

Correlation of Breath Holding Time with Spirometry in Health and Disease – A Cross Sectional Study

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Abstract:

Background: Spirometry is the most widely used and an important method of evaluating the pulmonary functions. Although an accurate and easy method of evaluating the pulmonary function, spirometry carries numerous disadvantages. The important limitations of spirometry include, requirement of a machine for performing spirometry, trained personnel to perform, patient effort and understanding to perform, equipment calibration, risk of transmission of infection, etc. Hence, there was a need for a technique which is simple, easy to perform, non-machine, non-technician dependent, and equivalent to spirometry. There are numerous bedside spirometry techniques. One among them is the Single Breath Holding Time, which measures the maximum time to which a person can hold his breath after maximal inspiration. Hence, an attempt was made to see the correlation of single breath holding time with spirometry, so that Single Breath Holding Time can be used as an alternative to spirometry in resource poor settings.

Aims and Objectives: To determine the correlation of breath holding time with standard measures, post bronchodilator FEV₁, FVC and FEV₁/FVC ratio, so that breath holding test can be taken as a non-machine, non-technician dependent, bedside surrogate test for lung function test.

Methods: It is a cross sectional study from July 2021 to June 2022 done at Government Stanley Medical College, Chennai and Government Hospital of Thoracic Medicine, Tambaram Sanatorium. 175 cases, who need to undergo spirometry for any clinical indication were included in the study. Single breath holding time was also done for those patients and its correlation with spirometry is analyzed in the study.

Results: The 175 patients were divided into normal, obstructive and restrictive, based on Spirometry. The Breath Holding Time showed strong correlation with FEV₁, FVC and FEV₁/FVC ratio in all three categories (p value < 0.005) - normal, obstructive and restrictive patients. Single Breath Holding Time is less than 16 seconds in patients with both obstructive and restrictive pattern, whereas Single Breath Holding Time is more than 25 seconds in normal individuals.

Conclusion: Single Breath Holding Time shows strong correlation with FEV₁, FVC and FEV₁/FVC. Hence, Single Breath Holding Time can be used as a simple bedside, non-machine, non-technician dependent alternative to spirometry in resource poor settings. It can also be used as a point of care test, to decide on further work-up of the patient.

Keywords: breathe hold time, spirometry, Copd, asthma, pulmonary function test, bedsidepft.

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Introduction

Pulmonary function testing is an important aspect of evaluating the respiratory diseases. Spirometry is the most commonly used method for assessing the pulmonary ventilatory functions. Spirometry measures the air flow during inspiration and expiration. Hence, spirometry is used for measuring the dynamic lung volumes and capacities. Though very useful in arriving at a diagnosis, this spirometry has numerous limitations. Few practically troublesome limitations of spirometry

include requirement of an equipment with proper software for its interpretation, technician to perform the test, proper patient understanding, risk of spread of infections, adequate training for the operating technician, and difficulty in carrying it to places like ICU or bedside of the patient. Hence, there was a need to find a test that could overcome the shortcomings of spirometry. Then came the concept of bedside spirometry. There are numerous bedside spirometry techniques like candle blowing,

single breath count, single breath holding time, etc. Among the various bedside spirometry tests, single breath holding time is one of the simple techniques which measures the maximal time a person can hold a breath after deep inspiration. Many studies are done all over the world comparing the various bedside pulmonary function tests with standard spirometry. Here, we have also attempted to see the correlation of single breath holding time with spirometry parameters of FEV₁, FVC and FEV₁/FVC. If there is a good correlation of breath holding time with spirometry, then it can be used as a simple, non-machine, non-technician dependent, bedside alternative to spirometry. Owing to its simplicity and cost effectiveness, Breath holding time has been suggested as reasonable alternative to spirometry in resource poor settings with the added advantages of independence from technician and machine. This study aims to verify this claim.

Aims & Objectives

The aim of the study is to determine the correlation of breath holding time with standard spirometry (post bronchodilator FEV₁, FVC and FEV₁/FVC ratio), so that breath holding time can be used as a non-machine, non-technician dependent, bedside, surrogate test for lung function test.

Materials and Methods

Type of Study: Cross Sectional Study

Study Population: Patients attending OPD in Government Stanley Medical College and Government Hospital of Thoracic Medicine, Tambaram Sanatorium, who need to undergo Spirometry for any clinical indication

Sample size: 175

Study Duration: 1 year

Inclusion Criteria:

All patients above the age of 18 years who need to undergo Spirometry for any clinical indication

Exclusion Criteria:

- Patients who are unwilling to participate in the study
- Patients whose spirometry are not acceptable and valid as per ATS standard
- Patients who have absolute or relative contra-indication for Spirometry
- Myocardial Infarction within the last one month
- Conditions which can lead to suboptimal test
- Chest, facial, oral or abdominal pain
- Stress Incontinence
- Dementia and confusion

- Active pulmonary tuberculosis
- Acute exacerbation of COPD
- Acute Severe Asthma
- Hemoptysis
- Post Covid patients

Ethical Considerations:

Study was approved by Institutional Ethical Committee. Informed written consent were obtained from all the study participants. Confidentiality of the participants in the study are maintained

Data Collection Tools: All study details are entered in a structured study proforma.

Methodology:

- 175 cases, who needed to undergo Spirometry for clinical indications.

Spirometry was performed in sitting position, with a nose clip attached.

- The ATS guidelines for Spirometry were followed.

Broncho dilation was achieved using a pMDI Salbutamol 400microgram

- Best of three successive test readings was taken as final result and the primary values, i.e., post bronchodilator forced vital capacity [FVC], forced expiratory volume in the first second [FEV₁] were noted.
- Then single breath holding time was performed by all participants within 3 minutes to avoid the waning effect of bronchodilator.
- Participants were asked to hold the breath after a normal tidal volume breath, till the breaking point.
- Breath hold test manoeuvre was performed 3 times with a gap of 5 minutes and the best of the 3 values were included for analysis.

Results

The collected data were analysed with IBM SPSS Statistics for Windows, Version 23.0. (Armonk, NY: IBM Corp). To describe about the data descriptive statistics frequency analysis, percentage analysis was used for categorical variables and the mean & S.D were used for continuous variables.

To find the significant difference in the multivariate analysis the one-way ANOVA with Tukey's Post-Hoc test was used. To find the significance in categorical data Chi-Square test was used.

In all the above statistical tools the probability value .05 is considered as significant level.

Table 1: Age distribution of our study population

Age distribution		
	Frequency	Percent
18 - 20 yrs	8	4.6
21 - 30 yrs	19	10.9
31 - 40 yrs	32	18.3
41 - 50 yrs	38	21.7
51 - 60 yrs	49	28.0
61 - 70 yrs	23	13.1
71 - 80 yrs	6	3.4
Total	175	100.0

Table 2: PFT pattern distribution of our study population

PFT pattern		
	Frequency	Percent
Normal	79	45.1
Obstructive	60	34.3
Restrictive	36	20.6
Total	175	100.0

Table 3: Comparison of Age with PFT pattern by Pearson's Chi-Square test

			PFT pattern			Total	χ^2 - value	p-value
			Normal	Obstructive	Restrictive			
Age	18 - 20 yrs.	Count	7	1	0	8	30.907	0.002 **
		%	8.9%	1.7%	0.0%	4.6%		
	21 - 30 yrs.	Count	14	1	4	19		
		%	17.7%	1.7%	11.1%	10.9%		
	31 - 40 yrs.	Count	15	12	5	32		
		%	19.0%	20.0%	13.9%	18.3%		
	41 - 50 yrs.	Count	20	9	9	38		
		%	25.3%	15.0%	25.0%	21.7%		
	51 - 60 yrs.	Count	14	21	14	49		
		%	17.7%	35.0%	38.9%	28.0%		
	61 - 70 yrs.	Count	9	12	2	23		
		%	11.4%	20.0%	5.6%	13.1%		
	71 - 80 yrs.	Count	0	4	2	6		
		%	0.0%	6.7%	5.6%	3.4%		
Total		Count	79	60	36	175		
		%	100.0%	100.0%	100.0%	100.0%		

** Highly Statistical Significance at $p < 0.01$ level**Table 4: Comparison of Gender with PFT pattern by Pearson's Chi-Square test**

			PFT pattern			Total	χ^2 value	p-value
			Normal	Obstructive	Restrictive			
Gender	Female	count	19	18	17	54	6.255	0.044
		%	24.1%	30.0%	47.2%	30.9%		
	Male	count	60	42	19	121		
		%	75.9%	70.0%	52.8%	69.2%		
	Total	count	79	60	36	175		
		%	100%	100%	100%	100%		

Statistical significance at $p < 0.05$ level

Table 5: Comparison of Smoking History with PFT pattern by Pearson's Chi-Square test

				PFT pattern				
			Normal	Obstructive	Restrictive	Total	x 2-value	p-value
Smoking History	No	count	79	19	36	134	102.627	0.0005**
		%	100%	31.7%	100%	76.6%		
	Yes	count	0	41	0	41		
		%	0.0%	68.3%	0.0%	23.4%		
Total		count	79	60	36	175		
		%	100%	100%	100%	100%		
Highly Statistical significance at p< 0.01 level								

Highly Statistical significance at $p < 0.01$ level**Table 6: Comparison of Biomass Exposure with PFT pattern by Pearson's Chi-Square test**

			PFT pattern			Total	χ^2 - value	p-value
			Normal	Obstructive	Restrictive			
Biomass Exposure	No	Count	77	52	36	165	10.132	0.006 **
		%	97.5%	86.7%	100.0%	94.3%		
	Yes	Count	2	8	0	10		
		%	2.5%	13.3%	0.0%	5.7%		
Total		Count	79	60	36	175		
		%	100.0%	100.0%	100.0%	100.0%		

** Highly Statistical Significance at $p < 0.01$ level**Table 7: Comparison of Chest X-ray with PFT pattern by Pearson's Chi-Square test**

			PFT pattern			Total	χ^2 - value	p-value
			Normal	Obstructive	Restrictive			
Chest X-ray	B/L HI	Count	0	45	0	45	119.531	0.0005 **
		%	0.0%	75.0%	0.0%	25.7%		
	Interstitial opacities	Count	0	0	1	1		
		%	0.0%	0.0%	2.8%	.6%		
	Normal	Count	79	15	35	129		
		%	100.0%	25.0%	97.2%	73.7%		
Total		Count	79	60	36	175		
		%	100.0%	100.0%	100.0%	100.0%		

** Highly Statistical Significance at $p < 0.01$ level**Table 8: Comparison of FEV1% with PFT pattern by One way ANOVA test**

Variable	PFT pattern	N	Mean	SD	F-value	p-value
FEV1%	Normal	79	81.20	11.33	150.379	0.0005 **
	Obstructive	60	43.68	16.38		
	Restrictive	36	48.75	13.01		

** Highly Statistical Significant at $p < 0.01$ level**Table 9: Comparison of FVC% with PFT pattern by One way ANOVA test**

Variable	PFT pattern	N	Mean	SD	F-value	p-value
FVC%	Normal	79	84.14	11.44	62.366	0.0005 **
	Obstructive	60	64.22	20.72		
	Restrictive	36	50.86	14.29		

** Highly Statistical Significant at $p < 0.01$ level**Table 10: Comparison of FEV1/FVC% with PFT pattern by One-way ANOVA test**

Variable	PFT pattern	N	Mean	SD	F-value	p-value
FEV1/FVC%	Normal	79	94.09	5.29	263.217	0.0005 **
	Obstructive	60	66.25	10.04		
	Restrictive	36	93.17	6.88		

** Highly Statistical Significant at $p < 0.01$ level

Table 11: Comparison of SBHT with FEV1, FVC & FEV1/FVC by Pearson's Chi-Square

Correlations				
		FEV1	FVC	FEV1/FVC
SBHT	r-value	.754**	.632**	.536**
	p-value	.0005	.0005	.0005
	N	175	175	175

** . Correlation is significant at the 0.01 level.

Table 12: Comparison of SBHT with PFT pattern by One way ANOVA test

Variable	PFT pattern	N	Mean	SD	F-value	p-value
SBHT	Normal	79	31.92	5.31	371.990	0.0005 **
	Obstructive	60	13.43	3.40		
	Restrictive	36	14.67	3.30		

** Highly Statistical Significant at $p < 0.01$ level

Discussion

The main aim of the study is to see the correlation of breath holding time with spirometry parameters in health and disease. Thus, 175 patients having any indication for spirometry are included in the study.

In our study, 49% of the study population was in the 51-60 years age group. 69% of the study population were males and 31% were females. Depending upon the PFT pattern, the study population was divided into normal, obstructive pattern and restrictive pattern. In our study, 79 patients (45.1%) showed normal spirometry pattern, 60 patients (34.3%) showed obstructive spirometry pattern and 36 patients (20.6%) showed restrictive pattern on spirometry.

The breath holding time was done for all the patients after spirometry. The mean breath holding time in normal patients was 31.92. Whereas the mean breath holding time in patients with obstructive pattern is 13.43 and the mean breath holding time in patients with restrictive pattern is 14.67. All the above obtained results are highly significant with a p value of 0.0005. Thus single breath holding time correlated significantly with spirometry.

The results of my study are similar to the study done by Vipin Aggarwal et al in 2018, which compared the Single Breath Holding Time with FEV1, FVC and PEFR. This study also showed similar results, which concluded that there was high significant correlation with post bronchodilator FEV1 and FVC, although there was low strength in cases of PEFR, particularly in patients with obstructive pattern.

Limitations

Single centre study Hospital based study Done in Indian population and hence a study population consisting of all ethnicity and races would help us in getting a better correlation

Conclusion

Thus, the study shows a significant correlation between single breath holding time and FEV1, FVC and FEV1/FVC. Breath holding time is reduced in patients with respiratory pathology, both obstructive and restrictive abnormalities. Whereas breath holding time is higher in normal individuals. Thus, single breath holding time can be used as a simple, bedside, non- machine, non-technician dependent test to assess if there is any respiratory abnormality, in resource poor settings. And it can also be used as the initial test in assessing the patients before spirometry is performed. Hence in resource poor settings, where spirometry is not available, single breath holding time can be used as a bed side surrogate test for spirometry.

References

1. Aggarwal V, Godbole G, Agawane S, Pophale H. Correlation of breath holding time with spirometry Amatya et al Nepal Medical College Journal 234 NMCJ test - An alternative non technician dependent surrogate test for spirometry. MedPulse Int'l J Med 2018; 5: 69-73.
2. American Thoracic Society/European Respiratory Society Task Force. Standardization of Lung Function Testing. Number 2: Standardization of spirometry. EurRespir J. 2005; 26:319-338
3. Schlegelmilch RM, Kramme R. Pulmonary Function Testing. In: Kramme R, Horffman K-P, Pozos R, editors. Springer Handbook of Medical Technology. Springer 2011: 95-118.
4. Calverley P. The clinical usefulness of spirometric information. Breathe 2009; 5: 214-20
5. Sharma J, Senjyu H, Williams L. Comparison of Chest Expansion Measurement in Clients with Ankylosing Spondylitis and Healthy Individuals. J Phys Ther Sci 2004; 15: 47-51.
6. Ruppel GL. What Is the Clinical Value of Lung Volumes? Respir Care 2012; 57: 26-38.
7. Bockenhauer SE, Chen H, Julliard KN, Weedon J. Measuring thoracic excursion: reli-

- ability of the cloth tape measure technique. *J Am Osteopath Assoc* 2007; 107: 191–6
8. Lanza F de C, Camargo AA de, Archija LRF, Selman JPaR, Malaguti C, Corso SD. Chest Wall Mobility Is Related to Respiratory Muscle Strength and Lung Volumes in Healthy Subjects. *Respir Care* 2013; 58: 2107–12.
 9. Parkes MJ. Breath-holding and its breakpoint. *Exp Physiol* 2006; 91:1–15.
 10. Lindholm P, Lundgren CEG. The physiology and pathophysiology of human breath-hold diving. *J Appl Physiol* 2009; 106: 284–92.
 11. Malaguti C, Rondelli RR, de Souza LM, Domingues M, Dal Corso S. Reliability of chest wall mobility and its correlation with pulmonary function in patients with chronic obstructive pulmonary disease. *Respir Care* 2009; 54: 1703–11.
 12. Taskar V, Clayton N, Atkins M, Shaheen Z, Stone P, Woodcock A. Breath-holding time in normal subjects, snorers, and sleep apnea patients. *Chest* 1995; 107: 959–62.
 13. Viecili RB. Real-Time Measurement of Maximal Voluntary Breath-Holding Time in Patients with Obstructive Ventilatory Defects and Normal Controls. *J Pulm Respir Med* 2013; 2: 2–4.
 14. Crapo RO. Pulmonary function testing. *N Engl J Med* 1994; 331: 25–30.
 15. Palaniyandi AK, Natarajan M, Chockalingam A, Karthick R, Professor A, Author C. Even a single breath counts. *IOSR J Dent Med Sci* 2017; 16: 70–2.
 16. Nepal GB, Das PKL, Bhaila A. Spirometric evaluation of pulmonary functions of medical students in Nepal. *Asian J Med Sci* 2014; 5: 82–6.
 17. Richards JA. Office spirometry - Indications and limitations. *South African Fam Pract* 2006; 48: 48–51.
 18. Viecili RB, Silva DeR, Sanches PRS, Muller AF, Silva DP da, Barreto SSM. Real-Time Measurement of Maximal Voluntary Breath-Holding Time in Patients with Obstructive Ventilatory Defects and Normal Controls. *J Pulm Respir Med* 2013; 2: 2–4.
 19. Morris JF, Koski A, Breese JD. Normal values and evaluation of forced end-expiratory flow. *Am Rev Respir Dis*. 1975 Jun; 111(6):755–762.
 20. BERNSTEIN L, KAZANTZIS G. The relation between the fast vital capacity curves and the maximum breathing capacity. *Thorax*. 1954 Dec; 9(4):326–339
 21. Jones JS, Renzetti AD, Jr, Mitchell MM. The maximal breathing capacity in extrathoracic airway obstruction. *Am Rev Respir Dis*. 1972 Dec; 106(6):925–927.
 22. Snider GL. Spirometric evaluation of ventilatory function. *JAMA*. 1966 Sep 26; 197(13):1095–1095.
 23. Weng TR, Levison H. Standards of pulmonary function in children. *Am Rev Respir Dis*. 1969 Jun; 99(6):879–894
 24. GAENSLER EA. Analysis of the ventilatory defect by timed capacity measurements. *Am Rev Tuberc*. 1951 Sep; 64(3):256–278.
 25. Morris JF, Temple WP, Koski A. Normal values for the ratio of one-second forced expiratory volume to forced vital capacity. *Am Rev Respir Dis*. 1973 Oct; 108(4):1000–1003.