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STUDIES ON MEDADUSTI IN PANDU W.S.R. TO ITS THERAPEUTIC MEASURES WITH SARAPUNKHA

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ABSTRACT

Introduction: Pandu is a disease mentioned in *Ayurveda* that can be correlated with anaemia. In *Pandu*, diminution of *Rakta Dhatu* (blood) & *Meda Dhatu* (fatty tissue and lipids) occurs. Due to intake of several causative factors, *Pitta Dosha* gets vitiated, due to which *Agni* (digestive fire) gets deranged, as a result of which *Ahara Paka* (the process of digestion of food) & *Dhatu Paka* (the process of metabolism) are also get altered which ultimately results in a diminution of *Rakta & Meda Dhatu*. *Meda Dhatu* comprises of two parts - *Sthayi Meda* (can be compared with adipose tissues) and *Drava Meda* (can be compared with circulatory lipids). In context with *Pandu* i.e., anaemia, the state of hypolipidemia should be evaluated and in context to *Meda Kshaya* in *Pandu* the decreased level of HDL should be the matter of concern and that hypothesis was verified through clinical study in the present study. **Aim & Objectives:** The present study was carried out to evaluate the role of *Meda Dhatu* in the pathogenesis of *Pandu* and to reveal the effectiveness of the stipulated drug i.e., *Sarapunkha* to combat *Pandu* as well as *Meda Dusti*. **Materials and Methods:** In selected 60 patients of *Pandu* having the *Medadusti Lakshan* based

on inclusion and exclusion criteria, the powder of the root of *Sarapunkha* (*Tephrosia purpurea Linn*) was administered in stipulated dose (6 grams in divided doses with plain warm water) for 90 consecutive days.

Observations & Results: The subjective parameters for *Meda Kshaya* are clinically present in a maximum number of *Pandu* patients. The result also reveals the significant efficacy of *Sarapunkha* on relevant subjective and objective parameters with a 'p-value <0.001 in the majority of subjective and objective parameters. **Conclusion:** In *Pandu,* the pathological phenomenon of occurrence of *Alpa Rakta* and *Alpa Meda* is present in the study sample which has been verified through subjective parameters & biochemically. The patients who are suffering from Pandu can be treated with the drug capable to correct *Meda Kshaya* like *Sarapunkha*.

Keywords: Pandu, Medadusti, Meda Kshaya, Sarapunkha

INTRODUCTION

Pandu is a disease that is clinically characterized by the generalized pallor of the skin& mucous membrane, produced as a resultant effect of *Dhatu Kshaya* (depletion of body tissues) specially *Rakta Kshaya* (depletion of blood) and *Meda Kshaya* (depletion of fatty tissue). ⁽¹⁾ *Pandu* can be clinically co-related with anaemia. Globally, anaemia affects 1.62 billion people which corresponds to 24.8 % of the population. The highest prevalence is in preschool-age children (47.4%) and the lowest prevalence is in men (12.7%). The population group with the greatest number of individuals affected is non-pregnant women that is 468.4 million. In India anaemia affects10 billion cases/year.⁽²⁾

In Charaka Samhita, very categorically it had shown that Meda Dusti (affliction of Meda Dhatu by vitiated Doshas) is an essential factor to cause Pandu. Pandu and Meda Dusti are interrelated as it is mentioned in Charaka Samhita that there are Kshaya of Rakta and Meda Dhatu takes place in the disease of Pandu. Grahani Dosa i.e., mal absorption is the main causative factor of Pandu. (3) It is the root cause of malnutrition in Pandu. In Charaka Samhita, the chapter of disease Pandu is described after the chapter of Grahani. There is a sequence of the pathogenesis of Pandu from Grahani Roga. According to scholar Chakrapani Dutta, the aggravation of Pitta in Grahani Roga constitutes a predominant factor in the causation of *Pandu*, therefore the description of the treatment of Pandu follows the treatment of Grahani Dosa.⁽⁴⁾ Mainly *Tikshnagni* is responsible for the aggravation of Pitta. Though Pitta is Ushna (hot) in

character, by the virtue of Drava Guna, it deactivates the function of Agni.⁽⁵⁾ In Charaka Samhita it has been emphasized that, in the disease Pandu along with a diminution of Rakta Dhatu, Meda Dhatu is also diminished.⁽⁶⁾ Hence, the patient becomes emaciated. Meda Dhatu is attributed in two forms, one is "Sthula Meda" i.e., "Sthayi Meda" or adipose tissue and the "Drava Meda" i.e., Circulating lipids. Hence, there is a chance of presenting Hypolipidemia in the patient of Pandu. This dictum Acharya Charaka has been taken as a hypothesis in the present study. (7) Pandu has been described as Rasa Pradoshaja Vikara; hence Rasa Dhatu gets involved in the initial stage of the disease. As Hridaya (heart)is the root of Rasavaha Srota, hence heart gets afflicted in Pandu Roga. (8) In Sushrut Samhita, Pandu Roga is described as a

In Sushrut Samhita, Pandu Roga is described as a sequence of Hrid Roga. In the treatise of Astanga Hridaya, the Pandu Roga Nidan is described after Udar Nidan. Scholar Arun Dutta has clarified the sequences of Dosha Sanghata in Udar Roga is theroot cause of Pandu.⁽⁹⁾ Also, some authors have mentioned Pandu as Upadrava of Udar Roga. In Madhava Nidan, the chapter of Pandu Roga is described after the Krimi Roga. Purishaja Krimi produces Pandu, hence Pandu Roga is described after Krimi Roga.⁽¹⁰⁾

The hypothesis of the study is based on the concept of "*Alpa Meda*" in *Pandu*. The study is based on the hypothesis, of whether the status of the circulatory lipids is reduced or not. In *Charaka Samhita*, it has been denoted that patient of Pandu suffer in diminution of *Rakta Dhatu* and *Meda Dhatu*. In *Sushruta Samhita* facts of diminution *of Rakta Dhatu* is described but status of *Meda Dhatu* is not mentioned. In *Astanga Hridaya*, *Acharya Vagbhatta* had followed the same conceptual principles of *Acharya Charaka*.

Meda definitely is the outcome of Dhatu Paka. Dhatu Paka takes place after effective Avastha Paka. If the Agni is altered and hypo functioning, then the process of Avastha Paka and Dhatu Paka is disturbed and as a result, the harmful effect of Nidan Sevana cannot be counteracted by the Agni and according to the type of Nidana Sevana the particular Doshas and Dhatus get affected. The Meda in Dhatu Paka stage is yielded from Meda Poshak fraction of Mamsa Dhatu under the governance of Mamsagni. ⁽¹¹⁾ In Agnimandya state, for the malfunctioning of both Jatharagni and Mamsagni formation of Meda also appears defective. In Pandu, it appears as Meda Kshava. Tvak(skin) gets afflicted in Pandu and as Tvak is the root of Mamsavaha Srota, (12) hence Mamsa is also afflicted. Aggravated Vata, Pitta andKapha causes various ailments being residing inMamsa i.e., Mamsavaha Srota. Following the hypothesis of transmission of Dhatu Paka (Kedarikulya Nyay) as expounded by scholar Chakrapani Dutta, certainly, the Mamsagni gets vitiated and Meda Poshak part inhibits *media* formation, $^{(13)}$ that's why the person becomes 'Alpa Rakta Alpa Medoska'.

The *Meda Dhatu* comprises two parts, one is the *Sthula Sthayi Dhatu* which remains in the pocket of fat in *Vapabaha* i.e abdominal fat. ⁽¹⁴⁾ The other part of *Meda* is *Drava Meda* which circulates through the body and different *Srotas* to provide nutrition to all the *Dhatus*. ⁽¹⁵⁾ The crude Meda may be corelated-with adipose tissue and the Drava Meda may be compared with circulatory lipids. So, the term *Poshan Krama* or nutrition to the Dhatus by Drava-Meda is likely to the function of circulatory lipids as lipids are present in every cell of the human body to construct the cell structure, storing energy, transporting hormones, digestion, insulation and protection. Adipose tissue is the body fat comprised of adipocytes, which is the loose tissue contains the stromal-

vascular fraction (SVF) of cells including pre adipocytes, fibroblast, vascular endothelial cells are a variety of immune cells such as adipose tissue macrophages. This information regards the connection of *Meda* with *Rasa, Rakta* and *Mamsa* and also support the concept of formation of *Oja* as adipose tissue produces cytokines.

The main function of adipose tissue is energy storage and regulation of body heat. Free fatty acids (FFAs) are liberated from lipoproteins by lipoprotein lipase (LPL) and enter the adipocyte, where they are reassembled into triglycerides by esterifying them into triglyceride by esterifying them onto glycerol. Human fat tissue contains about 87% lipids. In Pandu, the major pathological criteria that take place, are Rakta Kshaya and Meda Kshaya. The point of our discussion in this present study is Meda Kshaya i.e., likely to be correlated with hypolipidemia. Hypolipidemia is a state in which there is a decrease in plasma lipoprotein caused by primary and secondaryfactors refers to genetic and acquired hypolipidemia respectively. It is usually asymptomatic and diagnosed incidentally on routine lipid screening. Theterms hypolipidemia and hypocholesterolemia and hypo beta lipo proteinemia (HBL) are used interchangeably in the literature and refer to reducing plasma cholesterol. Most authors use total serum cholesterol (TC). Total cholesterol is the total amount of cholesterol in the blood. Total cholesterol includes low-density lipoprotein (LDL) and high-density lipoprotein (HDL). The LDL and HDL are known as bad cholesterol and good cholesterol respectively.

In context to *Meda Kshaya* in *Pandu* the decreased level of HDL should be the matter of concern and that hypothesis was verified through a clinical study. Hypolipidemia has been described in various types of chronic anaemia. ⁽¹⁶⁾ The exact cause of Hypolipidemia in an anaemic patient is not known. Some authors have been suggested that cholesterol deficiency leads to rigidity of erythrocytosis making them more prone for destruction. ⁽¹⁷⁾ *Rakta* provides nutrition to both the *Drava Meda* and *Sthula Meda*, render supported by the information of the physiology of blood where it is mentioned that blood boost up

the adipocytes and transport lipoprotein. Hence Rakta *Kshaya* is the root cause of *Meda Kshaya*.

In the above context, the present study was carried out with the following aims and objectives:

- 1. To study the diagnostic approach of *Pandu*.
- 2. To evaluate the role of *Meda* in pathogenesis of *Pandu*.
- 3. To reveal the effectiveness of the stipulated drug i.e., *Sarapunkha* to combat *Pandu* as well as *Meda Dusti*.

MATERIALS AND METHODS:

Study Settings:

The present study is an interventional, prospective, randomized clinical study with single group which has been treated with root powder of the plant *Sarapunkha (Tephrosia purpurea Linn.)* ⁽¹⁸⁾, 6 g / day in divided doses for ninety consecutive days.

Selection of Drug:

The powder of the root of Sarapunkha (Tephrosia *purpurea Linn.*) which was administered in patients, belongs to the family *Fabaceae*. It has *Tikta, Kasaya Rasa, Laghu, Ruksa, Tiksna Guna, Usna Virya, Katu Vipaka* andhaving *Karma* like *Kapha vata hara, Visaghna, Vranahara, Jvarahara, Krimighna* and *Rasayana*.⁽¹⁹⁾

Duration of Study:

The entire study was completed in two years.

Definition of Populations:

A small sample had been taken from the population of those who were suffering from *Pandu*, visiting the OPD & IPD of the Institute of Post Graduate Ayurvedic Education and Research at Shyamadas Vaidya Shastra Pith Hospital, Kolkata irrespective of their sex, occupation & religion. At first, the patients were selected based on subjective and objective criteria of *Pandu*, thereafter, signs and symptoms of *Meda Dusti* have been explored within them.

Inclusion Criteria:

- 1. Adult subjects of either sex between16-70 yrs. of age.
- 2. Presence of cardinal sign & symptoms of Pandu.
- 3. 3.Patients below haemoglobin level, incase of male 13.8g/dl & incase of female12.1g/dl.
- 4. Morphological characteristics of RBC sup-

porting nutritional anaemia including iron deficiency anaemia.

- 5. Patients below normal level of serum ferritin
- 6. Willingness to give written consent to participate in the study.
- 7. Patients those who are not receiving any other therapies except for research medicine.

Exclusion Criteria:

- 1. Severe anaemia where Hb% count is below 5mg/dl.
- 2. Anaemia introducing thalassemia, leukaemia, sickle cell anemia, aplastic anaemia, hereditary spherocytosis etc.
- 3. Any malignant condition, bone marrow disease presenting anemia.
- 4. Co-existing chronic diseases.
- 5. Patient receiving any other treatment for the same disease or any other systemic disease.
- 6. Anaemia due to Helminthiasis.

Sample Size:

A total of 70 patients were selected for the study of which about 10 patients were dropped out during the study. So, the complete study was carried out on 60 patients.

Methods of Data Collection:

The drug had been administered for 90 consecutive days for each patient and assessed after 90 days following the date of registration.

ScheduleofDataCollection:

In each group, the subject will be required at least 4 visits during studies:

- 1. Visit1–Screening & enrolment (baseline)
- 2. Visit 2 First follow-up (Day 30)
- 3. Visit3–Second followup (Day60)
- 4. Visit 4 Final follow-up (Day 90)

Subjective Parameters:

A. Subjective Parameters for Pandu: (20)

- 1. *Hatanala*. (Suppression of the power of digestion)
- 2. Durvala. (Weakness)
- 3. Annadvit. (Repugnance against food)
- 4. *Bhrama* (Feeling of Giddiness)
- 5. Shrama. (Fatigue)
- 6. Gatra Shula (Pain in the body)

- 7. Aruchi. (Anorexia)
- 8. Shunakshi Kota (Swelling of the orbital region)
- 9. *Shirna Loma* (Falling out of the small hairs of the body)
- 10. Hataprabha. (Losing of bodily luster)
- 11. Kopana. (irritability)
- 12. Nidralu. (Feeling of sleepy always)

B. Subjective Parameters for Meda Kshaya: (21)

- 1. Pleehabhi Vriddhi (Enlargement of spleen)
- 2. Ruksha Twak (Dryness of the skin)
- 3. *Medura Mamsa Prarthana*. (Craving for fatty meat)
- 4. *Karshya* (Emaciation)

5. *Sandhi Shunyata* (Feeling of emptiness of the joints/joint crepitus)

Objective Parameters:

- 1. Complete Blood Count (Hb%, TLC, DLC, ESR, MCV, MCH, MCHC, TIBC, Platelet)
- 2. Estimation of Serum Ferritin
- 3. Estimation of Serum Lipid Profile

statistical analysis:

The information gathered based on the observation made about various parameters was subjected to statistical analysis in terms of Mean, Standard Deviation (SD) and Standard Error (SE). Paired 't-test was carried out at P<0.05 and P<0.001. The obtained results were interpreted as -P<0.05 is significant & P<0.001 is highly significant.

Preparation of Clinical report File (CRF):

Prior to the administration of the drug, Informed

Consent Form (ICMR guideline) was filled by the willing patients. Clinical Report File (CRF) was prepared in which all the clinical and therapeutic data of individual patients along with dropout cases has been recorded.

OBSERVATIONS AND RESULTS:

The distribution of subjective parameters for Pandu among the 60 patients shows that cent percent of patients were suffering from the feature of Daurvalva & Hataprabha both. The next higher group is 96.6% were suffering from Hatanala & Annadvitboth. Other notable percentage are 93.3% & 91.6%, where patients were suffering from Srama & Sisiradwesi respectively (Table 1). Whereas the distribution of subjective parameters for Meda Kshaya among the patients of Pandu shows that 76.6%, 63.3%, 63.3% & 26.6% of patients are clinically assessed with the symptoms of Ruksha Tvak, Medura Mamsa Prarthana, Karshya and Sandhi Shunyata, hence satisfying the criteria of Meda Kshaya (Table 2). Statistical analysis of subjective and objective parameters in patients of Pandu as well as in subjective and objective parameters of Meda Kshaya in the patients of Pandu, before and after treatment shows that 'p-value <0.0001in the majority of the parameters, which indicates that the stipulated drug Sarapunkha is highly efficacious in the management of Pandu and also in combating Meda Kshaya in Pandu by its Tikta - Kasaya Rasa, Laghu - Ruksa -Tiksna Guna, Usna Virya &Katu Vipaka.

Sl.No.	SubjectiveCriteria	Noofpatients	Percentage (%)
1	Hatanala	58	96.6
2	Durvala	60	100
3	Annadvit	58	96.6
4	Shrama	56	93.3
5	Gatra Shula	48	80
6	Aruchi	53	88.3
7	SunakshiKota	8	13.3
8	Shirna Loma	15	25
9	Hataprabha	60	100
10	Kopana	15	25
11	Nidralu	6	10
12	Bhrama	12	20

Table 1: Distribution of subjective parameters of *Pandu* among the 60 patients:

Sl.No.	CriteriaofMedakshaya	No. of Patients	Percentage
1	PleehabhiVriddi	Nil	0
2	RukshaTvak	46	76.6
3	MeduraMamsaPrarthana	38	63.3
4	Karshya	38	63.3
5	SandhiShunyata	16	26.6

Table 2: Distribution of subjective parameters of Meda Kshaya in 60 patients of Pandu:

Table 3: Follow-up assessment of Subjective & Objective Parameters of Pandu before and after treatment:

Parameters	Mean BT	Mean AT	SD +/-	SE +/-	't' Value	'p-Value
Hatanala	2.18	0.75	0.83	0.16	9.1	< 0.001
Durvala	1.71	0.90	1.35	0.26	3.1	<0.05
Annadvit	2.04	0.63	0.56	0.107	13.08	< 0.001
Shrama	1.35	0.75	1.12	0.21	2.78	<0.05
Gatra Shula	1.86	0.9	0.57	0.10	8.76	< 0.001
Aruchi	1.34	0.37	0.55	0.105	9.21	< 0.001
SunakshiKota	2.86	2.5	0.41	0.12	3.0	< 0.05
Shirna Loma	1.57	0.93	0.68	0.19	6.38	< 0.001
Hataprabha	2.09	0.72	0.54	0.104	11.66	< 0.001
Kopana	2.86	2.5	0.47	0.12	3.0	< 0.001
Nidralu	2.56	0.91	0.663	0.12	13.75	<0.001
Bhrama	1.31	0.36	0.58	0.113	8.4	<0.001
Hb%	8.37	11.34	0.86	0.15	19.3	<0.001

Table 4: Follow-up assessment of Subjective & Objective Parameters of *Meda Kshaya* inpatients of *Pandu* before and after treatment:

Parameters	Mean AT	Mean BT	SD +/-	SE +/-	't' Value	'p-Value
RukshaTvak	3.71	2.07	0.50	0.133	12.33	< 0.001
MeduraMamsaPrarthana	3.438	3.142	0.462	0.124	2.3	< 0.05
Karshya	2.14	0.74	0.51	0.10	14.57	< 0.001
SandhiShunyata	3.438	3.142	0.462	0.124	2.3	< 0.05
Serum HDL Cholesterol	13.3 (Mean)		1.79	0.34	39.11	< 0.001

DISCUSSIONS

Table no.1 shows that the patients of the sample are satisfying the subjective criteria of *Pandu*. It reveals that the selected patients are suffering from *Pandu*. Cent percentage of the patients are satisfying the criteria of *'Hataprabha'* which implies the criteria of generalized pallori. the pale complexion of the skin. *Daurvalya* in cent percentage of people supports the evidence of *Oja Kshaya* and *Rakta Kshaya*. Feature of *Hatanala* in 96.6% represent *Agnimandya*. Other notable percentages are the identical feature of *Pandu* present in the sample together represents the evidence

of *Rasa, Rakta* and *Oja Kshaya.* Analysis of percentile data determines that all the patient included in the study are suffering from *Pandu.* Table no. 2 shows that the clinical criteria of *Meda Kshaya*, is present in the sample in the major patient, according to the percentile values. The literary information of *Meda Kshaya* in *Pandu* is verified by these data, and it reveals that sequences of *Meda Kshaya* take place in the disease process of *Pandu.* Regarding the point *Karshya*, it is to be said that no instrumental measurement has been taken, evaluation is done on the basis of macroscopic view of one observe only. The feature of *Pleehavi Vriddhi* i. e splenomegaly is absent in the sample, it may be due to exclusion of hepato-splenomegaly in this present study. Table no 3 and table no 4shows the improvement of subjective& objective criteria which denotes that Sarapunkha along with correcting the reduced level of Haemoglobin count combats Meda Kshaya internally which reduces the external manifestation of subjective criteria at a sustainable rate. In Ayurveda pharmacodynamics, Sarapunkha (Tephrosia purpurea Linn.) is found to be Kapha-Vata Hara (pacifies Kapha & Vata Dosha), Vrana Hara (heals ulcers), Jwara Hara (pacifies fever), Visaghna (anti-toxic), Krimighna (antihelminthic) and Rasayana (rejuvenating). Sarapunkha contains Tikta-Kasaya Rasa, Laghu-Ruksha-Tikshna Guna, Usna Virya, Katu Vipak, so it is advised as the good medicine in Pandu as well as in Meda Dusti, a hypothetical feature in the pathogenesis of Pandu. Hence the effectiveness of Sarapunkha is established in management of Pandu.

CONCLUSION

The study of the pathogenesis of Pandu reveals that the Pitta located in the heart is involved in course of the disease which vitiates the other fractions of *Pitta* through Dasavidha Dhamani, especially the pitta located in the skin. Agnimandya in Pandu causes altered Dhatupaka, which results in insufficient production of *Oja*, for this reason, the symptomatology of Indriya Shaithilya, Dourvalya, Sara Heenata etc. develop. All the patients in this study sample had satisfied the subjective criteria of *Pandu*, which denotes that every individual of the sample is suffering from Pandu irrespective of all doshas. Decreased Cholesterol level as a whole along with the mild low level of HDL Cholesterol in the sample shows that along with the decreased blood components the patients are also showing the evidence of low Lipid Components, may be due to nutritional origin, induced by the Aharaja Nidan. Declined level of HDL Cholesterol which may be interpreted as *Prakrita Meda*, rendering support to the dictum of 'Alpa Rakta Alpa Medaska.' Again, improvement of good Cholesterol levels implies that Sarapunkha (Tephrosia purpurea) can increase the Meda. As there is no literary reference of Sarapunkha to correct altered Meda, so it is to be interpreted that quantitative and qualitative improvement of Rakta i.e., blood and blood components are the indirect cause to improve Prakrita Meda. Therefore, at the end of the study it is to be concluded that in Pandu the pathological phenomenon of occurrence of Alpa Rakta and Alpa Meda is present in the study sample which had been verified biochemically and also Sarapunkha is an excellent remedy to treat Pandu and Meda Kshaya.

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