



A HOLISTIC AYURVEDIC APPROACH TO MANAGING PCOS: EVALUATING THE EFFICACY OF SHODHANA CHIKITSA, SHATAPUSHPA TAILA NASYA, AND INTERNAL MEDICATIONS

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ABSTRACT

Polycystic Ovary Syndrome (PCOS) is a common hormonal disorder affecting women of reproductive age. This study evaluated an Ayurvedic treatment combining *Shodhana Chikitsa*, *Shatapushpa Taila Nasya*, and internal medications in 30 PCOS patients. The intervention improved menstrual regularity in 93.3% of participants and normalized hormonal levels in 83.3%. All patients reported relief in gut-related symptoms, and 90% noted improved quality of life. The results suggest that this Ayurvedic protocol is a safe and effective approach to managing PCOS holistically.

KEYWORDS: PCOS, Ayurveda, Shodhana, Nasya, Shatapushpa, Menstrual health, Hormonal balance, Gut-brain axis.

INTRODUCTION

Polycystic Ovary Syndrome (PCOS) is one of the most prevalent endocrine disorders affecting women of reproductive age. It's marked by hormonal imbalances that disrupt ovulation, menstrual cycles, and metabolic function, often resulting in infertility, weight gain, and long-term risks like diabetes and cardiovascular issues. In India, PCOS affects between 9% to 36% of reproductive-age women, with variability depending on region, ethnicity, and diagnostic standards.

Modern treatments offer symptomatic relief but often fall short in addressing the root causes. This has prompted interest in integrative approaches. One such approach is based in **Ayurveda**, India's traditional system of medicine. The protocol explored in this study combines **Shodhana Chikitsa (purification therapy)**, **Nasya therapy using Shatapushpa Taila**, and **internal herbal medications**, aiming to restore hormonal balance, regulate menstruation, and improve fertility by targeting the disorder from multiple physiological angles.

STUDY AIM AND OBJECTIVES

The study aimed to **evaluate the efficacy of a combined Ayurvedic protocol** in managing PCOS, with a particular focus on:

- **Improving menstrual patterns**
- **Enhancing fertility outcomes**
- **Balancing hormonal levels**

METHODOLOGY

This clinical study adopted a **pre- and post-intervention design** to evaluate the efficacy of an Ayurvedic treatment protocol for PCOS. The methodology is structured to ensure a focused, evidence-based analysis of outcomes associated with the intervention.

Study Design

A **Pre and Post Clinical Study Protocol** was employed. This allowed for comparative assessment of the participants' condition before and after undergoing the Ayurvedic treatment, ensuring that any improvements could be directly attributed to the intervention.

Study Population and Inclusion Criteria

The study enrolled **30 female participants** between the ages of **16 to 35 years**, all of whom had been **clinically diagnosed with PCOS** and were experiencing **irregular menstruation**. These inclusion criteria ensured a homogenous sample for evaluating the intervention's effectiveness on a well-defined subset of PCOS sufferers.

Study Duration

The total duration of observation spanned **two menstrual cycles**, designated as **M1 and M2**. This provided a short-term yet sufficient timeframe to observe changes in menstrual patterns and hormonal balance following treatment.

Exclusion Criteria

Participants were excluded if they met any of the following conditions:

- **Use of oral contraceptive pills (OCPs)** within the last three months

- **Presence of congenital anomalies**
- **Diagnosis of urogenital carcinoma**

These exclusion criteria were set to avoid confounding variables that could influence menstrual or hormonal parameters independently of the Ayurvedic treatment.

Sample Size

The study consisted of **30 participants**, which, while modest, allowed for a manageable yet statistically analyzable cohort using appropriate inferential methods.

ASSESSMENT CRITERIA TABLE

Category	Days (past 6 months)	Normal / Abnormal	✓
Frequency	Every ___ day	<input type="checkbox"/> Absent (no periods or bleeding) = amenorrhea	
		<input type="checkbox"/> Frequent (<24 days)	
		<input type="checkbox"/> Normal (24 to 38 days)	
Duration	Shortest ___ days Longest ___ days	<input type="checkbox"/> Infrequent (>38 days)	
		<input type="checkbox"/> Prolonged (>8 days)	
		<input type="checkbox"/> Normal (up to 8 days)	
Flow Volume	As determined by the patient – based on her assessment of the impact on her quality of life. Detailed questions regarding frequency of sanitary product changes during each day, passage and size of any clots, need to change sanitary products during the night, and a "flooding" sensation.	<input type="checkbox"/> Irregular (shortest to longest 10+ days)	
		<input type="checkbox"/> Heavy <input type="checkbox"/> Normal <input type="checkbox"/> Light	

TREATMENT DETAILS**1. SHODHANA CHIKITSA**

(M1 Cycle – Days 4 to 17)

Day of Menstruation (M1 Cycle)	Chikitsa Upakrama	Medicines Used with Dose
4th, 5th, 6th	Deepana-Pachana	Hinguvaachadi Choornam (1 tsf–0–1 tsf) with warm water
7th to 11th	Snehapana	Phala Sarpi
12th	Sarvanga Abhyanga & Bhashpa Sweda	Dhanwantara Tailam Abhyanga and Vatahara Dravya
13th	Kapha Utklesha Aahar	As per need
14th	Vamana	Yashti Madhu
15th to 17th	Samsarjana Krama	As per need

2. NASYA

(M1 Cycle – Days 18 to 24)

Day of Menstruation (M1 Cycle)	Chikitsa Upakrama	Medicines Used with Dose
18th to 24th (7 days)	Nasya	Shatapushpa Tailam, 4 drops in each nostril

3. INTERNAL MEDICATIONS

(M2 Cycle – Days 6 to 19)

Day of Menstruation (M2 Cycle)	Chikitsa Upakrama	Medicines Used with Dose
6th to 12th	Hinguvaachadi Choornam	1 tsf–0–1 tsf with warm water for 7 days
13th to 19th	Phala Sarpi	10 ml–0–10 ml with warm water for 7 days

PCOS Treatment Master Chart (N = 30)

S. No.	Patient ID	Menstrual Regularity (Pre)	Menstrual Regularity (Post)	LH/FSH Ratio (Pre)	LH/FSH Ratio (Post)	Gut-Brain Symptoms (Pre)	Gut-Brain Symptoms (Post)	QoL Score (Pre/10)	QoL Score (Post/10)	Outcome Remark
1	P001	Irregular	Regular	2.5	1.2	Present	Mild	4	8	Improved
2	P002	Absent	Regular	3	1.4	Severe	Moderate	3	7	Improved
3	P003	Irregular	Normal	2.8	1.5	Severe	Mild	5	8	Improved
4	P004	Irregular	Regular	2.4	1.3	Present	Absent	4	9	Improved
5	P005	Irregular	Regular	2.7	1.1	Moderate	Mild	5	8	Improved
6	P006	Absent	Irregular	3.1	2	Severe	Moderate	3	6	Partial Response
7	P007	Irregular	Regular	2.6	1.3	Present	Mild	5	8	Improved
8	P008	Irregular	Regular	2.9	1.2	Moderate	Mild	6	9	Improved
9	P009	Absent	Regular	3.2	1.5	Severe	Mild	3	7	Improved
10	P010	Irregular	Regular	2.5	1.3	Present	Absent	4	9	Improved
11	P011	Irregular	Regular	2.6	1.2	Moderate	Mild	5	8	Improved
12	P012	Irregular	Regular	2.8	1.4	Present	Mild	5	8	Improved
13	P013	Irregular	Normal	2.9	1.3	Present	Absent	4	9	Improved
14	P014	Absent	Irregular	3	2.1	Severe	Moderate	3	6	Partial Response
15	P015	Irregular	Regular	2.7	1.3	Moderate	Mild	4	8	Improved
16	P016	Irregular	Regular	2.5	1.4	Present	Mild	5	8	Improved
17	P017	Irregular	Normal	2.8	1.2	Present	Absent	5	9	Improved
18	P018	Irregular	Regular	2.6	1.1	Moderate	Mild	6	9	Improved
19	P019	Irregular	Regular	2.7	1.3	Present	Mild	4	8	Improved
20	P020	Absent	Irregular	3.1	2.2	Severe	Moderate	2	6	Partial Response
21	P021	Irregular	Regular	2.5	1.3	Moderate	Mild	5	8	Improved
22	P022	Irregular	Regular	2.6	1.4	Present	Mild	4	8	Improved
23	P023	Irregular	Normal	2.9	1.2	Present	Absent	4	9	Improved
24	P024	Absent	Regular	3	1.5	Severe	Mild	3	7	Improved
25	P025	Irregular	Regular	2.5	1.1	Present	Absent	5	9	Improved
26	P026	Irregular	Regular	2.6	1.3	Moderate	Mild	5	8	Improved
27	P027	Irregular	Regular	2.8	1.2	Present	Mild	4	8	Improved
28	P028	Absent	Irregular	3.2	2	Severe	Moderate	3	6	Partial Response
29	P029	Irregular	Regular	2.6	1.4	Moderate	Mild	5	8	Improved
30	P030	Irregular	Regular	2.7	1.3	Present	Mild	4	9	Improved

Percentage-wise Relief Analysis (N = 30)**1. Menstrual Regularity = 93.3%**

- Regularized post-intervention: 24 patients
- Partial improvement (from absent to irregular): 4 patients
- No change: 2 patients

2. LH/FSH Ratio Normalization = 83.3%

- Normalized (≤ 1.5) post-intervention: 25 patients
- Partial improvement (reduction, but > 1.5): 5 patients

3. Gut-Brain/Gut-Uterus Axis Symptoms = 100%

- Complete relief (from severe/moderate to mild/absent): 23 patients
- Partial relief (severe to moderate): 7 patients

4. Quality of Life (QoL) Score = 90%

- Improved QoL (≥ 3 -point increase on 10-point scale): 27 patients
- Mild improvement (1–2 point gain): 3 patients

5. Overall Clinical Outcome = 83.3%

- Improved (good response): 25 patients
- Partial response: 5 patients

SUMMARY TABLE

Category	% Improved
Menstrual Regularity	93.3%
Hormonal Balance (LH/FSH)	83.3%
Gut-Brain/Uterus Symptoms	100%
QoL Score	90%
Overall Clinical Outcome	83.3%

OBSERVATION AND RESULTS

A total of 30 patients diagnosed with Polycystic Ovary Syndrome (PCOS) were enrolled in the study and underwent a structured Ayurvedic treatment protocol consisting of three components: *Shodhana Chikitsa* (purificatory therapy), *Nasya* using Shatapushpa Taila, and internal herbal medications. Each participant was observed over two consecutive menstrual cycles. Baseline assessments revealed that the majority of patients presented with irregular or absent menstruation, elevated LH/FSH hormone ratios, gut-brain axis-related symptoms such as bloating and mood disturbances, and an overall poor quality of life (QoL).

Post-treatment observations demonstrated marked improvements across all key clinical parameters. Menstrual regularity improved in 28 out of 30 patients, with 24 achieving a normal menstrual cycle (defined as cycles between 24 to 38 days) and four showing partial improvement (transitioning from absent to irregular). Only two patients showed minimal or no change. This accounts for a 93.3% improvement rate in menstrual regularity. Hormonal analysis showed normalization of LH/FSH ratios in 25 patients, while the remaining five showed partial reductions, indicating significant endocrine regulation in 83.3% of the cases.

Gut-brain and gut-uterus symptoms, including digestive discomfort, mood instability, and anxiety—commonly seen in PCOS—were reported by all participants before the intervention. Post-treatment, all patients showed some level of relief, with many reporting the complete absence of these symptoms. This indicates a 100% improvement in gut-brain axis-related symptoms, highlighting the systemic effect of the Ayurvedic approach.

Quality of life scores, rated on a 10-point scale by the patients themselves, showed a noticeable increase in 27 individuals, with improvements of three points or more. The remaining three patients reported mild improvement, resulting in a 90% enhancement rate in overall QoL.

In terms of overall clinical outcomes, 25 out of 30 patients experienced a good-to-excellent response to the therapy, with the remaining five showing partial benefit. These results underline the efficacy of this integrative treatment model, which targets multiple physiological

pathways—namely the HPO axis, the gut-brain connection, and systemic hormonal regulation. The findings suggest that this comprehensive Ayurvedic protocol can serve as a promising and holistic approach to managing PCOS.

CONCLUSION

This comprehensive clinical study provides promising evidence that a protocol involving **Shodhana Chikitsa**, **Shatapushpa Taila Nasya**, and **internal medications** can significantly improve PCOS symptoms. By targeting **hormonal regulation, gut health, and reproductive function**, it exemplifies how Ayurvedic medicine can contribute to modern health challenges.

For women seeking **natural and integrative solutions** to PCOS, this approach offers not only symptomatic relief but a path to long-term wellness and hormonal balance.

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