**Original Research Article** 

# Assessment of Body Composition and Heart Rate Variability in Bronchial Asthma

Md. Perwez Salam<sup>1</sup>, Adhiraj Singh<sup>2</sup>, Eema Chaudhary<sup>3</sup>, Kiran Singh<sup>4\*</sup>

<sup>1</sup>Ex JR Department of Physiology, Subharti Medical College Meerut, U.P., India <sup>2</sup>MBBS Final Year Student, Subharti Medical College Meerut, U.P., India <sup>3</sup>Professor & Head, Department of Pulmonary Medicine, Subharti Medical College Meerut, U.P., India <sup>4</sup>Professor & Head, Department of Physiology, Subharti Medical College Meerut, U.P., India

Received: 05-07-2020 / Revised: 03-09-2020 / Accepted: 12-09-2020

#### Abstract

**Background**: Previous studies were focused mainly either on autonomic modulation using heart rate variability (HRV) or body composition in the bronchial asthma patients, but our study has correlated both HRV and body composition indices in asthmatic patients. **Aim & Objective**:To evaluate the possible association of HRV and body composition parameters with main spirometric indices in newly diagnosed asthmatic patients & to compare these findings with healthy subjects. **Methods:** Thirty asthmatic patients (study group) and 30 apparently healthy subjects (controls) in age group of 20-55 years of both genders were included. BMI, waist-hip ratio (WHR) and body fat % were measured. Time and frequency domain parameters of HRV were recorded to determine sympathetic and parasympathetic autonomic modulations. Forced vital Capacity (FVC), Forced expiratory volume at 1st second (FEV<sub>1</sub>) & FEV<sub>1</sub>/FVC ratio parameters were recorded. **Results:** HF ms<sup>2</sup> & HF n.u were significantly (P<0.001) higher and LF n.u & LF-HF ratio were significantly (P<0.001) lower in asthmatics as compared to control. WHR was found to be significantly higher in study group as compared to control (P<0.001). We found a negative correlation of LF n.u & LF- HF ratio & positive correlation of HF n.u with FEV1(L), FEV1/FVC % & WHR which was not found statistically significant. **Conclusion:** In our study we observed altered autonomic activity with parasympathetic dominance as HF component was higher in asthmatic patients. We also found a significant association between WHR and asthma.

# **Key words:** Bronchial asthma, body composition, heart rate variability.

This is an Open Access article that uses a fund-ing model which does not charge readers or their institutions for access and distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0) and the Budapest Open Access Initiative (http://www.budapestopenaccessinitiative.org/read), which permit unrestricted use, distribution, and reproduction in any medium, provided the original work is properly credited.

## Introduction

Bronchial asthma has become more common in both children and adults around the world in recent decades, particularly because of its early onset. The increase in the prevalence of asthma has been associated with an increase in atopic sensitization, and is paralleled by similar increases in other allergic disorders such as eczema and rhinitis. [1] [2] A report estimated that 300 million people worldwide had asthma, and projected that this number would increase to 400 million by 2025, as countries became more urbanized. [3]

\*Correspondence

**Dr. Kiran Singh**Professor & Head
Department Of Physiology, Subharti Medical
College, Meerut – 250005, U.P., India.

The exact pathogenesis of bronchial asthma is multifactorial. The exaggerated broncho-constriction along with hyper-reactivity in bronchial asthma is associated with abnormal autonomic nervous system (ANS) control. [4] The parasympathetic nerves provide the dominant autonomic control of airway smooth muscle. They release acetylcholine onto muscarinic receptors, causing bronchoconstriction. [5] Studies have indicated that altered autonomic control of airway caliber in asthma is reflected by parallel changes in heart rate. Previous heart rate variability (HRV) studies in asthmatic subjects demonstrate predominance of parasympathetic drive concomitant with low HRV.[ 6, 7] which is against the general belief that enhanced parasympathetic modulation improves HRV. Other studies reported lack of association between bronchial asthma & ANS .[8, 9]

e-ISSN: 2590-3241, p-ISSN: 2590-325X

There are also studies suggesting that obesity represents a risk factor for the development of asthma, as both conditions are associated with underlying inflammatory pathway & thus seems to be associated. Increase in body mass index (BMI) values predisposes to new asthma development even in young adults .[10] A gender dependent influence has been observed with a significant association between body fat & asthma in women .[11] Overweight has been associated with an increase in inflammatory markers & with an imbalance in ANS. [12]

In the view of above data and lack of studies, the present study was designed to investigate HRV and the body composition parameters in newly diagnosed asthmatic patients and to compare these finding with healthy subjects. Study was also planned to see the possible association of cardiovascular autonomic changes using HRV and body composition parameters with patho-physiology of asthma by more elaborate and reliable pulmonary function tests.

#### Material and methods

After approval from institutional ethical committee on human experimentation, this observational study was conducted in the research lab of department of Physiology, in collaboration with department of pulmonary medicine Subharti medical college and associated Chhatrapati Shivaji Subharti hospital, Meerut. Informed consent from all the subjects was obtained before the start of the study.

## **Study Subjects**

The study group comprised of 30 patients of bronchial asthma in the age group of 20 to 55 years of both genders who were newly diagnosed (not taking any medication) with clinical symptoms along with positive PFT findings as defined by the American Thoracic Society recommendation. [13] Age & sex matched 30 apparently healthy volunteers were taken as control.

Exclusion criteria: having history of any other cardiorespiratory disease, hypertension, neuro-endocrine diseases, any psychiatric illness, allergy, atopy, urticaria, alcohol or any recreational drug and smoking in both groups.

After evaluating the past medical history including the bronchial-asthma treatment offered to the patient, all volunteers underwent clinical examination investigations during the same hours of the day preferably between 11 AM-1PM to avoid confounding, effects of circadian rhythm on asthma severity & HRV. All the participants were asked to report to the research laboratory of the department with following instructions:

- a) No eating or drinking 3 to 4 hours prior to the test
- b) No exercise 12 hours prior to the test.
- c) No caffeine consumption 24 hours prior to the test.
- d) To remove all metal objects.

#### and **Anthropometric** body composition measurements

Height was measured by stadiometer to nearest 1 cm and weight, by weighing machine (Krups) to the nearest 1 kg with subjects standing without shoes and wearing light clothes. Circumferences at waist (at the level of umbilicus) and hip (at the level of maximum extension of hips) were measured with a tape measure nearest to 0.1 cm. Participants were asked to lie in the supine position for 5 minutes breathing normally in a relaxed position. Body mass index, waist-hip ratio (WHR), body fat percentage (BF%) were measured by Multi-frequency impedance technique using Body stat QUADSCAN 4000 version 4/10(Isle of Man, UK) which employs multi-frequency (5, 50, 100 & 200 kHz) currents through a tetra polar hand-to-foot impedance model. Care was taken to place the arm well apart from the torso and the legs not touching one another.

### **Assessment of HRV**

Analysis of HRV from ECG recording is an important method for assessing cardio-vascular regulations. After 5 min of supine rest Lead II ECG recordings were done (25mm/sec & at voltage at 10 mm/mV) for 5 min. ECG was taken to record all three leads at a sampling frequency of 256Hz to PC and were analyzed offline after visual checking of abnormal ECG. The data recorded was subjected to time domain and frequency domain analysis using the HRV analysis (RMS Polyrite D software version Recommendation of Task Force on HRV followed. [14]The time domain parameters included was SDNN (standard deviation of the averages of NN intervals). Frequency domain analysis was performed using non-parametric method of Fast -Fourier Transformation. The power frequency spectrum was subsequently quantified into standard frequency domain measurements as Total power (TP) in ms<sup>2</sup>, Low frequency (LF) component (0.04-0.15 Hz) & High frequency (HF) component (0.15-0.4Hz) in normalized units (n.u) and power ms<sup>2</sup>& LF-HF ratio (LF/HF).

# **Pulmonary function tests**

Standardized protocol of performing spirometry was followed. Each subject performed 3 trials (with at least two reproducible and acceptable maneuvers) of all parameters according to the American Thoracic Society recommendation. Pulmonary function tests were recorded by Spirodoc (Spiro Pro 6.6 Spirometry standard mode, version no.A23-ow-06145). The

e-ISSN: 2590-3241, p-ISSN: 2590-325X

parameters included were Forced vital Capacity (FVC) in L, Forced expiratory volume at  $1^{st}$  second (FEV<sub>1</sub>) in L and FEV<sub>1</sub>/FVC % ratio. When the subject coughs or did not performed the technique correctly, the turn was ignored and repeated.

#### **Statistical Analysis**

Software graph pad Instat 6.07 version and Microsoft excel were used for statistical analysis. All values were expressed as Mean± SD. Unpaired Student's t test was used to find out the level of significance between the two groups. Pearson correlation coefficient (r) was used to assess any possible association between the variables. A value of P< 0.05 was considered statistically significant.

#### **Results**

In present study, BMI was within normal range (25.14  $\pm 2.25$  & (23.26  $\pm 5.57$ ) in study and control groups respectively. There was no significant difference (P>0.05) in BMI & body fat% between 2 groups.

However, WHR was found to be significantly higher (P<0.001) in asthmatics as compared to control group. FVC (L) and FEV $_1$  (L) were significantly (P<0.001) lower in asthmatics as compared to control. We did not find any significant difference in FEV1/FVC % between 2 groups.

In our study we observed altered autonomic activity with parasympathetic dominance in asthmatic patients. HF component of HRV was significantly higher (P <0.001) in asthmatics as compared to controls whether measured in absolute or normalized units. TP (ms²) & LF (ms²) were lower in asthmatics but the difference was not significant (P >0.05). However, the LF n.u. and LF-HF ratio were significantly (P<0.001) lower in asthmatics as compared to control. SDNN was lower in study group but this was statistically insignificant. LF n.u. & LF-HF ratio were negatively correlated while HF n.u. was positively correlated with FEV1 (L), FEV1/FVC % & WHR which were not found statistically significant.

Table-1: Body Composition & Spirometry Parameters in Study group & Control (n=30) in each group

	Study group	Control	P-value
Parameters	Mean±SD	Mean±SD Mean±SD	
BMI (Kg/m²)	25.14±2.25	23.26±5.57	0.092
WHR	1.01±0.03	0.96±0.04	< 0.001
BODY FAT%	26.62±8.48	24.16±5.24	0.182
FVC(L)	2.23±0.69	4.46±0.48	< 0.001
FEV1(L)	1.84±0.71	3.66±0.39	< 0.001
FEV1/FVC (%)	80.43±14.34	82.18±2.09	0.512

BMI-Body mass index, WHR–waist-hip ratio, FVC- Forced vital capacity, FEV $_1$ - Forced expiratory volume at  $_1$ st second , P<0.05 significant

Table-2: HRV parameters in Study group and Control (n=30) in each group

Table-2. The parameters in Study group and Control (11–50) in each group					
D	Study group	Control	P-value		
Parameters	Mean± SD	Mean± SD			
TP ms <sup>2</sup>	852.98±702.37	1086.74±916.18	0.272		
LF ms <sup>2</sup>	165.31±175.60	195.97±175.04	0.501		
HF ms <sup>2</sup>	296.23±307.43	111.57±112.31	< 0.001		
LF n.u	37.34±8.12	63.67±7.33	< 0.001		
HF n.u	62.66±8.12	36.30±7.34	< 0.001		
LF-HF ratio	0.62±0.21	1.87±0.61	< 0.001		
SDNN(ms)	37.03±13.77	40.69±21.37	0.433		

TP-Total power, LF-Low frequency, HF- High frequency, SDNN-standard deviation of the averages of NN intervals, P<0.05 significant

Parameters		FEV1	FEV1/FVC	WHR
LF n.u	r- value	225	193	181
Lr n.u	p-value	.231	.306	.338
HF n.u	r- value	.225	.193	.181
HF II.U	p-value	.232	.307	.339
LF-HF RATIO	r- value	228	181	211
	p-value	.225	.337	.264

#### Discussion

The present study was conducted to assess autonomic modulations based on short term HRV indices and body composition parameters in newly diagnosed asthmatic patients . The objective of the study was to determine whether HRV and body composition correlate with the main spirometric indices. We observed parasympathetic dominance as the HF ms² & HF n.u, were significantly (P<0.001) higher in asthmatic patients as compared to control . Our study is in accordance with Gupta et al who reported an enhanced central vagal tone, as exemplified by as high HF index of HRV in asthmatics. Similar findings were also reported by other studies. [15, 16] Conversely, some studies were unable to prove parasympathetic dominance in patients with bronchial asthma .

We found that TP ms² & LF ms² were lower in asthmatics and showed no significant (P>0.05) difference. However LF n.u, which reflects the fluctuation in sympathetic tone of HRV and LF-HF ratio, a marker of sympatho-vagal balance, were significantly low (P<0.01) in the asthmatic patients when compared with control group.

The results of our present study are consistent with those observed by Garrard et al. [17] and Pichon et al.[18] who also reported low sympathetic activity represented by LF n.u. in asthmatics as compared to controls. On the contrary, a study reported that there was no significant difference in the low frequency band amplitude and LF-HF ratio between the controls and the asthmatic children. [19]

In our study SDNN, which reflects all cyclic components of the variability in RR intervals, was lower in case but this was statistically insignificant. Kazuma et al examined the circadian rhythm of parasympathetic nervous function in asthmatic children and found that SDNN was lowest in the severe asthma group. A study has also documented a circadian pattern in the components of HRV, the LF component

having minimal power and the HF component having maximal power during the nocturnal period .[20] Vagal tone increases to the heart at night and and as changes in cardiac vagal activity reflect changes in bronchomotor tone this has led to the hypothesis that vagal activity may be important in the pathogenesis of nocturnal asthma .[21]

In the present study, BMI was within normal range in both groups. There was no significant difference (P>0.05) in BMI & body fat% between 2 groups. However, WHR, was found to be significantly higher (P<0.001) in asthmatics as compared to controls . A study suggested that obesity and overweight are linked with allergic diseases such as rhinitis and asthma, probably because of the immunological effects of adipose tissue.

Autonomic nervous system plays a central role in modifying energy expenditure and body fat content; however, the extent to which the ANS contributes to obesity remains uncertain. [22, 23] A disordered homeostatic mechanism may promote excessive storage of energy which may be promoted by decreasing sympathetic activity, while defending against weight gain by decreasing parasympathetic activity. [24]

In our study, LF n.u. & LF-HF ratio were negatively correlated while HF n.u. was positively correlated with FEV1, FEV1/FVC & WHR and this was not statistically significant. Behera et al also found positive correlations with the HF component and FEV<sub>1</sub>, and negative correlations with the LF component and peak expiratory flow in smokers.[25] A study reported that HF norm correlates negatively and LF norm correlates positively with both weight and BMI. In addition LF/HF correlates positively with weight but not BMI in healthy subjects. [26]

#### Conclusion

Our study provides the evidence of a relationship between WHR and asthma. We observed altered

autonomic activity with parasympathetic dominance which leads to an imbalance in sympatho-vagal interplay an essential feature of bronchial asthma. So, it can be stated without reservation from our study, that HRV is an absolutely safe and useful non-invasive method for assessing autonomic modulation in numerous disease states including bronchial asthma which will lead to better symptomatic control and reduction in number of asthmatic attacks. We have not evaluated gender difference as the sample size was small.

#### References

- 1. Wang T, Guan C, McLimont J S, Gershon AS. What is the lifetime risk of physician-diagnosed asthma in Ontario, Canada? Am J Respir Crit Care Med 2010; 181:337-343.
- **2.** Asher M I, Anderson H R, Stewart A W.et al Worldwide variations in the prevalence of asthma symptoms: International Study of Asthma and Allergies in Childhood (ISAAC). Eur Respir J 1998; 12: 315–335.
- **3.** Masoli M, Fabian D, Holt S, Beasley R. Global Initiative for Asthma (GINA) Program: The global burden of asthma: executive summary of the GINA Dissemination Committee report. Allergy 2004; 59(5): 469-478.
- **4.** Kallenbach JM, Webster T, Dowdeswell R, Reinach SG, Millar RN, Zwi S. Reflex heart control in asthma Evidence of parasympathetic overactivity. Chest 1985; 87(5): 644-648.
- **5.** Nadel J, Barnes P. Autonomic regulation of the airways. Ann. Rev. Med 1984; 35:451–467.
- **6.** Gupta J, Dube A, Singh V, Gupta RC. Spectral analysis of heart rate variability in bronchial asthma patients. Indian J. Physiol. Pharmacol 2012; 56(4): 330-336.
- **7.** Lutfi MF. Autonomic modulations in patients with bronchial asthma based on short-term heart rate variability. Lung India 2012; 29 (3):254–258.
- 8. Garcia-Araújo AS, Pires Di Lorenzo VA, Labadessa IG, et al. Increased sympathetic modulation and decreased response of the heart rate variability in controlled asthma. J Asthma 2014; 31: 1–8.
- **9.** Cabiddu R, Aletti F, Duarte Souza V, et al. Cardiorespiratory coupling during sleep in difficult-to-control asthmatic patients. Eng Med Biol Soc. 2012;;3652–3655.
- **10.** Ciprandi G, Pistorio A, Tosca M, Ferraro MF, Cirillo I. Body mass index, respiratory function and bronchial hyper-reactivity in allergic rhinitis

- and asthma. Respiratory Medicare 2009; 3(2): 289-295.
- **11.** Backett WS, Jacobs DR Jr, Yu X, Iribarren C, William OD. Asthma is associated with weight gain in females but not males independent of physical activity. Am J Respir Critical Care Med 2001; 164(11): 2045-506.
- **12.** Adam M, Imboden M, Schaffner E, et al. The adverse impact of obesity on heart rate variability is modified by a NFE2L2 gene variant: International Journal of Cardiology 228 .2017; 341–346.
- **13.** American Thoracic Society. Standardization of spirometry: 1987 update. Am Rev Respir Dis 1987; 136:1285–1298.
- **14.** Task Force of the European Society of Cardiology and the North American Society of Pacing and Electrophysiology Heart rate variability; Standards of measurements, Physiological interpretation and clinical use. Circulation 1996; 93: 1043-1065.
- **15.** Kazuma N, Shirase E, Matsuoka I, et al. Circadian rhythm of parasympathetic nervous function in asthmatic children Arerugi. 1998; 47(12):1248-1251.
- 16. Ostrowska –Nawarcyz L, Wronski W, Blaszczyk J, Buiczyliko, Nawarcyz T. The heart rate variability analysis in youth and children with asthma. Pol Merkur Lekarshi 2006; 20(118): 399-403.
- **17.** Garrard CS, Seidler A, McAlpine LE, Gordon D. Spectral analysis of heart rate variability in bronchial asthma. Clin Auton 1992; 2(2): 105-111.
- **18.** Pichon A, Bisschop CD, Diaz V, Denjean A. Parasympathetic airway response and heart rate variability before and at the end of methacholine challenge. Chest 2005; 127: 23–29.
- **19.** Fujii H, Fukutomi O, Inoue R, et al. Autonomic regulation after exercise evidenced by spectral analysis of heart rate variability in asthmatic children. Ann Allergy Asthma Immunol 2000; 85: 233–237.
- 20. Lombardi G, Sandrone A, Mortara M T, LaRovere E, et al. Circadian variation of spectral indices of heart rate variability after myocardial infarction Am Heart J 1992;123 (6): 1521-1529.
- **21.** Morrison J F, Pearson S B, Dean H G. Parasympathetic nervous system in nocturnal asthma. Br Med J (Clin Res Ed) 1988: 296 (6634):1427-1429.

- 22. Vanderlei L, Pastre C, Júnior Ismael Freitas F, Fernandes M. Analysis of cardiac autonomic modulation in obese and eutrophic children. Clinics (Sao Paulo) 2010; 65(8): 789-792.
- Romer G. Heart autonomic function overweight adolescents. Indian Pediatrics 2005; 42(5): 464-469.
- 24. Peterson H, Rothschild M, Weinberg C, et al. Body fat and the activity of the autonomic
- 23. Guízar J, Ahuatzin R, Amador N, Sánchez G,
- nervous system. N Engl J Med 1988; 318: 1077-1083.
- 25. Behera JK, Sood S, Kumar N, Sharma K, Mishra R, Roy PS. Heart rate variability and its correlation with pulmonary function test of smokers. Heart Views 2013; 14: 22-25.
- 26. Lutfi MF. Relationship of height weight and body mass index to heart rate variability. Sudan Med J 2011; 47(1):14-19.

**Source of Support:Nil Conflict of Interest: Nil**