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# Evaluation of efficacy of *Dhoopana* (polyherbal nasal fumigation) versus polyherbal steam inhalation in children with *Pratishyaya* (rhinitis): A pilot study

**Research Article** 

# Monika1\*, Renu B Rathi<sup>2</sup>, Bharat Rathi<sup>3</sup>, Deepthi Balakrishnana<sup>4</sup>

1. Ph.D. Scholar, 2. Professor & H.O.D, Department of Kaumarabhritya; 3.Professor, Department of RSBK, Mahatma Gandhi Ayurveda Medical College, Hospital & Research Centre, Salod (H), Wardha (Maharashtra). India. 4. Professor & H.O.D, Dept. of Kaumarabhritya, PNNM Ayurveda Medical College, Cheruthuruthy, Kerala, India.

## Abstract

Background: According to Ayurveda, Pratishyaya is characterized by the continuous outward flow of Vata, Pitta, and Kapha from the nostrils, resembling rhinitis, where the nasal mucosa becomes inflamed due to infection, allergy, or injury. The study aims to evaluate the efficacy of polyherbal steam inhalation versus nasal fumigation (Dhoopana) in children with rhinitis. Material and methods: Randomized reference controlled open label equivalence pilot study. Patients after initial screening were subjected to randomization and were included in two groups; Group C (Control group) and Group T (Trial group). 10 subjects in each group. Group C (Control group) Polyherbal steam inhalation, Arka prepared from Tulsi, Nirgundi, Vasa and Nilgiri were used for steam inhalation. Group T (Trial group) Dhoopana (Polyherbal nasal fumigation). Dhoomvarti prepared from dry leaves of Tulsi, Nirgundi, Vasa and Nilgiri. Results: From the observations of the clinical trial, it can be concluded that efficacy of Dhoopana (polyherbal fumigation) is better as compare to polyherbal steam inhalation. Statistically significant difference is seen in objective criteria of nasal patency and symptoms of nasal itching and coughing whereas no significant difference is observed in symptoms rhinorrhoea, nasal obstruction, headache and anorexia. Conclusion: Rhinitis in acute stage is one of the frequently troubling conditions in children, though many medicines are available in the market in every system of medicine for its management, but it is difficult to administer them orally in pediatric population. Polyherbal Dhoopana (Nasal Fumigation) is safe and easy to administer, so, can be used in general pediatric practice in patients suffering from acute condition of rhinitis.

**Keywords:** *Ayurvedic treatment, Dhoopana,* Inflammatory response, Pediatric rhinitis, Polyherbal steam inhalation, Therapeutic efficacy.

# Introduction

As per Ayurveda, Pratishyaya (rhinitis) is one among Nasagata Roga (nasal disorders) in which Kaphadi Doshas(principal constituents of the body) are continuously eliminated through the nose. These vitiated Doshas get accumulated in the head and their further movement towards the nose causes Pratishyaya. According to Acharya Charaka, Definition of Pratishyaya is "Pratikshnamshyayatiitipratishyaya" means continuous outward movement of Vata. Pitta and Kapha doshas from nostrils is Pratishyaya.(1) Pratishyaya is classified into 5 types as Vataja, Pittaja, Kaphaja, Sannipataja and RaktajaPratishyaya.(2) Similar clinical presentation of *Pratishvava* can be seen in Rhinitis. In rhinitis, nasal mucosa gets inflamed due to any infection, allergy or injury.(3) The prevalence of non-allergic rhinitis is about 40%.(4) The prevalence of allergic rhinitis in India was reported "11.3% in children

\* Corresponding Author:

#### Monika

PhD Scholar Department of Kaumarbhritya. MGACH&RC, Salod, Wardha, Maharashtra, India, Pin code-442001 Email Id: <u>ayurmonika@gmail.com</u> aged 6-7 years and 24.4 % in children aged 13-14 years. (5) Acharya Videha explained clinical presentation of Pratisyaya as Excessive secretions from nasal cavity and eyes, fever, generalized weakness with severe headache. (6) Nasal mucosae have rich blood supply, stimulation of sympathetic nervous system causes vasoconstriction which further result in shrinkage of nasal mucosa on the other side, stimulation of parasympathetic system is responsible for excessive secretion from the nasal mucosa along with dilatation of local vessels. Emotional disturbance also plays a significant role as autonomic nervous system supply of nasal mucosa is under the control of hypothalamus. If no any treatment is given at early stage it may get complicated and further lead to other comorbid conditions like chronic rhinitis, cough, or breathing difficulty with debility.(7) There is no use of Antibiotics in acute rhinitis.(8) Antihistamine-decongestants are frequently used in cough and cold. But in some studies, these were not found effective in the management of rhinitis.(9) There is limited data regarding safety of Pseudoephedrine 4 and phenylephrine in rhinitis.(10) Various Ayurveda interventions are mentioned in Avurveda classics as protocol for the management of Pratishyaya, out of which Dhoopana is one of the interventions, could effectively manage the disease.



Dhoopana is a natural way to sterilize the environment; Fumes generated from various herbs works through their antimicrobial properties. Dhoopana is also having therapeutic benefits and used in various disorders like Fever, cough, cold, headache, wound healing and epilepsy. Many Dhoopana drugs are excellent environmental cleansers but present study focuses on Pratishyaya (Rhinitis), Inflammation of the nasal cavity. The aim of the study is to test the efficacy of Dhoopana (Polyherbal nasal fumigation) versus Polyherbal steam inhalation in children with Pratishyaya (Rhinitis).

## Materials and methods

Study type and study design: Randomized reference controlled open label equivalence pilot study

#### **Population:**

- Inclusion criteria: Patients of either sex aged between 7 to 14 years. Patients suffering from common cold, presenting with features of *Pratishyaya* (Rhinitis) for 7-10days. Patients willing and able to participate in the study.
- Exclusion criteria: Patients suffering from common cold for more than 10days. Patients with Chronic Allergic Rhinitis, *Dushta Pratishyaya, Raktaja Pratishyaya, Sannipataja Pratishyaya* and Infectious diseases like T.B. Patients suffering from Cleft Palate, Deviated nasal septum and Nasal Polyps. History of hypersensitivity to the trial drug or any of its ingredients.

**Place of study/source of data:** O.P. and I.P. Department of Kaumarabhritya, Mahatma Gandhi Ayurveda College and Hospital Wardha

Sample size: 10 children in each group.

**Sample selection technique:** Computer generated random number table.

**Drug:** The trial drugs i.e., *Dhoopanavarti* and *Arka* were prepared in *Rasa Shastra* lab of Mahatma Gandhi Ayurveda college Hospital and Research Centre.

**Grouping and Posology**: Patients after initial screening were subjected to randomization and were included in two groups; Group C (Control group) and Group T (Trial group).

- Group C (Control group) Polyherbal steam inhalation: Arka prepared from Tulsi, Nirgundi, Vasa and Nilgiri. In mild symptoms, 2.5 ml Arka was added in 500ml of water for 1minute steam inhalation and in case of moderate to severe symptoms, 5ml Arka was added in 500ml of water for steam inhalation for 1.5minute.
- Group T (Trial group) *Dhoopana* (Polyherbal nasal fumigation): *Dhoomvarti* (fumigation stick) was prepared from the powder of dry leaves of *Tulsi*, *Nirgundi*, *Vasa and Nilgiri*. At the time of administration, *Dhoomvarti* was dipped in cow ghee then it was ignited using matchbox, after few seconds

it was extinguished and then its fumes were instilled through nasal route. Nasal fumigation was given for 1 minute in mild symptoms and for 1.5 minutes in moderate to severe *Pratishyaya*.

**Treatment period:** 7 days, twice daily, once in morning and evening. Assessment on 1<sup>st</sup>, 3<sup>rd</sup>, 5<sup>th</sup> and 7<sup>th</sup> day.

Follow up period: 14 days after the treatment period.

**Outcome:** Outcome of the study was assessed using subjective and objective criteria.

**Subjective criteria:** Sign and symptoms of *Pratishyaya* (Rhinitis) by using TNSS (Total Nasal Symptom Score).

Symptom	0 (None)	1 (Mild)	2 (Moderate )	3 (Severe)
Nasal Congestion	No congestion	Occasional, minimal	Frequent, noticeable	Constant, severe
Rhinorrhea	No	Occasional,	Frequent,	Constant,
(Runny Nose)	discharge	minimal	moderate	profuse
Sneezing	No	Occasional,	Frequent,	Severe,
	sneezing	mild	moderate	persistent

Table 1: TNSS (Total Nasal Symptom Score)

**Total TNSS Score** = Sum of individual symptom scores (Range: **0-12**).

mild

Occasional, Frequent,

Severe,

moderate bothersome

Higher scores indicate more severe rhinitis symptoms.

Nasal Itching No itching

**Objective criteria**: Assessment of nasal blockage by using modified cold spatula test.

**Statistical analysis:** The data underwent analysis employing suitable descriptive and inferential statistical methods. Quantitative variables were analyzed by the "Student t-test" whereas the Chi-square test was employed for subjective parameters.

**Ethical Consideration-** The study was approved by Ethics Committee of MGACHRC

#### Consort flow chart is as follows: Figure No. 1: CONSORT Flow chart



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#### **Observations and Results** Weight characteristics

In the control group (N=10), the mean weight is 24.30 with a standard deviation of 7.13 and a standard error mean of 2.25. In the experimental group (N=10), the mean weight is 21.00 with a standard deviation of 6.21 and a standard error mean of 1.96. A t-test was conducted to compare the means of weight between the control and experimental groups, resulting in a t-value of 1.103 and a p-value of 0.285. The p-value suggests that there is no statistically significant difference in weight between the two groups.

#### **Height characteristics**

In the control group (N=10), the mean height is 138.40 with a standard deviation of 14.03 and a standard error mean of 4.43. In the experimental group (N=10), the mean height is 137.40 with a standard deviation of 12.86 and a standard error mean of 4.06. A t-test was conducted to compare the means of height between the control and experimental groups, resulting in a t-value of 0.16 and a p-value of 0.87. The p-value

suggests that there is no statistically significant difference in height between the two groups.

#### Prakriti (body constitution)

- *Pitta-Kapha*: There is 1 individual in the control group and 1 individual in the experimental group. This represents 10.0% of the experimental group and 5.0% of the control group.
- *Vata-Kapha*: There are 7 individuals in both the control and experimental groups, accounting for 70.0% of each group.
- *Vata-Pitta*: There are 3 individuals in the control group and 2 individuals in the experimental group, making up 30.0% of the control group and 20.0% of the experimental group. Overall, they represent 25.0% of the total sample.

#### Interpretation

Both before and after treatment, there is no significant association between rhinorrhoea grades and group membership. In other words, the distribution of rhinorrhoea grades does not differ significantly between the control and experimental groups before and after treatment. Detail in Table No.1 and Figure No.2.

#### Table 2: Comparative evaluation of Rhinorrhoea severity Control against Experimental on basis of TNSS grading scale

		Befor	e treatment	After treatment			
Parameters		<b>Control- frequency</b>	<b>Experimental- frequency</b>	<b>Control- frequency</b>	<b>Experimental- frequency</b>		
	Grade Zero	10.0%	0.0%	70.0%	90.0%		
Rhinorrhoea	Grade One	50.0%	70.0%	20.0%	10.0%		
	Grade Two	30.0%	30.0%	10.0%	0.0%		
	Grade Three	10.0%	0.0%	-	-		
	Chi square value		2.333	1.583			
	P value		0.506	0.453			
		Befor	e treatment	After treatment			
		Control- frequency Experimental- frequency		Control- frequency Experimental- frequenc			
	Grade Zero	20%	20%				
	Grade One	70%	80%				
Nasal Itching	Grade Two	10%	10%				
	Chi square value		1.067	NA			
	P value		0.58	NA			
		Befor	e treatment	After treatment			
		Control- frequency Experimental- frequency		Control- frequency	Experimental- frequency		
	Grade Zero	10.0%	0.0%	70%	80%		
	Grade One	70.0%	70.0% 60.0% 3		20%		
Nasal obstruction	Grade Two	20.0%	40.0%	-	-		
	Chi square value	1.744		0.267			
	P value		0.4182	0.606			
		Before treatment		After treatment			
		<b>Control- frequency</b>	<b>Experimental- frequency</b>	<b>Control- frequency</b>	<b>Experimental- frequency</b>		
Sneezing	Grade Zero	10.0%	10.0%	30.0%	60.0%		
	Grade One	50.0%	80.0%	70.0%	40.0%		
	Grade Two	40.0%	10.0%	-	-		
	Chi square value		2.492	1.818			
	P value		0.288	0.178			
		Befor	e treatment	After treatment			
		Control- frequency Experimental- frequency		<b>Control- frequency</b>	y Experimental- frequency		
Cough	Grade Zero	40.0%	10.0%	70.0%	100.0%		
	Grade One	40.0%	70.0%	30.0%	0.0%		
	Grade Two	20.0% -		-	-		
	Chi square value		2.618	3.529			
	P value		0.27	0.06			



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				Before treatment					After treatment			
				Contr	ol- frequency	Experime	ntal- frequen	cy	Control- frequenc	y Experi	mental- frequency	
Headache severity		Grad	e Zero	30.0%		60.0%			90.0%		90.0%	
		Grad	le One		60.0%	30.0%			10.0%		10.0%	
		Grad	e Two		10.0%	10.0%			-		-	
		Chi squ	are value	2					0			
		P v	alue	0.368				1				
			Before treatment						After treatment			
				Contr	ol- frequency	Experimental- frequency		cy	<b>Control- frequence</b>	ontrol- frequency Experimental- freq		
Anorexia severity	Grad	e Zero		60.0%	60.0%			80.0%		100.0%		
	Grad	le One		40.0%	40.0%			20.0%		0.0%		
	enty	Chi squ	are value			0			2.22			
		P v	alue	1				0.13				
					Nost	ril size (mn	1 square)					
					Mean	Ν	Std Devia	l. tion	Std. Error Mean	T-test	P-value	
Control Rt no sc Lt Tot	Rt nost	tril (mm Before		•	436.6000	10	97.53	427	30.84304	4 10500	0.002224	
	square)		After		602.0000	10	118.85	5752	37.58605	-4.19509	0.002324	
	T 4 m	t nostril Before After		•	465.8000	10	145.37	7140	45.97047	5 04274	0.000/0/	
	LUN				760.0000	10	156.42	2677	49.46649	-5.04574	0.000696	
	Total	al (mm Before uare) After			857.0000	10	217.72	2409	68.85040		<0.01	
	squ				1362.0000	10	227.06	5044	71.80282	-5.5509	<0.01	
F Experimental	Rt nost	nostril (mm Before square) After		e	472.0000	10	118.38	3919	37.43795	2 0776	0.004	
	squ				631.2000	10	149.44	4995	47.26023	-3.8770	0.004	
	Lt nostril		Before	•	442.0000	10	137.96	5296	43.62772	4 01241	0.002	
			After		657.6000	10	96.64	965	30.56330	-4.01241	0.003	
	Total	(mm	Before	e	919.6000	10	234.76	5097	74.23794	5.042	0.000608	
	square)		After		1288.8000	10	185.21	1147	58.56901	-3.042	0.000098	

#### Follow up

Post treatment follow was taken for the next 14 days in two visits and no any recurrence of rhinitis was noted in all the 20 patients. No any side effect was noted in both the groups.

# Discussion

The disease *Pratishyaya* is very common in Pediatric population, many medicines are available for its management but no any standardized Ayurvedic therapy is available which can be given locally in nostrils in its acute stage. So, this study was planned to work upon acute rhinitis. Local fumigation and steam inhalation are simple and fast effective as compare to oral medication.

Kashyapa Samhita is mainly devoted to *Kaumarbhritya* branch (Ayurvedic Pediatrics) of *Ashtanga Ayurveda* and *Dhoopana* is described in the treatment protocol of *Pratishyaya*.(11) In this mode, various herbs are made into a stick known as *Varti* and its fumes are used for local fumigation of the affected area. It can be considered as an ayurvedic therapy that delivers the medicines directly into the airways which provide relief and protection to the local regions. (12)

#### Discussion on mode of Action of drugs

In Pratishyaya, treatment should be aimed to relieve the Avarodha (obstruction) created by the Dosha. Drugs having Ushna (hot) and Tikshna (pungent) Properties are indicated in Dhoopana. Tulsi(Ocimum tenuiflorum L.), Nirgundi (Vitex negundo L.), Vasa (Justicia adhatoda L.) and Nilgiri (Eucalyptus obliqua L'Hér.) are popular Dhoop dravyas

(material for fumigation) for Respiratory problems. (13) Dhoopana karma has Rukshana properties because of which it is helpful in nasal discharge and cough. Fumes of these herbs may help in relieving the sign and symptoms of Pratishyaya due to their Ushna Virya (hot potency) and Kapha Vata Shamak (Dosha relieving) properties. (14) Vasa (Justicia adhatoda L.) is known effective in Respiratory tract related disorders. It may help in combating Feverish sensation present in Pratishyaya due to its Kapha Pitta Shamak and Jwaraghana (antifebrile) Properties. (15) Moreover, in one recent In-silico study, Phytochemicals of Nirgundi, Vasa and Nilgiri were found effective against protein targets of Covid-19. (16) Due to its Ushna (hot) and Tikshna (pungent) properties and ability to reach minute Srotasa (channels of the body) it is beneficial in nasal obstruction, sneezing, headache and anorexia. Hence, these four herbs have proven efficacy for steam inhalation. (17)

There is a vast description of *Tulsi (Ocimum tenuiflorum* L.) in all the ayurvedic classics in respiratory disorders. *Tulsi* along with curing viral, bacterial, fungal infections gives relief from congestion and helps in smoother breathing. It has antiallergic, antibacterial, antifungal, disinfectant properties. A study shows that *Tulsi* given in the form of aerosol through nebulization is highly efficient. (18). *Neem (Azadirachta indica Juss.)* is also frequently employed as one of the primary constituents in most formulations of *Dhoopana.*(19) Azadirachtin, found in Neem seed oil, has been shown to have insecticidal effects. In a study, Staphylococcus aureus showed a high level of inhibition to the fumes of *Azadirachta Indica* L., with a



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maximum of 90% inhibition in 10 minutes and 50% in 5 minutes.(20)

#### **Discussion on results**

Rhinorrhoea of grade zero was present in 5% patients before treatment which then improved to 80% patients; this shows significant improvement in rhinorrhoea. Nasal itching of grade zero was present in 20% patients before treatment which then improved to 100% patients after treatment. Nasal obstruction of grade zero was present in 5% patients before treatment which then improved to 75% patients after treatment. Sneezing of grade zero was present in 10% patients before treatment which then improved to 45% patients after treatment. Cough of grade zero was present in 25% patients before treatment which then improved to 85% patients after treatment. Headache of grade zero was present in 55% patients before treatment which then improved to 90% patients after treatment. Anorexia of grade zero was present in 60% patients before treatment which then improved to 90%.



# Conclusion

Polyherbal *Dhoopana* (Nasal Fumigation) is safe and more effective over Polyherbal steam inhalation, so, can be used in general pediatric practice in patients suffering from acute condition of rhinitis. It is concluded that the intervention had a notable impact on removing nasal blockage and increasing nasal patency during rhinitis, as indicated by the significant differences in measurements before and after the intervention in both groups. Further large-scale studies can be more resourceful in drawing fruitful results.

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