46	14.72	0.08	10.16
SE of mean	0.11	2.63	0.12	2.33	0.09	2.94	0.02	2.03

d) PFT in patients group

Among patients group of 27, 10 (37%) had a normal pre-operative baseline PFT. Another 10 (37%) and 7(25.9) had mild and moderate obstruction. Mean Vital capacity of the patients was 74.59% of predicted. The mean FVC was 81.15% of predicted value. FEV₁ mean was 72.89% of the predicted value. FEV₁ /FVC ratio had a mean 90.16% of the predicted value for patients. The data is represented in table 4.

Table 4: Pulmonary function in patients group	Table 4:	Pulmonary	function	in	patients	group
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Statistics								
Statistics	VC (L)	VC %*	FVC(L)	FVC%	FEV ₁ (L)	FEV₁%	FEV ₁ /FVC	FEV₁/FVC%
Minimum	1.29	46.00	1.12	41.00	0.77	33.00	0.35	43.00
Maximum	3.96	97.00	5.00	112	3.57	105.00	0.95	115.00
Mean	2.60	74.59	2.75	81.15	2.08	72.89	0.73	90.19
Median	2.53	73.00	2.68	81.00	2.03	76.00	0.75	96.00
Std.deviation	0.80	14.41	0.90	18.21	0.70	19.01	0.14	18.38
S.E of mean	0.15	2.77	0.17	3.50	0.13	3.66	0.03	3.54

Our study observed that among controls out of 25 majority of the subjects i.e 23 (92%) had a normal PFT with only 1 patient (4%) had mild and another 1 patient had moderate restriction on PFT. Whereas among the patients group out a total of 27, 10 (37%) had a normal pre-operative baseline PFT. Another 10

(37%) and 7(25.9) had mild and moderate obstruction respectively.

The mean difference between the VC of controls and patients was 0.47 Litres with a p value of 0.018.The mean difference between the FVC of controls and patients was 0.48 litres with a p- value of 0.030. The mean difference between the FEV1 of controls and FEV1 of patients was 0.55 Litres with a p-value of 0.002. The mean difference between the FEV1 / FVC ratio of controls and patients was 0.55 with a p- value of 0.001.

Thus our study found a statistically significant differences between the PFT of the controls and patients with CRS among all the parameters studied.

IV. DISCUSSION

Respiratory tract infections are guite commonly managed by health care providers. A recent report from the Centers for Disease Control, respiratory infections (upper respiratory tract infections, otitis media, and lower respiratory tract infections) accounted for 16% of all outpatient visits of patients to physicians[6]. The nose functions as the air conditioner of human body delivering inspired air to the tracheobronchial tree which comes into contact with large surface area of the nasal mucosa permitting three physiological functions of: filtration, warming and humidification[7]. Inhalation of warm, moist air into the lower airway is required for the optimal functioning of the bronchopulmonary system hence chronic nasal dysfunction can contribute significantly to lower respiratory diseases such as asthma and chronic bronchitis.

The present study is an observational and comparative study including a total of 52 subjects with 25 healthy controls and 27 patients of CRS whose pulmonary function tests were evaluated with the objective to evaluate and compare the pulmonary function among the two groups. There have been quite a few studies demonstrating association of upper airway diseases with asthma and other lower airway conditions in past[8] but to the best of our knowledge this is one of the first study done in the Himalayan state of Himachal Pradesh to study the association of lower airway involvement in CRS.

Out of 27 CRS patients maximum number of patients n = 27 (55.6%) belonged to the age group of 41-60 years with mean age of 40.59 years. Similarly Sokol in a 2000 study[9] evaluated 16,000 patients suffering from CRS and found the mean age of patients comparable i.e 44 years.

In our present study majority of patients were male with n = 17 (63%) whereas females constituting 37% with n = 10. With a male to female ratio of 1.7:1. This is in accordance with a similar study done by Ahmed M Youssef[10] in 2017 on 25 patients with CRS in which male constituted a majority 60% (n = 15). Another Indian study conducted by Ziauddin Ahmed[11] in 2017 found a male preponderance 32 out of a total of 48 with males constituting 66.6% of the total cases.

a) Pulmonary function tests

Our study found that CRS patients had significantly poorer pulmonary functions compared to healthy controls. Similarly results of a 2004 study by Ragab A et al[12] showed that 60% of patients with CRS had different kinds of lower airway involvement. Some are manifest such as asthma and some are hidden such as small airway disease and bronchial hyper-reactivity.

Our study found significant differences in pulmonary functions of healthy controls and the patients suffering from CRS. A total of 17 patients (63.9%) out of 27 having degranged pulmonary functions. with n = 10(37%) of patients having mild pulmonary obstruction and n= 7 (25.9%) of patients having moderate pulmonary obstruction. Whereas compared to control group where only 2 patients (8%) out of a total of 25 had deranged pulmonary function test with n = 1 patient (4%) having mild and n = 1 (4%) having moderate obstruction. The mean vital capacity in patients group was 2.60 litres (74.59%) as compared to the mean vital capacity in the control group which was 3.07 I (82.80%). This difference was statistically significant with a p value of 0.018. The mean FVC in patients group was 2.75 litres (81.15%) whereas in controls it was 3.23 litres (90.08%) this difference was also statistically significant with a p value of 0.03. The mean FEV_1 in patients group was 2.08 litres (72.89%) whereas in control group the mean FEV_1 was 2.63 (84.80%) the difference between patients and controls was highly significant statistically with a p value of 0.02. The mean FEV₁/FVC ratio in the patients group was 0.73(90.19%) as compared to control group which had a mean of 1.28(97.12%). Table 11,12,13,14. Findings of our present study matched the findings of a study done by Kariva S et al[13] in 2014 to evaluate the pulmonary function of 273 patients with CRS and 100 age-matched normal control subjects .CRS patients had significant obstructive lung function changes regardless of the presence of asthma. Similar results were observed in a retrospective study done on 284 subjects by Suh Young Lee et al[14] in 2010 to evaluate the influence of chronic sinusitis and nasal polyp on the lower airway of subjects without lower airway diseases. This study provided evidence that the presence of CT findings suggestive of chronic sinusitis is associated with a subclinical airflow limitation in subjects without lower respiratory disease. A unified airway disease hypothesis[15] may be valid for subjects even without definite lung disease.

The results of our study showed that majority of the patients had derranged pulmonary functions even if they had not been diagnosed with an overt lower airway disease these results are in accordance with the results of a study done by GottliebMJ[16] in a 1925 on 117 cases of asthma, found that in all there were 31 cases of disease in one or more of the paranasal sinuses, a correlation was found between lower airway and upper airway diseases.

This nonsymptomatic lower airway involvement in patients with CRS can be explained by the small lower airway dysfunction that involves the terminal and respiratory bronchioles. Another entity of lower airway functional involvement is the inflammation of the lower airways resulting in bronchial hyperreactivity[17][18]. Nasal obstruction can induce a blockage of the sinus ostia, with a reduction in the availability of nitric oxide in the upper and lower airways, as reported for patients with chronic sinus disease[19].

V. Conclusion

Coexistence of upper and lower respiratory diseases is common. Our study found a significant derangement in the pulmonary functions of patients with CRS not previously diagnosed with any lower airway diseases. A united airway disease hypothesis may thus be valid. How exactly the upper airway disease effects lower airway is still a matter a of debate and beyond the scope of present study. However several hypotheses exist, like nasopharyngo bronchial reflexes that may lead to airway hyper responsiveness. Also sinonasal mucosal inflammation may lead to pulmonary inflammation by releasing chemotactic factors and leukocytes. Oral breathing caused by CRS may also lead to inhalation of cold dry air and environmental pollutions bypassing the protective nasal function. Although our study is limited by a relatively small number of patients with most of the patients having the disease since many months if not years, our comparative study, with its well-defined outcome and criteria included for patient selection, helps to clarify the true value of lower airway assessment in these difficult-to-treat patients and to emphasize that the underuse of objective testing, such as spirometry for patients with CRS may lead to underdiagnosed lower airway problems.

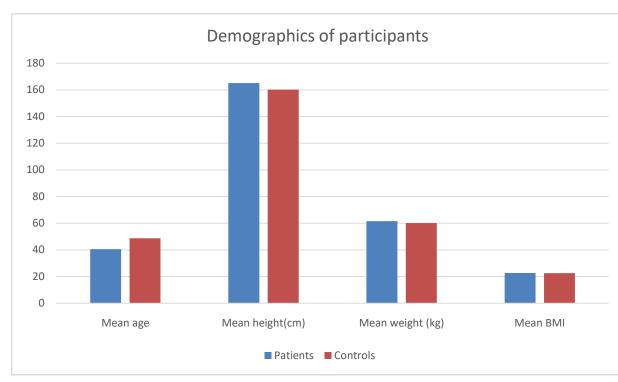


Figure 1: Demographic data of participants

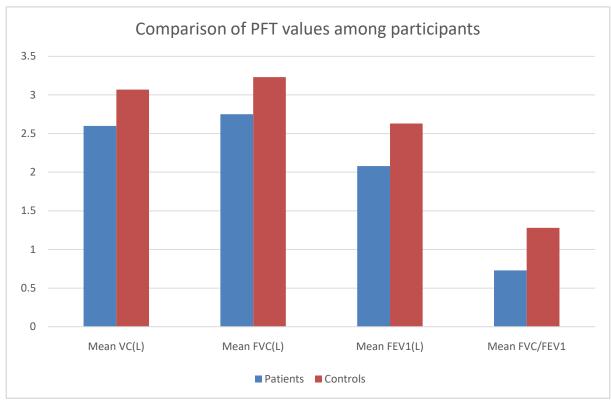


Figure 2: PFT comparison between patients and controls

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