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#### **Review Article**

EFFECT OF AYUASMO CAPSULES (AYURVEDIC FORMULATION) IN THE MANAGEMENT OF TAMAKA SAWASA (BRONCHIAL ASTHMA) BY ASSESSMENT OF PULMONARY FUNCTION TESTS: A SINGLE CASE STUDY

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#### **ABSTRACT**

Introduction: - Bronchial Asthma is a Chronic Obstructive Pulmonary Disease affecting a large population across the world. In ayurveda, bronchial asthma can be classified under *Sawasa Roga. Tamaka Sawasa* is a subtype of *Sawasa* described as an independent disease having its own *Nidan* (etiology), *Samprapti* (pathophysiology), and *Chikitsa* (management). Main Clinical Findings- A female patient aged 37 years with a history of *Tamaka Sawasa* (bronchial asthma) for the last 7 years. The chief complaints of the patient were cough and dyspnoea since last 7 years. Assessment of Pulmonary Function Test through Spirometer at baseline was FEV1 of 50%. Diagnosis: - Bronchial Asthma. Interventions: - Ayuasmo capsule (Ayurvedic Formulation) - a combination of 14 ingredients (*Vasa, Kantakari* etc). Ayuasmo Capsules was administered in a dose of 2 Capsules twice a day for 3 months. No other medicine was being taken or given other than the above-mentioned protocol. Outcome: - Along with clinical improvement in symptoms of dyspnoea, wheezing, significant improvement was also observed in the lung capacity (FEV1 50% at baseline to FEV1 90% at the end of 3 months). Conclusion: Ayuasmo can be considered as an effective medication in the management of Bronchial Asthma.

Keyword: - Ayuasmo, Tamaka Swasa, FEV1

Case report

Introduction

In modern times, the prevalence of respiratory diseases has increased dramatically due to a combination of lifestyle factors and environmental pollution. Management of this condition is done with the use of Bronchodilators, anti-inflammatory and also corticosteroids in modern medicine. Ayurveda recommends the use of various herbs and formulations that have been proven to be scientifically useful in the management of Bronchial Asthma (*Tamak Swasa*).

A 37-year-old female came to O.P.D. of NIA Jaipur, India on September 28, 2018 (screening visit). The chief complaints of the patient were cough and dyspnoea since last 7 years. Auscultation of chest wheezing was present bilaterally. The patient had history of Bronchial Asthma since 7 years and was taking treatment for the same on and off. Written informed consent was taken from the patient. All laboratory related parameters including CBC, LFT, RFT, Lipid Profile, Urine (Routine and Microscopic) were within normal limits. Spirometery examination showed FEV1% as 50% (moderate case of asthma).

Subject was recruited in the study and was asked to take Ayuasmo capsules in a dose of 2 Capsules twice a day for 3 months. Subject was regularly followed up every 15 days when clinical examination was done. At the interval of every 30 days assessment was also done on Spirometer for changes in the lung capacity. At the end of the study all the laboratory parameters checked at baseline were assessed.

Drug:

Ayuasmo (Ayurvedic Formulation) capsule is a combination of 14 ingredients which have been used in Ayurveda in the management of various types of respiratory diseases. Ayuasmo capsules support the respiratory system and strengthen the lungs' and bronchial tubes' fragile tissues.

Ingredients: Vasa<sup>[2]</sup> Extract{Adhatoda vasica} (50mg), Yastimadhu<sup>[3]</sup> Extract{Glycyrrhiza glabra} (50mg), Kulanjan<sup>[4]</sup> Extract{Alpinia galangal} (30 mg), Puskaramula<sup>[5],[6]</sup> Extract{Inula racemosa} (30 mg), Aamalaki<sup>[7],[8]</sup> Extract{Phyllanthus emblica} (30 mg), Katphala<sup>[9]</sup> Extract{Myrica Nagi}(30mg), Bala<sup>[10]</sup> Extract {Sida cordifolia} (30mg), Bharangi<sup>[11]</sup> Extract {Clerodendrum indicum} (25 mg), Shati<sup>[12]</sup> Extract{Hedychium spicatum} (25mg), Karkatashringi<sup>[13]</sup> Extract{Pistacia integerrima} (25 mg), Kutki<sup>[14]</sup> Extract {Picrorrhiza kurroa} (20mg), Shunthi<sup>[15]</sup> Extract {Zingiber officinale} (20mg), Tulsi<sup>[16],[17]</sup> Extract{Occimum sanctum} (15mg), Talispatra<sup>[18]</sup> Extract {Abies webbiana} (10mg).

Indications: 1. Bronchial asthma 2. Cold, Cough 3. Rhinitis 4. Bronchitis 5. Sinusitis

Dosage: 2 Capsules twice a day with warm water for 3 months.

#### **Results:**

Assessment of effect of consumption of Ayuasmo Capsules was done on both clinical parameters and also through Pulmonary Function tests (Spirometry).

# i) Assessment of FEV1 over a period of 90 days:

FEV1 is the amount of air, which can be forcibly exhaled from the lungs in the first second of a forced exhalation.

At Day 0 (Baseline visit) FEV1 predicted at day 0 (2.2 L), best effort by patient (1.1 L), % Prediction (50%) signifying moderate case of asthma. At Day 30 there was a significant improvement where the FEV1 predicted was (2.2 L) best effort by patient (1.8 L) % Prediction (81%). At Day 60 further improvement was observed with FEV1 predicted at day 60 (1.8 L) best effort by patient (1.59 L) % Prediction (70%) signifying moderate case of asthma. At the end of the study i.e. Day 90, FEV1 predicted was (2.2 L) best effort by patient (2.04 L) % prediction (90%) signifying only mild Asthma. Refer Table 1.1.

# ii) Assessment of FVC over a period of 90 days.

**FVC** Forced vital capacity (L) the amount of air that can be forcibly exhaled from your lungs after taking the deepest breath.

At Day 0 (Baseline visit) FVC predicted at day 0 (2.84 L) best effort by patient (1.83 L) % Prediction (64%). At the end of day 30 significant improvement was observed with FVC predicted at day 30 (2.84 L) best effort by patient (2.72 L) % prediction (96 %). Further at day 60 improvement continued with FVC predicted at day 60 (2.84 L) best effort by patient (2.27 L) % Prediction (80 %). At the end of the study i.e. day 90 was much better than Day 0 FVC predicted At day 60 (2.84 L) best effort by patient (2.81 L) % prediction (99 %). Refer table 1.2 for details.

# iii) Assessment of FEV1/FVC over a period of 90 days

The FEV1/FVC ratio, also called Tiffeneau - Pinelli index, is a calculated ratio used in the diagnosis of obstructive and restrictive lung disease. Acc. to GINA (Global Initiative For Chronic Obstructive Lung Disease) Post-bronchodilator FEV1/FVC ratio less than 0.70 should be used to indicate the presence of airway obstruction and this is applied to individuals of all ages, genders, heights and ethnicities.

Results of FEV1/FVC are as follows:

- Day 0 the FEV1/FVC ratio % Prediction is 77%.
- Day 30 the FEV1/FVC ratio % Prediction is 85%.
- Day 60 the FEV1/FVC ratio % Prediction is 90%.
- Day 90 the FEV1/FVC ratio % Prediction is 92%.

#### iv) Assessment of FEF 25-75 over a period of 90 days:

Forced expiratory flow over the middle one half of the FVC; the average flow from the point at which 25 percent of the FVC has been exhaled to the point at which 75 percent of the FVC has been exhaled.

Results of FEF 25-75 are as follows:

- Baseline Visit (Day 0): % Prediction 24%.
- Day 30: % Prediction 48% (Significant improvement over baseline)
- Day 60: % Prediction 45% (Significant improvement over baseline)
- Day 90: % Prediction 54% (Significant improvement over baseline)

# v) Assessment of ACQ (Asthma Control Questionnaire) over a period of 90 days:

The ACQ is able to identify the adequacy of asthma control in individual patient. In general, subjects with a score below 1.0 have adequately controlled asthma and above 1.0 their asthma was not well controlled. It was done on every visit.

- Baseline (Day 0) the total score was 1.85 (uncontrolled Asthma)
- Day 30 the total score 0.28 (adequately control of asthma).
- Day 60 the total score 0.42 (adequately control of asthma).
- Day 90 the total score 0.14 (adequately control of asthma).

# vi) Assessment of cough in daytime and nighttime:

Assessment of daytime and nighttime cough was done on the basis of subject reported diary. The diary captured episodes, their severity and duration. There was a gradual reduction in cough from baseline to the end of 90 days at which there was no daytime and nighttime cough.

# vii) Assessment of Laboratory related parameters:

Laboratory parameters like CBC, Liver function tests, Renal function tests, Lipid profile were observed to be within normal range both at baseline and at the end of the study. There was no significant change on these parameters.

### **Discussion & Conclusion:**

Bronchial Asthma requires continuous care through the use of bronchodilators and also cortisteroids. Though these medications are found to be effective they have potential side effects. The present study showed that Ayuasmo capsules was effective when used in a dose of 2 Capsules twice a day in a female patient suffering from Bronchial Asthma. Vital lung function tests like FEV1, FVC showed significant improvement over a period of 90 days of the study.

AyuAsmo capsules contains ingredients like *Vasa, Kulanjan, Puskaramula* which are anti-inflammatory and bronchodilator, *Yastimadhu* is a known mucolytic that helps to expel sputum, *Bharangi, Shati, Karkatashringi* have strong anti-histaminic action that help to prevent allergic reactions commonly observed in bronchial Asthma. *Bala* and *Amalaki* are proven anti-oxidants that help to prevent long term damage to the lung tissue.

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# TABLE 1 SPIROMETERY ASSESMENT

(TABLE N	O 1.1) ASSESSMENT	OF FEV1 MON	NTH WISE
VISIT SCHEDULE	PREDICTION	BEST	% PREDICTION
BASELINE VISIT	2.22	1.1	50%
DAY 30	2.22	1.8	81%
Day 60	2.22	1.59	70%
<b>Day 90</b>	2.22	2.04	90%
(Tab	ole No1.2) Assessment	of FVC Month	wise
Visit Schedule	Prediction	Best	% Prediction
Baseline Visit	2.84	1.83	64%
Day 30	2.84	2.72	96%
<b>Day 60</b>	2.84	2.27	80%
<b>Day 90</b>	2.84	2.81	99%
(Table I	No 1.3) Assessment of	FEV1/FVC Mor	nth wise
Visit Schedule	Prediction	Best	% Prediction
<b>Baseline Visit</b>	78.17	60.11	77%
Day 30	78.17	66.18	85%
Day 60	78.17	70.04	90%
<b>Day 90</b>	78.17	72.6	92%
(Table No	1.4) Assessment of FE	CF 25-75 (L/S) M	Ionth wise
Visit Schedule	Prediction	Best	% Prediction
Baseline Visit	2.92	0.69	24%
Day 30	2.92	1.4	48%
Day 60	2.92	1.3	45%
Day 90	2.92	1.57	54%

**Table 2 Dairy Card Assessment** 

(Table No 2.1) Assessment of ACQ (Asthma Control Questionnaire) Month wise		
Visit Schedule	Best	
Baseline Visit	1.85	
Day 30	0.28	

<b>Day 60</b>		0.42	
Day 90	0.14		
(Table no 2.2) Assess	sment of cough in day time	and Severity	
Visit	Cough (Day time)	Severity	
Day 15	11	Mild	
Day 30	20	Mild	
Day 45	11	Mild	
Day 60	0	No	
Day 75	0	No	
D. 00	0	No	
<b>Day 90</b>	Ŭ		
·	ment of Cough At night tin	ne and severity	
•	ment of Cough At night tin	ne and severity	
(Table no. 2.3) Assess:	ment of Cough At night tin	•	
(Table no. 2.3) Assess Visit	ment of Cough At night tin	Cough	
(Table no. 2.3) Assess.  Visit  Day 15	ment of Cough At night tin	Cough 0	
(Table no. 2.3) Assess  Visit  Day 15  Day 30	ment of Cough At night tin	Cough  0 0	
(Table no. 2.3) Assess.  Visit  Day 15  Day 30  Day 45	ment of Cough At night tin	Cough  0 0 0	

**Table 3 Laboratory parameter** 

(Table no 3.1) Assessment of Hematology					
Hematology	BT	AT			
RBC	4.41	4.3			
WBC	7100	9000			
НВ	13.4	12.4			
Platelets	1.89	2.5			
ESR	4	25			
(Table no 3.	(Table no 3.2) Assessment of Renal Function test				
RFT	BT	AT			
Sr. Creatinine	1	0.7			
BUN	13.95	15.16			
(Table no. 3.3) Assessment of Liver function test					
LFT	BT	AT			

T Bilurubin	0.8	0.4		
AST	24	26.3		
ALT	24	26.3		
ALI	24	20.3		
ALP	198	166.3		
Total Protien	6.2	7.9		
A/G Ratio	1.5:1	1		
(Table no.	3.4) Assessment of L	ipid Profile		
Lipid Profile	BT	AT		
Total Cholestrol	190	184.3		
Triglycride	148	95.1		
LDL	113.4	116		
HDl	47	49.3		
VLDL	29.6	19		
(Table no. 3.5) Assessment of Blood sugar (Fasting)				
Blood Sugar	BT	AT		
Fasting	95	82		
Table no 3.6) Assessment of urine examination				
Urine Examination	BT	AT		
Protien	Trace	Absent		
Glucose	Nil	Absent		
Ketone	-	Absent		

#### Report FEV1 %Pred COPD SEVERITY FVC%Pred Interpretation 150 F(Litres/Sec) 16 OBS NORM OBS NORM 125 125 14 100 100 75 75 12 MODERATE 50 SEVERE 10 VERY SEVERE MIXED RES 8 PEFR 25 50 75 100 125 150 25 50 75 100 125 150 ofef25% 6 (FEV1/FVC)%Pred (FEV1/FVC)%Pred □FEF50% Spirometry(FVC Results) 4 Parameter Pred M.Pre %Pred M.Post %Pred %Imp □FEF75% 2 FVC 02.84 01.83 064 02.81 099 (L) +54 V(Litres) FEV1 (L) 02.22 01.10 050 02.04 092 +85 0 (왕) FEV1/FVC 78.17 60.11 077 72.60 093 +21 FEF25-75 (L/s)02.92 00.69 024 01.57 054 +128 -2PEFR (L/s)06.82 01.77 026 03.74 055 +111 -4 FIVC (L) 02.01 02.65 +32 FEV.5 (L) 00.71 01.41 +99 -6 02.76 02.81 102 FEV3 01.75 063 +61 (L) 01.40 02.18 +56 PIFR (L/s)-8 FEF75-85 00.68 +119 (L/s)00.31 \_\_\_ FEF.2-1.2(L/s) 05.28 00.92 017 02.96 056 +222 -10 FEF 25% (L/s)06.02 01.31 022 03.14 052 +140 V(Litres) 8 FEF 50% (L/s)04.67 00.80 017 01.73 037 +116 FEF 75% 00.84 035 02.42 00.38 +121 (L/s)016 7 POST FEV.5/FVC (왕) 38.80 50.18 +29 FEV3/FVC (왕) 97.18 95.63 098 100.00 103 +05 FET (Sec) 03.54 02.67 6 ExplTime (Sec) 00.09 00.08 Lung Age (Yrs) 038 057 150 041 108 -28 5 FEV6 02.84 (L) 01.29 02.02 FIF25% (L/s)------+57 4 FIF50% (L/s)01.29 02.09 +62 01.07 FIF75% 01.94 +81 (L/s)FV 3 oPEV1 2 Pre Test COPD Severity Restrictive stage COPD as FEV1/FVC >= 70% and FEV1 < 80% 1 Post Test COPD Severity Test within normal limits 0 T(Seconds) Medication Report Indicates Early Small Airway Obstruction as FEF 25-75 %Pred or PEFR %Pred < 70 Mixed Blockage as (FEV1/FVC) Pred < 95 and FVC Pred < 80Post Medication Report Indicates Early Small Airway Obstruction as FEF 25-75 %Pred or PEFR %Pred < 70 Mild Obstruction as (FEV1/FVC) %pred <95 and FVC%Pred >80 yogesh

The contents of this report require clinical co-relation before any clinical action.

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