

WORLD JOURNAL OF PHARMACEUTICAL RESEARCH

SJIF Impact Factor 8.084

Volume 9, Issue 8, 1600-1608.

Case Study

ISSN 2277-7105

PARINAMSHULA (DUODENAL ULCER) AYURVEDIC MANAGEMENT- A CASE STUDY

Madhu Rani*¹, M. B. Nikumbh², Sumitra Mane-Deshmukh³ and Snehal M. Yadav⁴

^{1,3,4}P.G. Scholar, Department of Rachana Sharir, Govt. Ayurvedic College, Osmanabad, 413501, Maharashtra, India.

²Professor & H.O.D., Department of Rachana Sharir, Govt. Ayurvedic College, Osmanabad, 413501, Maharashtra, India.

Article Received on 05 June 2020,

Revised on 25 June 2020, Accepted on 15 July 2020

DOI: 10.20959/wjpr20208-18157

*Corresponding Author Dr. Madhu Rani

P.G. Scholar, Department of Rachana Sharir, Govt. Ayurvedic College, Osmanabad, 413501, Maharashtra, India.

ABSTRACT

Duodenal ulcer a type of peptic ulcer is very common health issue in all countries of world especially developing countries. There are a lot of causes of it including lifestyle. It is very common in people who are always in hurry, eat spicy curry and do worry. A male patient of 30 years came with complain of excessive retrosternal burning, sour eructation's, headache, epigastric pain and indigestion. These symptoms were felt after 1.5-2 hours of meals. According to Ayurveda diagnosis was made as 'Parinamshula' in which the pain or symptoms appear during the time of digestion or transformation of food. In modern medical sciences it can be correlated with duodenal ulcer and the treatment for it is only symptomatic by painkillers and

antacids. The chronic consumption of these medicines causes hepatotoxicity and disturbance to intestinal flora. Ayurvedic *panchakarma* treatment like *virechana*, *vamana* followed by *shamana* have miraculous result in these types of cases. A try is done to present this case by ayurvedic *panchakarma* treatment.

KEYWORDS: Duodenal ulcer, *parinamshula*, panchakarma treatment.

Aim and objectives

- To understand the *samprapti* (pathogenesis) of *parinamshula*.
- To assess the role of *virechana* & *vamana* (purification of body) in treatment of *parinamshula*.
- To assess the role of *dincharya* in treatment of lifestyle disorders like *parinamshula*.

INTRODUCTION

Duodenal ulcers are a type of peptic ulcers. There are more than 1 million cases per year in India. Male to female ratio of for duodenal ulcers varies from 5:1 to 2:1. [2] So, the ulcers are more common in males. Duodenal ulcers occur in the upper part of small intestine i.e. in duodenum. In duodenal ulcer pain or gastric discomfort occurs after 1-2 hours of eating. The symptoms are like pain in abdomen, sour eructation, bloating, nausea, vomiting, headache etc. The main cause of the ulcer is H. Pylori bacterial infection. The faulty life style, eating habits, stress, medicines like NSAIDs also have a major role in causation of peptic ulcers.

In Ayurveda it can be corelated with the disease entity *parinamshula*. *Parinamshula* means the *shula* or pain occur in the *parinaman kaal* (digestion or transformation) of food. [1]

During this stage of digestion of food *pitta* is dominant that's why the symptoms are more severe during this stage. In this disease the vitiated *vata* gets mixed with morbid *pitta* and *kapha* and later gets consequently blocked by these 2 *doshas* causing severe pain.^[3]

The main clinical features described in Ayurveda are pain abdomen during digestion of food, epigastric pain, pain in flank region, para-umbilical pain, pain in sternal area and over bladder area. The pain occurs during digestion process. *Amlapitta* symptoms like sour eructation, burning and pain in chest & epigastric region may be present along with it.

According to dominancy of specific *dosha* the symptoms may be indigestion, bloating, constipation, anxiety etc.(*vata*),^[5] thirst, burning, excessive sweating etc.(*pitta*),^[6] vomiting, nausea, drowsiness etc.(*kapha*)^[7] Or the symptoms may be mixed if two or the all three *dosha* are present in the pathogenesis of the disease.^[8]

As this is a lifestyle disorder, so firstly we have to correct *dincharya* (daily schedule), eating habits of the patient. It helps in the correction of biological clock.

In Ayurveda *panchakarma* treatment is like main axis of chariot. In *panchakarma* the treatment is done by detoxification of body by performing different purification techniques like *vamana* (emesis), *virechana* (purgation), *basti* (medicated enema), *raktamoksha* (bloodletting) and *nasya* (nasal medication).

In pathogenesis^[9] of the *parinamshula* there is *aavarana* of vitiated *pitta* and displaced *kapha* by vitiated *vata*. This vitiated *vata* causes pain and the *vikrut pitta* causes *amlapitta* like

symptoms. As this *pitta* is increased in quantity but doesn't have *pachana* or digestion property. There is increment in *drava guna* of the *pitta*. This causes *agnimandya* which leads to indigestion of food, nausea, vomiting. The *doshas* are so much aggravated and ready to come out of body. *Virechana* is the best treatment described in Ayurveda for the *pitta* dominant ailments. According to dominancy of *dosha* and *ritu vamana* also shows wonderful results. So, *vamana*, *virechana*^[10] along with lifestyle management was taken under consideration in this case.

MATERIAL AND METHODS

Case presentation

A male patient of 30 years came with complain of excessive epigastric and retrosternal burning, epigastric pain, headache after 1.5-2 hours of intake of meals along with bloating, indigestion and constipation.

Patient used to feel cough cold symptoms along with all these symptoms.

Patient was suffering from above complains for 2 years.

Past history

No history of any medical disease like hypertension, diabetes, thyroid, COPD etc.

No history of any surgical intervention.

Drug history: History of intake of anta-acids and antihistamines.

Family history: No relevant history.

Dietary history: Mixed diet, daily consumption of spicy, oily and non-vegetarian food.

Occupation: Primary school teacher.

Examination of patient

Examination	Findings
	No scar mark or other abnormality seen
P/A	Soft, mild tenderness over epigastric, left and right hypochondriac regions.
	Bowel sounds normally audible.
CVS	S1, S2 normal audible, no murmurs
	Chest clear bilateral
R/S	No added sound
	Air entry equal and adequate bilateral.
	Conscious and Well oriented to time, place and person.
CNS	E4V5M6
	Pupils normal in size and reactive to light bilaterally.

Madhu et al.

Ashtavidha pariksha

Nadi: 84/min, regular, VP

Mala: Unsatisfactory, once a day

Mutra: Samyak pravriti, 7-8 times a day

Jihva: Ishat-sama
Druka: Shweta

Sparsha: Anushan-sheeta

Aakriti: Madhyam Shabda: Prakruta

Investigations

ECG: Within normal limits

CBC: Hb: 14.1 gm/dl **TLC:** 8.7*10³/ul **PLT:** 293* 10³/ul **RBC:** 5.85 *10⁶/ul **HCT:** 45.1% **MCV:** 77.1 Fl

MCH: 24.1 l pg **MCHC:** 31.3L gm/dl

Coagulation profile

Bleeding time: 1 minute 45 sec. **Clotting time:** 5 minutes 5 sec.

Rft

Serum creatinine: 0.79 Blood urea: 26.4

Blood sugar: Fasting: 100 mg%

Lipid profile

Total cholesterol: 288.4 mg/dl Serum triglyceride: 164.4 mg/dl HDL Cholesterol: 40.0 mg/dl LDL Cholesterol: 215.5 mg/dl

VLDL Cholesterol: 32.88 mg/dl T.C./HDL ratio: 7.2

0

LDL/ HDL ratio: 5.3

Treatment

Virechana was planned. First of all, deepana- pachana was given with Avipattikar churana 3gm mixed with goghrita with first bite of meal -twice a day and Goghruta 2 tsf empty stomach in morning and evening for 5 days.

For *virechana*, *Snehapan* was given with *murchit til taila* (upto *samyak snigdha lakshana*). *Til taila* was used in view of elevated cholesterol levels. After that *purvakarma- snehan*, *swedana* was given. *Abhyadi modaka-*2 tablet -stat followed by 2tab after 2 hrs and *Aargvadhadi kadha-* 500ml divided in 2 doses, *Munakka phant-* 200ml divided in 2 doses was given on day of *virechana*.

Total *Vega*- 30 (*pravara shudhi*), so *samsarjana karma* was planned for 7 days with *peya/lahya -vilepi- akrita yusha/mamsa rasa-krita mamsa rasa/krita yusha*.

Sr.no	Procedure	Medicine	Dose	Frequency	Duration
1.	Deepana- pachana	Avipattikar churna	3gm with goghrita	B.D.	1 st -5 th day
2.	Snehapana	Murchhit til taila	Varedhman quantity upto samyak snigddh	O.D.	6 th -10 th day
3.	Sarvang- Snehana- swedana	Snehana- til taila, swedana- dashmool kwatha	Upto samyak sweda	O.D.	11 th -14 th day
4.	Virechana	Abhyadi modaka, munakka phant, aargvaddhadi kadha,	A.M 4 tab, Phant- 200ml, A.K 500ml	In divided doses	15 th day
5.	Samsarjana karma	Peya-vilepi-mamsa rasa			For 7 days

The patient was asked to visit OPD after completion of *samsarjana karma* or any other emergency. He was advised to do lipid profile.

Effect

- No constipation and bloating
- Lightness in body
- Headache and cold relieved
- Sour eructation relieved
- No abdominal pain
- Decreased levels of serum cholesterol
- Decrease in weight.

Shaman chikitsa was given on follow up

- Yashtimadhu ghruta 10ml before meals.
- Swarnasutshekar 125mg twice a day.
- Aarogyavardhini vati 500mg bd



1604

Again, he felt sour eructation and retrosternal burning after somedays. So, *vamana* was planned according to state of *dosha* and *kala* (*vasant ritu*).

All routine examinations along with lipid profile and ECG were done.

Snehapana: with yashtighrita was done upto the samyak snigdha lakshana. (6 days) followed by rest on 7th day, on 8th day – snehana- swedana and kaphavardhak aahara (rice+milk, banana, halwa etc.) in dinner and on 9th day- vamana was given. On vamana day Snehana, swedana followed by dugdha pana- vamak dravya chatan (madanphal pimpli+vacha+ honey) was given. Waited for 1 muhurata. Then, vega started in 30 minutes. Yashtimadhu phant – according to patients' capacity (about 3 litre) was given.

About – 12 *vega* (*pravara shudhi*) of *vamana* and *Apitdarshnata* with lightness in body.

Paschat karma was given with dhumpana (haldi+karpur). Patient was kept under observation for 1 day. There was not any fresh complain. On discharhge samsarjana karma for 7 days was advised (peya- vilepi-akrita mamsa rasa/yusha - krita mamsa rasa/yusha). After that Rasayana karma- vardhman pipli rasayan (14 days). Patient was asked for Follow-up after this.

On next visit no fresh complain, all symptoms were relieved. Patient was feeling healthy and energetic. Dyslipidemia was arranged. There was also decrease in weight.

DISCUSSION

On consumption or getting exposed to the causative factors which causes vitiation of *vata* out of proportions. This vitiated *vayu* gets mixed or associated with morbid *pitta* and *pitta* and subsequently get blocked by these two *doshas* causing severe colic called *parinamshula*.^[4] It can be named according to the dominancy of *dosha* like *vataj*, *pittaj*, *kaphaj* etc.

According to alternative pathogenesis^[5] of the *parinamshula* there is *aavarana* of vitiated *pitta* and displaced *kapha* by vitiated *vata*. This vitiated *vata* causes pain and the *vikrut pitta* causes amlapitta like symptoms. As this *pitta* is increased in quantity or *dravta* but doesn't have *pachana* property. This causes *agnimandya* which leads to indigestion of food or *amotpatti*.

This increased and vitiated *pitta* goes from its normal place to other places causing sour eructation, belching, burning chest, epigastric burning etc. The *ama* produced causes blockage of channels leading to headache, cold, bloating and constipation along with nausea, vomiting etc.

So, for *amotpatti* first of all *deepana pachana* was prescribed to increase the digestive fire. After that *virechana* was planned as it is the best treatment described in Ayurveda for the pitta dominant ailments. According to dominancy of *dosha* and *ritu vamana* also shows wonderful results. Here *murcchit til taila* was used instead of *ghritapana* in view of dyslipidemia.

All this treatment showed marvelous result in patient.

CONCLUSION

As *parinamshula* related to lifestyle, increased stress so first of all patient's lifestyle and dietary habits must be changed. *Virechana* and *Vamana* are *panchakarma* treatment which cleanses the body so, removes all the vitiated *dosha*. For the remaining *dosha* palliative treatment can be given. For more specific results with statistical data the study should be done on large sample size. Cholesterol decreased as can be seen by reports: -

iame Of Patient : SYED AHEMAD		Date 24-12-019			Date:30-12-0
IGE: yrs SEX	: Male		Name Of Patient : SYED AHME	ED	
Ref.By : DR : SELF			AGE: yrs	SEX: Male	
		••••••	Ref.By : DR :SELF		
BIO-CHEMISTRY	" LIPID PROFILE "	•	BIO-CHEMISTRY	*** LIPID PROFILE ***	
TEST	OBS VALUE	: NORMAL VALUE	TEST	OBS,VALUE	: NORMAL VALUE
TOTAL CHOLESTEROL	248.4 mg/dl	<200 mg/dl	TOTAL CHOLESTEROL	228.4 mg/dl	<200 mg/dl
SR.TRIGLYCERIDE	153.3 mg/dl	<150 mg/dl	SR.TRIGLYCERIDE	137.4 mg/dl	<150 mg/dI
HDL CHOLESTEROL	42.0 mg/dl	>45 mg/dl	HDL CHOLESTEROL	40.0 mg/dl	>45 mg/dl
LDL CHOLESTEROL	175.7 mg/dl	<130 mg/dl	LDL CHOLESTEROL	160.9 mg/dl	<130 mg/dl
VLDL CHOLESTEROL	30.66 mg/dl	0-30 mg/dl	VLDL CHOLESTEROL	27.48 mg/dl	0-30 mg/dl
T.C./HDL CHO.RATIO	5.9 Ratio	3.0-5.0 Ratio	T.C./HDL CHO.RATIO	5.7 Ratio	3.0-5.0 Ratio
LDL/HDL CHO.RATIO	4.1 Ratio	1.0-3.5 Ratio	LDL/HDL CHO.RATIO	4.0 Ratio	1.0-3.5 Ratio

		Date:6 -1-20					
			CLIENT NAME : NHM		REG. LARNAUM : 280		: 28/01/2020 04:1
				BIOCHE			: OSMANARAD
AGE: 30 yrs	SEX: Male		Investigation	Result	Units	Bio. Ref. Inter	al
Ref.By : DR SACHIN TIKE [G.A.C	c.os,Bad]		Serum Total Cholesterol	: 165	mg/dl	Desirable Borderline High High	< 200
BIO-CHEMISTRY	*** LIPID PROFILE *	**	Method CHOD-APP Reference Ranges as per NCEP ATP III Guide Aiert III 8-10 hours fasting is mandatory for lipic	lines. parameters. If not,	values might fluct	vate.	
TEST	OBS.VALUE	: NORMAL VALUE	Serum Triglycerides	: 214.1	mg/dl	Borderline High:	: < 150 150 - 199 : 200 - 499
TOTAL CHOLESTEROL	288.4 mg/dl	<200 mg/dI	Method. Enzymatic GPO Trinder.				>= 500
SR.TRIGLYCERIDE	164.4 mg/dI	<150 mg/dI	*References Range as per NCEP ATP IIf Guide *Alert III 8-10 hours fasting is mandatory for lipid		values might fluctu	rata.	
HDL CHOLESTEROL	40.0 mg/dl	>45 mg/dI	Serum VLDL-Cholesterol Method Calculated	42.82	mg/dl	10 - 35	
LDL CHOLESTEROL	215.5 mg/dl	<130 mg/dl	*References Range as per NCEP ATP III Guidel *Alert III 8-10 hours fasting is mandatory for lipid	nes. parameters, If not, v	ralues might fluctu	ate.	
VLDL CHOLESTEROL	32.88 mg/dl	0-30 mg/dI	Serum HDL-Cholesterol	: 42.4	mg/dl	Low <40	
T.C./HDL CHO.RATIO	7.2 Ratio	3.0-5.0 Ratio	• Method Direct			High >60	
LDL/HDL CHO.RATIO	5.3 Ratio	1.0-3.5 Ratio	"References Range as per NCEP ATP III Guideling "Alert III 8-10 hours fasting is mandatory for lipid p		alues might fluctu	ale.	

REFERENCES

- 1. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 472(1): 26-16.
- 2. Davidson Santley, Davidson's principles & practice of medicine, alimentary tract & pancreatic disease, 2008; 885: 20-22.
- 3. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 472(1): 26-15.
- 4. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 472(1): 26-16.
- 5. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 473(1): 26-17.
- 6. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 473(1): 26-18.
- 7. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 473(1): 26-19.
- 8. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 473(1): 26-20.
- 9. Sastri Sri Sudarsana, Madhavanidanam of Shri Madhavkara, Chaukhamba Sanskrit sansthan publication, Varanasi, chapter shuladinidanam, 472(1): 26-15-16.
- 10. Shastri vidya laxmipati, Yogaratnakara, Chaukhambi Sanskrit sansthan, Varanasi, volume 2, *uttrardha*, *shulnidana*.

Articles referred:

11. Parinamshula, definition, types, treatment, medicines by Dr Raghuram Y.S.

12. A holistic approach to management of peptic ulcer by Dr. Ram Kumar Agarwal et al., wjpls, 2(6): 428-434.