A CLINICAL STUDY TO EVALUATE THE EFFICACY OF PICCHABASTI AND KAPIT-TASHTAKACHURNA IN THE MANAGEMENT OF IRRITABLE BOWEL SYNDROME W.S.R. TO PRAVAHIKA

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ABSTRACT

Irritable bowel syndrome is a functional bowel disorder characterized by abdominal pain or discomfort and altered bowel habits in the absence of detectable structural abnormalities. Throughout the world about 10-20% of adult and adolescents have symptoms consistent with IBS and most studies show a female predominance. Women are diagnosed with IBS 2 to 3 times as often as men and make up to 80% of the population with severe IBS. Considering this, the study was undertaken to evaluate the efficacy of picchabasti (one of the treatment modality), followed by Kapittashtakachurna as shamanoushadhi in the Management of Irritable bowel syndrome vis-à-vis Pravahika. Materials and Methods; It was an observational clinical study with pre and post-test design, consisting of Picchabasti (niruha), pippalyaditaila (anuvasana) and Kapittashtakachurna as shamanoushadhi. The subjects fulfilling the diagnostic criteria of IBS vis-à-vis Pravahika were assigned in to single group consisted of 20 subjects. The symptoms like, Tenesmus (kunthanasheela mala pravritti), constipation (Baddhamalapraavritti), abdominal pain/discomfort (Udarashoola), mucus in stool, Gas/Flatulence, and increased frequency of defeation on grading were taken as parameters of assessment. The study consisted of 3 assessments i.e on 0th day, 8th day and on 31st day. Results: Statistical results on parameters showed highly significant results on symptoms of abdominal pain/discomfort, tenesmus, mucus in stool, flatulence and increased frequency of defeation with p value 0.000 and statistically significant result on symptom of constipation with p value 0.015.

Keywords: IBS, Pravahika, kapittashtakachurna, picchabasti, pippalyaditaila.

INTRODUCTION
Irritable Bowel Syndrome (IBS) is a functional GI disorder characterized by abdominal pain and altered bowel habits in the absence of a specific and unique organic pathology, rarely associated with microscopic inflammation. Studies have revealed that IBS is a disorder that affects all ages, although most patients have their first symptom before the age of 45. Older individuals have a lower reporting frequency. In south Asia and in India, most IBS reporting patients are young men. Almost all the symptoms of IBS are similar to that of Pravahika. Pravahika is a disease related to pakvashaya (colon) characterized by pravahana (straining) during defecation associated with shoola, dahaand vibaddhavatavcharas. The pureesha (stool) will be picchila (slimy), saphena (frothy), sakapha (mucus) and excreted in little quantity with frequent episodes.

For Pakvashayagatavikaras and vataja disorders, Bastichikitsa is considered to be one of the best methods of treatment. As IBS is a motility disorder related to colon (pakwashaya) which is chronic in nature and the main dosha involved in the pathogenesis of pravahika is vatadosha, this study was undertaken to evaluate the efficacy of picchabasti, followed by Kapittashtakachurna as shamanoushadhi (palliative treatment) in the Management of Irritable bowel syndrome vis-à-vis Pravahika.

A total number of 20 subjects completed the intervention. Subjective improvement in IBS patients with statistical analysis of results has been explained in clinical study. All the observations were recorded in a specially designed proforma.

**Aim and Objective**
To evaluate the combined effect of Picchabasti and Kapittashtakachurnaas shamanoushadhi in IBS vis-à-vis Pravahika.

**Materials & methods**

**Source of data**
Subjects were selected from the O.P.D. and I.P.D. of Government Ayurveda Medical College & Hospital, Mysuru and from the Special camp conducted for the study at Government Ayurveda Medical College & Hospital, Mysuru for 1 month.

**Sample size and Sampling method:**
A total of 23 subjects irrespective of gender, socio-economic status and religion, having the signs and symptoms of Irritable Bowel Syndrome vis-à-vis Pravahika fulfilling the inclusion criteria were registered for the study. The selected subject’s detailed profile was prepared as per the detailed proforma designed for the same purpose, which incorporates relevant data like symptomatology, physical signs, laboratory investigation reports as well as assessment criteria after taking informed Consent of the subject.

**Inclusion Criteria**
Patients with symptoms of IBS vis-à-vis pravahika viz.
- Abdominal pain associated with altered bowel habit that consists of constipation, diarrhoea or both. (Sashoolamalapravritti /Vibaddhamala pravritti).
- Mucus in stool (Sakapha mala pravritti).
Altered stool passage (straining, urgency or feeling of incomplete evacuation/ kunthanasheela pravritti/ Kruteapiakruta sangya).

Cases with or without dyspepsia, nausea, vomiting, flatulence were taken for the study.

Patients between the age group of 16-60 years were selected.

Patients of all gender, religion, occupation were selected for the study.

Both fresh and treated cases were taken for the study.

Exclusion Criteria

Patients suffering from any other systemic disorders which interfere with the course of the intervention were excluded.

Patients with complications of IBS like haemorrhoids, depression and weight loss were excluded.

Patients with upadrava of Pravahika like Gudabramsha (rectal prolapse), Gudapaka (ano-rectal inflammation), Gudashotha (rectal swelling), Gudenaraktasrava (rectal bleeding) were excluded.

Pregnant and lactating women were excluded.

Diagnostic Criteria

Rome III criteria for functional bowel disorders.

The diagnosis was made based on the Rome III criteria. Recurrent abdominal pain or discomfort lasting at least 3 days a month in the last 3 months associated with any two below mentioned features

- Relieved with defecation
- Onset associated with a change in frequency of stool
- Associated with a change in form (appearance) of stool

Symptoms that support the diagnosis of IBS

- Altered stool frequency (may be defined as greater than 3 bowel movements/day and less than 3 bowel movements/week).
- Altered stool form (lumpy/hard or loose watery stool)
- Altered stool passage (straining, urgency or feeling of incomplete evacuation- kunthanasheela, Sashoola mala pravritti, Kruteapiakrutasaangya).
- Passage of mucus – Sakapha/saphena mala pravritti.
- Bloating or feeling of abdominal distension.

Research design

The present study was an observational clinical trial with Pre and Post-test design.

Intervention:

The interventions were as follows.

1. Picchabasti³ in yogabasti pattern for first 8 consecutive days.

   • Anuvasana-PippalyadiTaila⁴
   • Niruha – madhu (honey)- 30gms, siandhava lavana (rock salt)- 5gms, pippalyadi-tail-70ml, yashtimadhu (glycyrrhiza glabra) kalka- 12gms, mocharasa (Salmalia malabarica) ksheerapaka- 350ml

2. Kapittashtakachurna⁵ 12gms in two divided doses of 6gms in the morning and 6gms in the night after food with takra (butter milk) as anupana (drink to be given
along with medicine) for 22 days from 9th day of intervention.

Period of intervention- 30 days

Assessment -
Assessment schedule
In this Study, total three assessments of the subjects were done. Pre-test assessment was done on 0th day, next assessment was done on 8th day (after basti therapy) and post-test assessment i.e. after the completion of intervention was done on 31st day

Statistical methods
The results were analysed statistically by using Cross tabulation analysis using Service product for statistical solution (SPSS) for windows software.

Investigations:
Haematological investigations namely Haemoglobin %, TC, DC, ESR, Random blood sugar, and Urine examinations namely Sugar, Albumin &Microscopic, Stool for Ova, Cyst and Microscopic were carried out to rule out other systemic diseases in all the cases.

OBSERVATION AND RESULTS
Pre test observation;
Out of 20 patients, in pre- test assessment, 1(5.0%) subject had Continuous abdominal pain not relieved by passage of flatus & stool, 19(95.0%) had intermittent lower abdominal pain relieved by passage of flatus. 1(5.0%) subject had No tenesmus, 2(10.0%) subjects had occasional tenesmus, 11(55.0%) subjects had tenesmus 1-2 times/day and in 6(30.0%) subjects tenesmus was frequently present. 17(85.0%) subjects had No constipation, in 3 (15.0%) subjects constipation was once in 2 days. 5 (25.0%) subjects had Passage of large amount of mucus in stool, 12 (60.0%) subjects had Passage of mucus with frequent stool, 3 (15.0%) subjects had visible mucus in stool. 9 (45.0%) subjects had Rumbling / Gurgling sound in abdomen, 11 (55.0%) subjects had Frequent abdominal distension with increased Flatulence & belching. In 4 (20.0%) subjects, frequency of defecation was > 6 times a day, in 15 (75.0%) subjects frequency of defecation was 4-6 times a day, in 1 (5.0%) subject frequency of defecation was 2-4 times day.

Observation after basti therapy
Out of 20 patients, in mid- test assessment, 3(15.0%) subjects had intermittent lower abdominal pain, 11 (55.0%) subjects had occasional abdominal pain, 6 (30.0%) subjects had no abdominal pain. In 1 (5.0%) subject tenesmus was frequently present, in 4(20.0%) subjects tenesmus was 1-2 times/day, 10(50.0%) subjects had occasional tenesmus and 5(25.0%) subjects had no tenesmus. 3 (15.0%) subjects had alternative day constipation and 17(85.0%) had no constipation. 2 (10.0%) subjects had passage of large amount of mucus in stool, 3(15.0%) subjects had passage of mucus with frequent stool, 10(50.0%) subjects had passage of visible mucus with stool and 5(25.0%) subjects had no visible mucus in stool. 1(5.0%) subject had rumbling or gurgling sound in abdomen, 6(30.0%) subjects had frequent abdominal distension with increased flatulence and belching, 9(45.0%) subjects had occasional abdominal distension and 4 (20.0%) subjects had no abdominal flatu-
ulence. In 3(15.05) subjects frequency of defecation was 4-6 times/day, in 9(45.0%) subjects frequency of defecation was 2-4 times/day and in 8(40.0%) subjects frequency of defecation was twice daily.

**Post test assessment:**
In Post-test assessment, 1(5.0%) subject had Occasional abdominal pain and 19(95.0%) subjects were completely relieved from abdominal pain.17(85.0%) subjects had no tenesmus, 2 (10.0%) subjects had occasional tenesmus, and in 1(5.0%) subject tenesmus was 1-2 times a day.20 (100.0%) subjects had no constipation, 0 (0.0%) subjects had alternative day constipation.17(85.0%) subjects had no visible mucus in stool, 2(10.0%) subjects had visible mucus in stool and 1 subject had Passage of mucus with frequent stool.15(75.0%) subjects had No abdominal distension, 4(20.0%) subjects had Occasional abdominal distension, 1 (5.0%) subject had frequent abdominal distension with increased Flatulence & belching. In 19(95.0%) subjects frequency of defecation was once daily, In 1(5.0%) subject frequency of defecation was twice daily.

**Table 1:** Showing results on abdominal pain/discomfort

<table>
<thead>
<tr>
<th>Day</th>
<th>Abdominal pain/discomfort</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No abdominal pain</td>
<td>Occasional abdominal pain</td>
</tr>
<tr>
<td>0th day</td>
<td>0(0.0%)</td>
<td>0(0.0%)</td>
</tr>
<tr>
<td>8th day</td>
<td>6(30.0%)</td>
<td>11(55.0%)</td>
</tr>
<tr>
<td>31st day</td>
<td>0(0.0%)</td>
<td>0(0.0%)</td>
</tr>
</tbody>
</table>

The result on abdominal pain/discomfort showed statistically highly significant result with p value 0.000

**Table 2:** Showing the results on Tenesmus

<table>
<thead>
<tr>
<th>Day</th>
<th>Tenesmus</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No tenesmus</td>
<td>Occasional tenesmus</td>
</tr>
<tr>
<td>0th day</td>
<td>1(5.0%)</td>
<td>2(10.0%)</td>
</tr>
<tr>
<td>8th day</td>
<td>5(25.0%)</td>
<td>10(50.0%)</td>
</tr>
<tr>
<td>31st day</td>
<td>1(5.0%)</td>
<td>0(0.0%)</td>
</tr>
</tbody>
</table>

The result on Tenesmus showed statistically highly significant result with p value 0.000

**Table 3:** Showing the results on Constipation

<table>
<thead>
<tr>
<th>Day</th>
<th>Constipation</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No Constipation</td>
<td>Alternative day constipation</td>
</tr>
<tr>
<td>0th day</td>
<td>17(85.0%)</td>
<td>0(0.0%)</td>
</tr>
</tbody>
</table>
The result on constipation showed statistically significant result with $p$ value 0.015

**Table 4:** Showing the results on Mucus in stool

<table>
<thead>
<tr>
<th>Day</th>
<th>Mucus in stool</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No visible mucus in stool</td>
<td>visible mucus in stool</td>
</tr>
<tr>
<td>0th day</td>
<td>0(0.0%)</td>
<td>3(15.0%)</td>
</tr>
<tr>
<td>8th day</td>
<td>5(25.0%)</td>
<td>10(50.0%)</td>
</tr>
<tr>
<td>31st day</td>
<td>17(85.0%)</td>
<td>2(10.0%)</td>
</tr>
</tbody>
</table>

The result on Mucus in stool showed statistically highly significant result with $p$ value 0.000

**Table 5:** Showing the results on Flatulence

<table>
<thead>
<tr>
<th>Day</th>
<th>Flatulence</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No abdominal distension</td>
<td>Occasional abdominal distension</td>
</tr>
<tr>
<td>0th day</td>
<td>0(0.0%)</td>
<td>0(0.0%)</td>
</tr>
<tr>
<td>8th day</td>
<td>4(20.0%)</td>
<td>9(45.0%)</td>
</tr>
<tr>
<td>31st day</td>
<td>15(75.0%)</td>
<td>4(20.0%)</td>
</tr>
</tbody>
</table>

The result on Flatulence showed statistically highly significant result with $p$ value 0.000

**Table 6:** Showing the results on increased frequency of defecation

<table>
<thead>
<tr>
<th>Day</th>
<th>Increased frequency of defecation</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Normal once daily</td>
<td>twice daily</td>
</tr>
<tr>
<td>0th day</td>
<td>0(0.0%)</td>
<td>0(0.0%)</td>
</tr>
<tr>
<td>8th day</td>
<td>0(0.0%)</td>
<td>8(40.0%)</td>
</tr>
<tr>
<td>31st day</td>
<td>19(95.0%)</td>
<td>1(5.0%)</td>
</tr>
</tbody>
</table>

The result of increased frequency of defecation showed statistically highly significant result with $p$ value 0.000.

**DISCUSSION**

Ingredients of *pippalyaditaila* possess *deepana* (appetizer), *pachana* (digestive), *vatanulomana* (carminative), *grahi* (absorbent) and *shoolahara* (analgesic) properties. *Taila* (Sneha) in general is Vatahara, removes the obstruction produced by *mala*. Owing to the *SnigdhaGuna* (unctuous nature), it produces
unctuousness in the body and by Sukshma-Guna (minuteness) it helps the potency of the drug to reach into the micro channels. Mocharasa is having kashaya(astringent) rasa, snigdha (unctuous), sheeta (cold), grahi (absorbent) and sthambanaguna (with-holding property). Ksheera (milk) acts as vata pitta shamaka, rasayana, balya by virtue of its properties and when processed with mocharasa it acts as stambhaka, grahi, vatanulomaka and hence helps in the management of Pravahika. Mocharasa also possesses the properties like antibacterial and antimicrobial, thus helps in managing post infectious IBS.

The main ingredient of kapittashtatakachurna is phalamajja (fruit pulp) of unripen kapitta fruit (Limonia acidissima). The phalamajja of unripen fruit is used here because it possesses Grahi and sthambaka action. Other ingredients of this yoga have deepana, pachana, vatanulomana, shoola prashamana and tridoshashamaka properties.

CONCLUSION

Pravahika is a vatakaphapradhanavyadhi characterised by pravahana (straining), sakapha mala pravritti (mucus in stool), alpalpa mala pravritti (little quantity of stool) and muhurmuhur mala pravritti (increased frequency of stools). All these correlates with the symptoms of IBS. It is clear from the observation made on 20 subjects that, maximum number of subjects had history of irregular food habits, which can be considered under ahitashana (improper diet), which on long run vitiates vata and kapha doshas. So the piccha basti chikitsa helps in treating vata as well as the vatasthanagata kapha. The overall assessment has shown that; Complete relief was found in 7 (35.0%) subjects, marked improvement was found in 5 (25.0%) subjects, Moderate improvement was found in 2(10.0%) subjects, Mild improvement was found in 5(25.0%) subjects, No improvement was found in 1(5.0%) subject. From the results obtained it can be concluded that both the interventions used in the study are effective in the management of IBS vis-à-vis Pravahika. If the same line of treatment is continued in more number of subjects, effectiveness of the intervention can be assessed accurately.

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