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A Study on Aetiopathogenesis of *Anartava* w.s.r to PCOS and its *Upashayatmaka Adhyayana* with *Tila Kwatha*

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ABSTRACT

Anartava is absence of artava, that is may be antar-artava(ovum) or bahir-artava(menstrual blood). The process of releasing ovum is known as ovulation, after ovulation if it is not fertilized then it flows out with cyclic menstrual blood. In Anartava this normal process of ovulation and bleeding hampers and we call it as anovulation, but it is not mandatory that both the processes must hamper together. This anovulation condition is one of the diagnostic criteria of PCOS and is one of the burning problem concerning the women health. In India, nearly 1 million women diagnosed with PCOS yearly⁵. In Bhaishajya Ratnavali, Tila kwatha is explained to induce menstruation in rakta gulma rogi¹, under the context of Gulma Chikitsa. In present study Tila kwatha is selected to restart the menstruation in PCOS patients. Percentage of improvement in Group B on amenorrhea is 81%, acne vulgaris is 65% and USG is 77% and significant comparatively. Thus it can be concluded that Tila kwatha as Upashaya in the management of Anartava will enhance the quality of treatment.

Key Words Anartava, Tila kwatha, Anovulation, Artava vyapad, Upashaya

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INTRODUCTION

Nirukti of artava is "Rutou bhavam artavam"² that means a substance which flows out at specific time. Anartava explained as "Ayogya rutushu utpannah"³ means, which may not be seen in proper period or unseasonable. So simply here in this study Anartava is a condition of menstrual irregularities and its side effects.

OBJECTIVES

To know the benefits of *Upashaya* dravya through the use of *Tila kwatha* in *Anartaya* w.s.r to PCOS.

To understand the aetiopathogenesis of *Anartava* w.s.r to PCOS in modern era.

MATERIALS AND METHODS

Female patients with 2-3 classical symptoms of *Anartava* w.s.r to PCOS were selected randomly from OPD & IPD of Shri Jayachamarajendra institute of Indian medicine hospital, Bengaluru without bias of social, economic, educational or religious status.

STUDY DESIGN





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| Group A | Group B | |
|--|--|--|
| Analyses of the lakshanas of PCOS to | Analyses of the lakshanas of PCOS to | Analyses of the lakshanas of PCOS to |
| elicit the aetiopathogenesis in selected | elicit the aetiopathogenesis in selected | elicit the aetiopathogenesis in selected |
| patient | patient | patient |
| Advise to take Tila kwatha, prepared | Advise Tila kwatha as Upashaya | Advise placebo medicine prepared by |
| by given tila choorna | dravya along with medicines prescribed | rice powder |
| | by gynecologist | |
| Observation of changes in the patient | Observation of changes in the patient | Observation of changes in the patient |
| condition | condition | condition |

Assessment criteria:

Patients who has 2 or more than 2 parameters including both subjective or objectives criteria were selected.

| Subjective parameters: | Objective parameters: |
|------------------------|--|
| Amenorrhea | Abdominal obesity |
| Infertility | Acne vulgaris |
| Weight gain | Skin thickening and darkening (at nape of neck & inner thighs) |
| Pimples | USG or TVS showing multiple cyst appearance in ovaries |
| Hircuticm | |

DIAGNOSTIC CRITERIA:

Signs and symptoms of PCOS

USG and TVS showing polycystic ovaries

Assessment protocol:

| Symptoms : (Subjective) | Grade | Score |
|--|----------------|-------|
| Amenorrhea- Absent | G_0 | 0 |
| Amenorrhea- Asymptomatic, Occasionally | G_1 | 1 |
| Amenorrhea- Symptomatic, since 6 months | G_2 | 2 |
| Oligomenorrhea-3days/25-30days | G_0 | 0 |
| Oligomenorrhea-2days/25-30days | G_1 | 1 |
| Oligomenorrhea-1day/25-30days | G_2 | 2 |
| Acne vulgaris- Absent | G_0 | 0 |
| Acne vulgaris- <15 Inflammatory Lesions | G_1 | 1 |
| Acne vulgaris- 15 to 50 Inflammatory Lesions | G_2 | 2 |
| Weight gain- As before | G_0 | 0 |
| Weight gain- Up to 5kgs | ${\sf G}_1$ | 1 |
| Weight gain- 5-10kgs | G_2 | 2 |
| Hirsutism- FG score < 8 | G_0 | 0 |
| Hirsutism- FG score = 8 | G_1 | 1 |
| Hirsutism- FG score 8-16 | G_2 | 2 |
| Infertility- Absent | G_0 | 0 |
| Infertility- Secondary | G_1 | 1 |
| Infertility- Primary | G_2 | 2 |

| Symptoms: (Objective) | Grade | Score |
|--|----------------|-------|
| Abdominal Obesity- 30.5 to 33.5 inches of abdominal circumference | G_0 | 0 |
| Abdominal Obesity- > 33.5 inches of abdominal circumference | G_1 | 1 |
| Abdominal Obesity- > 40 inches of abdominal circumference | G_2 | 2 |
| Skin Thickening and Darkening- Smooth to touch, | G_0 | 0 |
| Skin Thickening and Darkening- Rough to touch | G_1 | 1 |
| Skin Thickening and Darkening- Coarseness can be observed visually | G_2 | 2 |
| USG/TVS- Normal Pelvic scan | G_0 | 0 |
| USG/TVS-Presence of < 12 immature follicles 2-9 mm in | G_1 | 1 |
| diameter. | | |
| USG/TVS- Presence of 12 or more immature follicles 2-9 mm in | G_2 | 2 |
| diameter or an increased ovarian volume > 10 ml without | | |
| a cyst or dominant follicle in either ovary. | | |





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OBSERVATIONS

Table 1 Distribution of Anartava patients according to observation parameters

| PARAMETER | OBSERVATION | PERCENTAGE |
|------------------------------|-------------|------------|
| Age | 16-26 years | 51% |
| Occupation | House wife | 44% |
| Agni dushti | Vishamagni | 47% |
| Addiction | Tea | 56% |
| BMI | 25-30 | 67% |
| Chinta as prime nidana | In 28 cases | 62% |
| Abhishyandi & guru ahara | 31 cases | 69% |
| Ruksha & katurasayukta ahara | 21 cases | 47% |
| | | |

RESULTS

Effect on Amenorrhea

Effect of tila kwatha as upashaya on amenorrhea in group A is given in detail in Table No.02. Statistical analysis showed that the mean score which was 1.60 before the treatment was reduced to 0.67 after the treatment with 58% improvement and there is a statistically highly significant. (P<0.001)

Tila kwatha effect in Group-B on Amenorrhea revealed in detail in Table No.02. Statistical

analysis showed that the mean score which was 1.73 before the treatment was reduced to 0.33 after the treatment with 81% improvement and there is a statistically highly significant change. (P<0.001) Group-C changes in Amenorrhea revealed are given in detail in Table No.02. Statistical analysis showed that the mean score which was 1.53 before the treatment was not reduced to after the treatment with 0% improvement.

Table 2 Effect on Amenorrhea of ANARTAVA in Group-A,B&C

| AMENORRHEA | Mean sc | ore | | % | S.D (±) | S.E (±) | t value | p value |
|------------|---------|------|-------|----|---------|---------|---------|---------|
| | BT | AT | BT-AT | | | | | |
| Group A | 1.60 | 0.67 | 0.93 | 58 | 0.458 | 0.118 | 4.09 | < 0.001 |
| Group B | 1.73 | 0.33 | 1.40 | 81 | 0.514 | 0.133 | 8.10 | < 0.001 |
| Group C | 1.53 | 1.53 | 0.00 | 0 | 0.000 | 0.000 | 0 | 0 |

Table 3 Effect on Acne vulgaris of ANARTAVA in Group-A, B&C

| Acne Vulgaries | Mean se | core | | % | S.D (±) | S.E (±) | t value | p value |
|----------------|---------|------|-------|----|---------|---------|---------|---------|
| | BT | AT | BT-AT | | | | | |
| Group A | 0.60 | 0.40 | 0.20 | 33 | 0.41 | 0.107 | 1.08 | >0.05 |
| Group B | 1.13 | 0.40 | 0.73 | 65 | 0.47 | 0.121 | 3.47 | < 0.001 |
| Group C | 0.60 | 0.60 | 0.00 | 0 | 0.00 | 0.000 | 0 | 0 |

Effect on Acne Vulgaries

In this work of *Anartava* 15 patients were studied with Group-A, **changes in Acne Vulgaries** revealed in detail in Table No.03. Statistical analysis showed that the mean score which was 0.60 before the treatment was reduced to 0.40 after

the treatment with 33% improvement and there is a statistically not significant. (P>0.05)

Changes in Group-B in-concern with **Acne Vulgaries** given in detail in Table No.03.
Statistical analysis showed that the mean score which was 1.13 before the treatment was reduced

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to 0.40after the treatment with 65% improvement and there is a statistically highly significant change. (P<0.001)

Group-C **Acne Vulgaries changes** revealed are given in detail in Table No.03. Statistical analysis

showed that the mean score which was 0.60 before the treatment was not reduced to after the treatment with 0% improvement.

Table 4 Effect on Skin Thickening and Darkening of ANARTAVA in Group-A,B&C

| Skin Thickening | and | Mean sc | ore | | % | S.D (±) | S.E (±) | t value | p value |
|-----------------|-----|---------|------|-------|-----|---------|---------|---------|---------|
| Darkening | | BT | AT | BT-AT | _ | | | | |
| Group A | | 0.40 | 0.33 | 0.07 | 17 | 0.258 | 0.067 | 0.36 | >0.05 |
| Group B | | 0.40 | 0.00 | 0.40 | 100 | 0.497 | 0.128 | 3.05 | < 0.05 |
| Group C | | 0.53 | 0.53 | 0.00 | 0 | 0.00 | 0.000 | 0 | 0 |

Effect on Skin Thickening and Darkening

Changes in **Skin Thickening and Darkening of** Group-A **patients** revealed in detail in Table No.04. Statistical analysis showed that the mean score which was 0.40 before the treatment was reduced to 0.33 after the treatment with 17% improvement and there is a statistically no significant change (P>0.05). Effect of tila kwatha in Group-B on **Skin Thickening and Darkening** given in detail in Table No.04. Statistical analysis

showed that the mean score which was 0.40 before the treatment was reduced to 0.00 after the treatment with 100% improvement and there is a statistically significant change (P<0.05). Group-C **Skin Thickening and darkening changes** revealed in detail in Table No.04. Statistical analysis showed that the mean score which was 0.53 before the treatment was not reduced to after the treatment with 0% improvement.

Table 5 Effect on USG/TVS of Anartava in Group-A

| USG/TVS | Mean sc | ore | | % | S.D (±) | S.E (±) | t value | p value |
|---------|---------|------|-------|----|---------|---------|---------|---------|
| | BT | AT | BT-AT | | | | | |
| Group A | 2.00 | 1.80 | 0.20 | 10 | 0.414 | 0.107 | 1.87 | < 0.05 |
| Group B | 1.73 | 0.40 | 1.33 | 77 | 0.617 | 0.159 | 6.61 | < 0.001 |
| Group C | 1.67 | 1.67 | 0.00 | 0 | 0.000 | 0.000 | 0 | 0 |

Effect on USG/TVS

In this work of 15 patients studied in *Anartava* with Group-A **USG/TVS** revealed are given in detail in Table No.05. Statistical analysis showed that the mean score which was 2.00 before the treatment was reduced to 1.80 after the treatment with 10% improvement and there is a statistically significant change. (P<0.05)

Group-B on **USG/TVS** revealed is given in detail in Table No.05. Statistical analysis showed that

the mean score which was 1.73 before the treatment was reduced to 0.40 after the treatment with 77% improvement and there is a statistically significant change. (P<0.001)

Group-C **USG/TVS** revealed are given in detail in Table No.05. Statistical analysis showed that the mean score which was 1.67 before the treatment was not reduced to after the treatment with 0% improvement.





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ASSESSMENT OF TOTAL EFFECT OF THERAPY

Table 6 Overall effect of Group-A

| EFFECT OF TREATMENT IN GROUP - A | | | | |
|----------------------------------|----------------------|----------------|--|--|
| Class | Grading | No of patients | | |
| 0-24% | No Improvement | 9 | | |
| 25% -49% | Mild Improvement | 1 | | |
| 50% - 74% | Moderate Improvement | 5 | | |
| 75% - 100% | Marked Improvement | 0 | | |

Table 7 Overall effect of Group-B

| EFFECT OF TREATMENT IN GROUP – B | | | |
|----------------------------------|----------------------|----------------|--|
| Class | Grading | No of patients | |
| 0-24% | No Improvement | 0 | |
| 25% -49% | Mild Improvement | 0 | |
| 50% - 74% | Moderate Improvement | 7 | |
| 75% - 100% | Marked Improvement | 8 | |

Table 8 Overall effect of Group-C

| EFFECT OF TREATMENT IN GROUP - C | | | | | |
|----------------------------------|----------------------|----------------|--|--|--|
| Class | Grading | No of patients | | | |
| 0-24% | No Improvement | 0 | | | |
| 25% -49% | Mild Improvement | 0 | | | |
| 50% - 74% | Moderate Improvement | 0 | | | |
| 75% - 100% | Marked Improvement | 0 | | | |

Table 9 Comparative results of Group-A, Group-B and Group-C (Symptom wise)

| Characteristics | Group-A | | | Group-B | | | Group-C | | |
|---------------------|---------|-------|------------------|------------|------|------------------|------------|------|------------------|
| Signs and Symptoms | Mean | score | % of — relief | Mean score | | % of — relief | Mean Score | | % of — relief |
| | BT | AT | | BT | AT | — Tener | BT | AT | — Tellel |
| AMENORRHEA | 1.60 | 0.67 | 58 | 1.73 | 0.33 | 81 | 1.53 | 1.53 | 0 |
| ACNE VULGARIS | 0.60 | 0.40 | 33 | 1.13 | 0.40 | 65 | 0.60 | 0.60 | 0 |
| HIRSUITISM | 0.20 | 0.20 | 0 | 0.47 | 0.20 | 57 | 0.40 | 0.40 | 0 |
| ABDOMINAL OBESITY | 0.60 | 0.47 | 22 | 0.47 | 0.20 | 57 | 0.67 | 0.67 | 0 |
| SKIN THICKENING AND | 0.40 | 0.33 | 17 | 0.40 | 0.00 | 100 | 0.53 | 0.53 | 0 |
| DARKENING | | | | | | | | | |
| USG/TVS | 2.00 | 1.80 | 10 | 1.73 | 0.40 | 77 | 1.67 | 1.67 | 0 |

Result of group A

The percentage of improvement in Group A on Amenorrhea is 58%, Acne Vulgaris is 33%., Hirsuitism is0%, Abdominal Obesity is 22%, Skin thickening and Darkening is 17% and USG/TVS is 10%.

Result of group B

The percentage of improvement in Group B on Amenorrhea is 81%, Acne vulgaris is 65%., Hirsuitism is 57%, Abdominal Obesity is 57%,

Skin thickening and Darkening is 100% and USG/TVS is 77%.

Result of group C

The percentage of improvement in Group B on Amenorrhea is 0%, Acne Vulgaris is 0%., Hirsuitism is 0%, Abdominal Obesity is 0%, Skin thickening and Darkening is 0% and USG/TVS is 0%.

Table No. 10: Comparative results of Group A and Group B(Overall)

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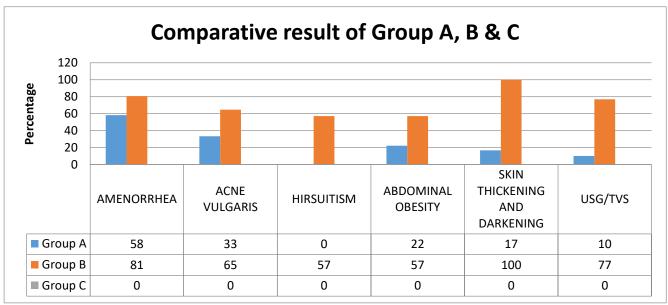




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Comparative analysis of the overall effect of the treatments in both the groups was done by statistically with paired t test. The test shows that the treatment is statistically significant in Group B

when compared to Group A. Group C is not getting any value to compare their two groups. Group A overall result is 30.03% and Group B overall result is 76.43%.



Graph No. 01 Comparative result of Group A, B & C

DISCUSSION

DISCUSSION ON RESULTS:

Marked response was found in 8 cases of group B only. Moderate response was observed in 7 cases of group B and 5 cases of group A. Mild response was observed in 1 patient of group A. Poor response was observed in 9 cases of group A and 15 cases of group C.

Highly significant difference was observed with amenorrhea in the first 2 groups (Group A and Group B) and between the groups significant difference was observed in USG. No significant difference was observed in between groups in abdominal obesity.

Based on the overall response, there was 76.43%% overall results in group B which was administered the *Tila kwatha* along with gynecological

medicines and 30.30% overall results in group A which was kept on only *Tila kwatha*.

This clearly states the superiority of *Upashaya* (*Tila kwatha*) in the management of disease (*Anartava*) and showed that it enhance the quality of treatment and make it successful.

DISCUSSION ON UPASHAYA:

Upashaya is just the reverse of the Nidana (cause) of the disease. Anything whatsoever that relieves or tends to relieve the patient of his suffering is Upashaya. The knowledge of Upashaya has a greater importance in the treatment of a disease rather than its ascertainment, but its value in diagnosing a disease cannot altogether be ignored. Here the Upashaya taken for the study is Hetu vipareeta. Tila kwatha is a combination of Tila, Pippali, Shunti, Maricha, Bharangi, Hingu, Guda



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whose properties balance each other and enhance the *vata kapha hara*⁴ properties seen in all of them. Most of the drugs have *katu tikta pradhana rasa*, *deepana*, *pachana*⁴ properties which do *agni deepana*, prevent further formation of *aama* and does *pachana* of already formed *aama*, thereby controlling the root cause of the disease. Once *saamata* of *doshas* is relieved, they gain their originality, start functioning normally and thus symptoms get relieved.

Most of the drugs are *Ushna veerya yukta*⁴ and act against the *sheeta pradhayata* of both *Vata* and *Kapha* and thus by performing their *Agni guna pradhana* action, they help in *utpatti of artava* by the action of "*VRIDDHI SAMANAI*". They also act as *Vatanulomaka* and *Kapha nissaraka*.

Few drugs have property which helps in removing sroto sangha caused by aama and mala rupi kapha. Few are Artava janaka⁴, shula prashamana⁴, krumighna⁴ and they act accordingly. Few also have Rasayana property which help in overall improvement.

Numerous research works done on these drugs have proven their blood purifier, lipophilic antioxidant, Anti-tumor, Anti-inflammatory and bioavailability enhancer effects which are all of helps to relieve the symptoms of *Anartava*.

CONCLUSION

Anartava is a disease where in Kapha and Vata play a major role in the pathogenesis along with aama.

- ❖ Disturbance in the *Agni* is first and foremost factor which leads to Production of *aama*, *Dosha dushti* and *Sroto dushti*.
- ❖ 45 patients who were selected for the study were grouped into 3 − Group A, Group B and C, where in group A was administered with *Tila kwatha*. Group B was administered with *Tila kwatha* along with gynecological medicines. Group C kept only on starch powder.
- Most of the *nidanas* explained in the texts were observed even clinically.
- ❖ Clinically Agni vikriti, Laghu bhojana, Ati bhojana, Adhyashana were also observed which substantiates the involvement of anna vaha srotas in the disease.
- ❖ Sthoulya, Ati santarpana, Dhatu kshaya substantiate the involvement of Rasavaha, Raktavaha, Medavaha srotas.
- Abhishyandhi, Guru, Atikatu ahara sevana are observed as prime aharaja nidanas while Diwaswapna, Avyayama has been observed as a main viharaja nidana.
- Chinta is one of main manasika nidana and sthoulya is anya nidana observed in this study
- ❖ Group B which was administered with *Tila kwatha* and Gynecological medicines showed highly significant results in Amenorrhea and Acne-vulgaris and significant results in Skin thickening and darkening and USG/TVS.
- Thus it can be concluded that *Tila kwatha* along with gynecological medicines play a significant role in the treatment of *Anartava/PCOD*.





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