



## Accuracy of Pulmonary Asthma Score in Predicting Severity of Asthma Exacerbation in Mild and Moderate Asthma

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### Abstract

**Objectives:** The objective of the study was to determine the accuracy of pulmonary asthma score in predicting severity of asthma exacerbation and comparison of efficacy between levosalbutamol and ipratropium combination over levosalbutamol nebulization in reversing airflow obstruction and improving oxygenation, evaluated using the pulmonary asthma score, SaO<sub>2</sub>, and PEFR in mild and moderate asthma.

**Methods:** A prospective, randomized, study was performed in RMMCH pediatric emergency department. Children between 6 and 12 years of age who presented with mild to moderate asthma exacerbations were included in the study. They were randomly allocated into two different groups: one nebulized with levosalbutamol alone and another with addition of ipratropium bromide to levosalbutamol. Baseline Peak expiratory flow rate and final absolute values or change from baseline 60–120 minutes after the inhalation are measured. Patients were evaluated using the pulmonary asthma score<sup>4</sup>.

**Results:** After treatment there is improvement in the mean pulmonary asthma scores and PEFR percentage in A+B group than A group, but it is not statistically significant (p value >0.05). There is statistically significant improvement in pulmonary asthma score and PEFR in each of the groups before and after nebulization and pulmonary asthma score has a sensitivity of 66.7% and 65.6% in diagnosing severity of asthma in relation to PEFR.

**Keywords:** Levosalbutamol, Ipratropium bromide, peak expiratory flow rate, pulmonary asthma score (PAS).

### Introduction

The administration of  $\beta_2$  agonists is the standard treatment for mild to moderate asthma crisis in emergency departments<sup>(1,2)</sup>. Current guidelines suggested the use of a combination of inhalation of  $\beta_2$  agonists plus anticholinergics for pediatric patients with acute severe or life-threatening asthma in the emergency setting. But its role in mild to moderate asthma exacerbation is still

inconclusive. It is practically difficult to assess lung function in children under 5 years of age because most lung functions in diagnosing and predicting severity of asthma are subjective and needs individual understanding and cooperation. Recent studies of acute asthma management in the emergency department have focused on objective assessment of the severity of asthma and rapid provision of effective treatment. Care is guided by

the clinical evaluation, including a directed history and a physical examination focused on key warning signs of rapid deterioration or severe exacerbation and measurement of oxygen saturation by pulse oximetry. Various clinical scoring systems are effective in predicting the severity of asthma in emergency room. In our study PAS is used in assessing the severity of asthma in relation to PEF.

### Materials and Methods

A prospective, randomized, study was performed in RMMCH pediatric emergency department between the months of August 2016 and August 2018. The study was previously approved by the hospital ethics committee. Children between 6 and 12 years of age who presented with mild to moderate asthma exacerbations were included in

the study. They were randomly allocated into two different groups: one nebulized with levosalbutamol alone and another with addition of ipratropium bromide to levosalbutamol. Baseline Peak expiratory flow rate and final absolute values or change from baseline 60–120 minutes after the inhalation were measured). Because the peak bronchodilator effect after the administration of anticholinergics occurs within 1–2 hours, it is reasonable to expect significant improvement during this time. Patients were evaluated using the pulmonary score<sup>3</sup>. The oxygen saturation (SaO<sub>2</sub>) was measured using a pulse oximeter. Peak expiratory flow rate was performed using a portable peak flow meter. The predicted PEF was adjusted for sex and height and the percentages of the PEF before and after nebulization was obtained.<sup>4,5</sup>.

**Table:** Pulmonary Asthma Score

Component score	0	1	2	3
Respiratory rate	14-20	21-26	27-30	>31
Oxygen saturation	>98% at room air	95-97% on room air	90-94% on room air	<90% on room air
Auscultation	Normal breath sounds	End expiratory wheezing	Expiratory wheezing	Inspiratory and expiratory wheeze
Retractions	None	Intercostal	Intercostal and substernal	Intercostal and substernal and supraclavicular
Dyspnea	Speaks in complete sentences	Speaks in short sentences, coos, and babbles	Speaks in partial sentences, short cry	Speaks in single words, short phrase/grunting

The severity of the asthma crisis was classified according to the pulmonary asthma score as mild (score 5–7 and PEF > 70% of the predicted value according to the age and sex; moderate (score 8–11) and PEF 50–70% or FEV<sub>1</sub> 60–80% of the predicted value and severe (score 12–15) and PEF < 50% or FEV<sub>1</sub> < 60% of the predicted value. The PAS was the main criteria to rank severity of asthmatic crisis.

### Results

Group 1: A-levosalbutamol

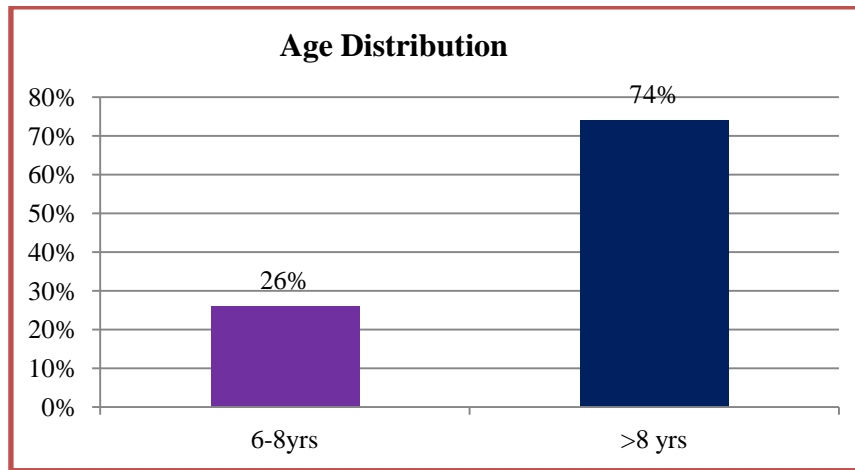
Group 2: A+B - levosalbutamol + ipratropium bromide

**Table 1:** Name of the Medication Used

	Frequency	Percent
A	25	50.0
A+B	25	50.0
Total	50	100.0

**Table 2:** Age distribution among study population

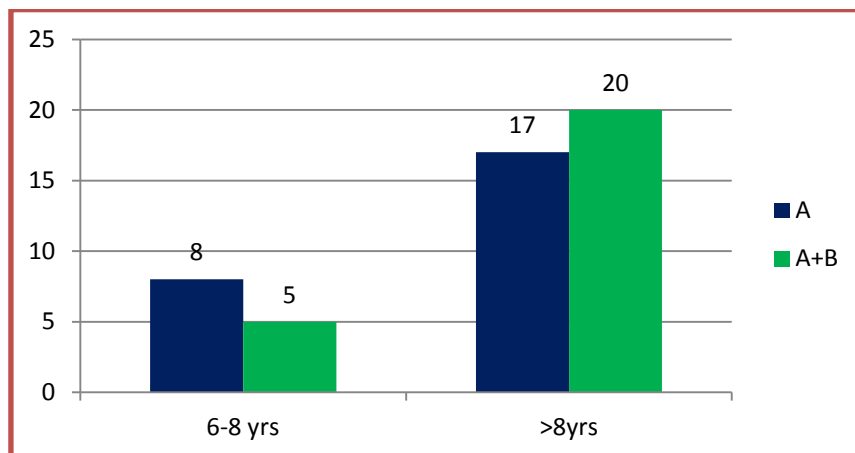
	Frequency	Percent
6-8yrs	13	26.0
>8 yrs	37	74.0
Total	50	100.0



Comparison of age in both groups

**Table 3:** Age Range and Name of the Medication Used Cross tabulation

Age range	Name of the Medication Used		Total	P value
	A	A+B		
6-8 yrs	8	5	13	0.260
>8yrs	17	20	37	
Total	25	25	50	



**Table 4:** Sex distribution among study population

	Frequency	Percent
Male	26	52.0
Female	24	48.0
Total	50	100.0

**Table 5:** Pulmonary asthma score before nebulization in both groups

PAS range	Name of the Medication Used		Total	P value
	A	A+B		
Mild	10	8	18	0.509
Moderate	15	17	32	
Total	25	25	50	

**Table 6:** Distribution of PEFR before nebulization

	Frequency	Percent
>70%	18	36.0
50 -70%	32	64.0
Total	50	100.0

**Table 7:** Means of HR, RR, SPO2

	N	Mean	Std. Deviation
<b>Before</b>			
B-HR	50	130.84	23.191
B-RR	50	35.74	8.799
SPO2	50	94.5000%	3.01865%
<b>After</b>			
A-HR	50	128.40	23.181
A-RR	50	29.44	7.843
A-SPO2	50	96.2800%	2.57967%

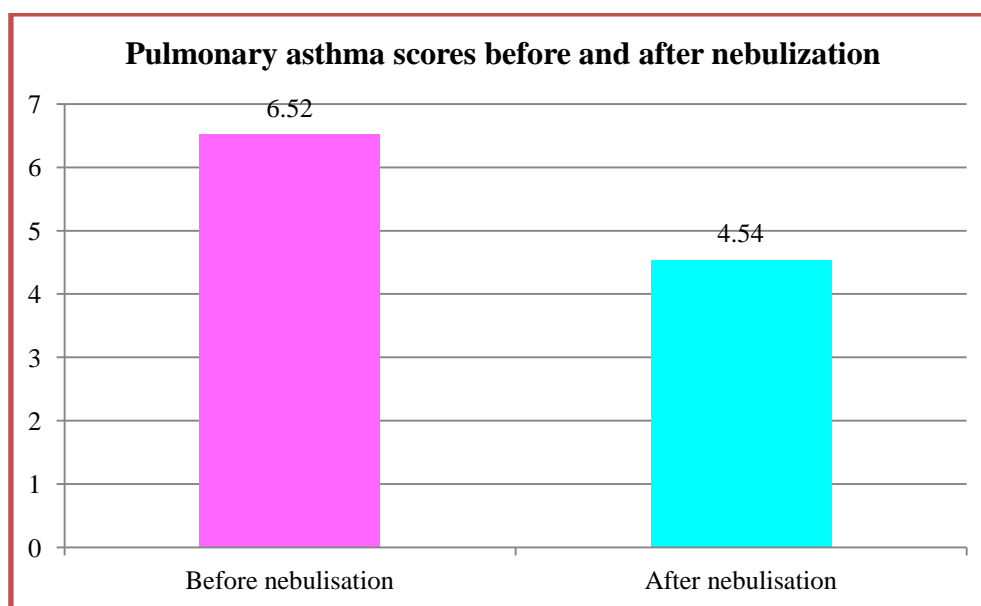
Comparison of PEFR before nebulization between the groups

**Table - 8:** Baseline PEFR and Name of the Medication Used Cross tabulation

		Name of the Medication Used		Total	P value
		A	A+B		
Pefr %	Mild	10	8	18	<b>0.384</b>
	Moderate	15	17	32	
Total		25	25	50	

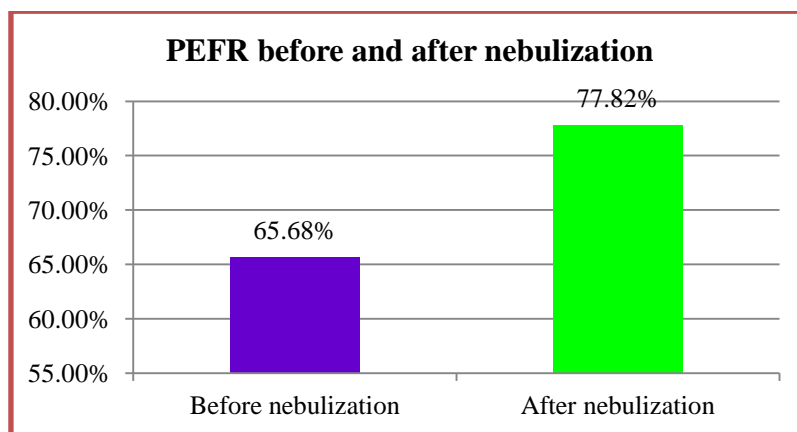
**Table -9:** Pulmonary asthma scores before and after nebulization

	Mean	N	Std. Deviation	T value	P value
Before Pulmonary asthma score	6.52	50	2.443	15.697	<0.05 *
After PAS	4.54	50	2.451		

**Table 10:** PEFR before and after nebulization

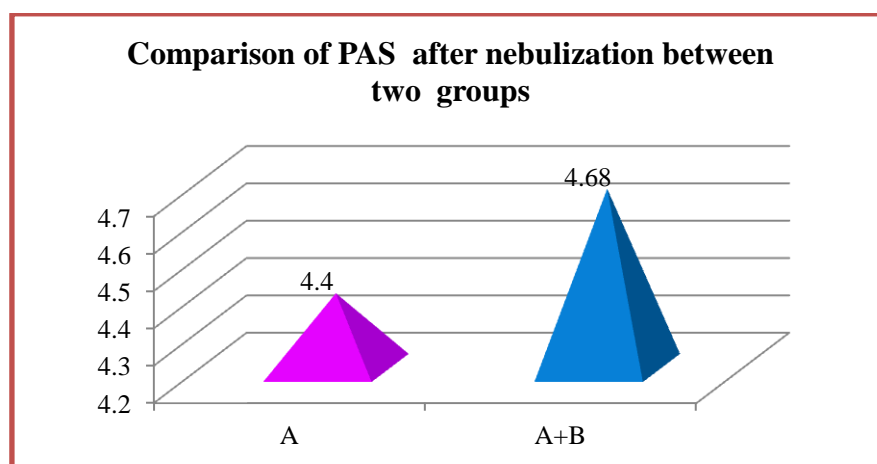
		Mean	N	Std. Deviation	T value	P value
Pair 1	% of Predicted PEFR baseline	65.6800%	50	9.29437%	-15.588	<0.05 *
	% of Predicted PEFR after neb	77.8200%	50	7.93003%		

There significant improvement in PAS and PEFR after nebulization in both the treatment regimes.



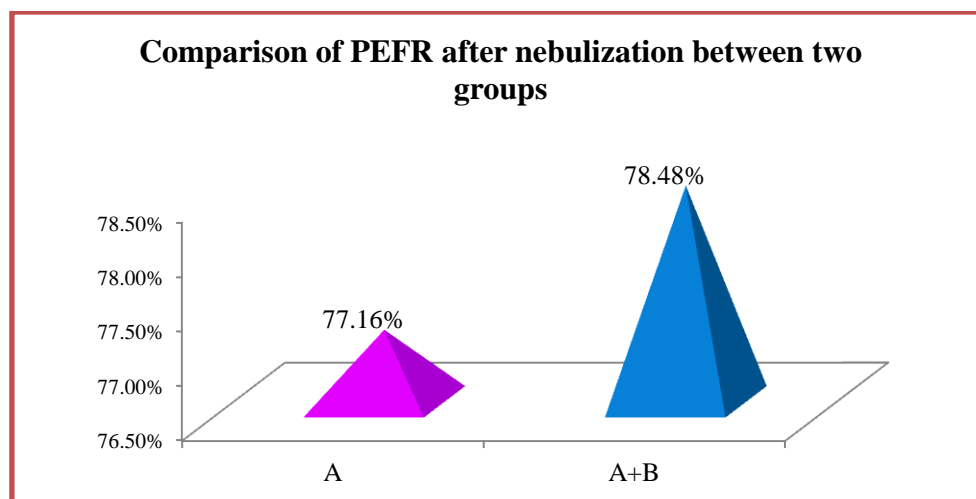
**Table 11:** Comparison of PAS after nebulization between two groups

Name of the Medication Used		N	Mean	Std. Deviation	T value	P value
PAS	A	25	4.40	2.449	-0.40	0.691
	A+B	25	4.68	2.495		



**Table 12:** Comparison of PEFR after nebulization in two groups

	Name of the Medication Used	N	Mean	Std. Deviation	T value	P value
% of Predicted PEFR after neb	A	25	77.1600%	8.80663%	-0.585	0.562
	A+B	25	78.4800%	7.06588%		



Chi-Square Tests			
	Value	df	Asymp. Sig. (2-sided)
Pearson Chi-Square	17.645 <sup>a</sup>	2	.000
Likelihood Ratio	20.428	2	.000
Linear-by-Linear Association	16.589	1	.000
N of Valid Cases	50		
a. 1 cells (16.7%) have expected count less than 5. The minimum expected count is 3.60.			

**Sensitivity :**

66.7%

**Specificity :**

65.6%

### Discussion

Out of 50 children included in the study, 52 percent were males and 48 percent were females. 26 percent were less than 8 years of age and 74 percent were more than 8 years of age.

Both groups were matched for age, sex, oxygen saturation and asthma severity. Regarding baseline severity score, in group A it was 5-7 in 10 and 8-11 in 15 cases. In group A +B it was 5-7 in 7 and 8-11 in 18 cases.

Based on PEFR, out of 50 children, 36% had mild exacerbation and 64% had moderate exacerbation of asthma respectively.

Chi square test was used for statistical analysis, paired and unpaired T test were used for comparison between two groups.

There is statistically significant improvement in PAS and PEFR in each of the groups before and after nebulization and pulmonary asthma score has a sensitivity of 66.7% and 65.6% in diagnosing severity of asthma in relation to PEFR. After nebulization there is significant improvement in respiratory rate and oxygen saturation in both groups but there is not much difference between both groups.

After treatment there is a greater improvement in the mean PAS and PEFR percentage in A+B group than A group, but it is not statistically significant (p value >0.05). This may be attributed

to small sample size in both groups. Yet the effects of Ipratropium with levosalbutamol seems to be equal to the levosalbutamol alone group.

In a study of 125 children with severe asthma, by Quershi et al<sup>6</sup>, found that (FEV1) improved to a greater extent in children receiving salbutamol and ipratropium than in those receiving salbutamol placebo, but there is no effect on overall rates of hospitalization. In a subgroup analysis of children in whom FEV1 was less than 30 percent of the predictive value, the hospitalisation rate among those receiving the combination therapy was significantly lower than the rate with salbutamol alone.

In a study of 434 children by Schuh et al<sup>7</sup> with moderate and severe asthma exacerbation showed that the addition of ipratropium bromide had a significant effect on improvement of asthma score, but there was no significant improvement of PEFR.

In a study done in India by Sharma A et al<sup>8</sup>, in 2004 showed that frequent combined nebulization with salbutamol and ipratropium significantly improved percentage of PEFR starting at 30 minutes and lasting for 4 hours in 50 children (6-14 years) with moderate exacerbation

## Conclusion

Ipratropium combination therapy has a definitive role in severe asthma crisis. But its role in mild to moderate asthma exacerbation is still inconclusive. Statistical significance has not been obtained in our study due to relatively smaller number of cases in each group. PAS holds good in assessing the severity of asthma exacerbation and hence can be used for rapid clinical evaluation of acute asthma in emergency settings and also to assess the response to treatment and to decide on hospital admission

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