



Case Study

MANAGEMENT OF RIGIDITY DOMINANT PARKINSON'S DISEASE THROUGH AYURVEDIC PROTOCOL

Jesny V Jose^{1*}, Shylamma T M², Aparna V K³

*¹Post Graduate Scholar, ²Professor and Head of the Department, ³Assistant Professor, Department of Kayachikitsa, Government Ayurveda College, Tripunithura, Kerala, India.

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ABSTRACT

Parkinson's Disease (PD) is a disease with insidious onset and slow progression. It is a neurologic condition that causes motor manifestations namely, bradykinesia, rigidity, resting tremor, postural instability and non-motor symptoms such as depression and dementia. The modern treatment provides some symptomatic relief but any proven means for slowing the progression have not been found yet. This case is of a 64-year-old male patient who presented with complaints of slowness in daily activities, difficulty in raising left upper arm along with heaviness, tremors in both hands, difficulty in speech and movement for 3 years. Complaints were increasing progressively hindering his routine activities. He underwent our IP management for 60 days with follow-up after every 20 days. In every follow-up, the patient reported significant relief in his symptoms and after two months, he was able to perform his routine activities without any help.

INTRODUCTION

Parkinson's disease is the second most common neurodegenerative disorder characterized by loss of dopaminergic neurons in the substantia nigra pars compacta and locus coeruleus[1]. It is included in ICD10. G20-G26 extra pyramidal movement disorders[2]. It is manifested by a combination of rigidity, bradykinesia, postural instability with or without resting tremor. Globally Parkinson's disease affects approximately 1% of those more than 55 years and the incidence doubles by the age of 65 years[3]. Direct reference of parkinsonian movement disorders in ancient Ayurvedic literature is sparse and refers only related symptoms such as Kampa, Sthambha, Gatisanga, Chestasanga etc. For treating numerous neurodegenerative disorders, nowadays clinicians depend upon the treatment principles of diseases mentioned under Vatavyadhi prakarana of classical texts. There is no precise treatment principle available parkinsonian movement disorders in any Ayurvedic literature. Bhasavaraja the author of Bhasavarajeeyam coined the term Kampavata with



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features like *Karapadathala kampa*, *Deha bhramana*, *Matiksheena* and *Nidrabhanga*^[4]. It is better to take such description as a condition seen among Parkinson's disease patients than to compare *Kampavata* with Parkinson's disease.

Patient Information

A 64-year-old moderately built male patient retired as deputy superintendent (desk job), presented with complaints of slowness in daily activities, difficulty in raising left upper arm along with heaviness for the past three years. Tremors developed gradually that were aggravating at rest and subsiding during activities. He consulted a nearby modern hospital and was diagnosed with Parkinson's disease. They advised medicines for the same, which he is continuing even now. Gradually heaviness over upper limb and walking difficulty also developed. Recently he started experiencing night mares and increased anxiety which disturbs his night sleep. For the past one year, the patient noticed a tendency to fall forward while walking, difficulty in speech particularly during initiating sentences and expressing emotions, difficulty in raising left arm and holding objects etc.

His appetite and thirst were normal but he complained of hard stools with incomplete evacuation once in two days for past four years. He was also diagnosed with co-morbidities like diabetes mellitus and hypertension. He started to notice difficulty in

initiating day-to-day activities such as buttoning and unbuttoning, slowness in eating food, difficulty in holding objects and recollecting recent fact for the past five months. For this he consulted a neurologist and was prescribed medicines Tab. Gabapentin 100mg (0-0-1), Tab. Syndopa plus (1-0-1), Tab. Rasalect 0.5mg (1-0-1) Tab. Cilacar 10mg (1-0-1), Tab. Melmet 500 SR (1-0-1), Tab. Glizid MR- 60 (1-0-1). No one in his family had similar complaints and he had no history of exposure to chemical toxins, poison, head injury etc. He took all the prescribed modern medication but no significant relief was seen. Gradually there was an increase in symptoms which compelled him to visit our hospital.

Clinical Findings

A moderately built, well-dressed hygienic, cooperative male patient having erect posture and masked - anxious facies with 21.77kg/m² body mass index. His gait was festinating. His blood pressure was 150/90mmhg and pulse regular with rate 74/min. The respiratory system, gastro-intestinal system, cardiovascular system, remote memory and other higher mental functions were within normal limits. On examination, his olfactory nerve was affected, distant vision was impaired and glabellar tap was positive. No relevant muscle bulk reduction noticed. Muscle power found to be within G4 and G5 grades. Speech was monotonous. Deep tendon reflex over triceps and biceps were diminished for left upper limb, all other reflexes were found to be normal.

On examining the cerebellar functionsdysdiadochokinesis was absent but finger-nose test and heel shin test were impaired on left side.

Romberg's test (eye closed) and tandem walking were not possible. Finger tapping was also impaired on left side. His saccades and pursuits were slower than normal limits. On examining the involuntary system, the tremor was slow-repose with rate 22-24/min, rhythmic, multifocal and non-specific direction in nature. On further examination, progressive micrographia, cog-wheel rigidity and hypokinesia were present.

Ashtasthana Pareeksha

Nadi (pulse) - Sarpagati
Mutra (urine)- Atisrishta
Mala (faeces)- Badham (constipated)
Jihwa (tongue) - Upalipta (coated)
Sabda (voice) - Manda and Asphashta
Sparsa (tactile examination) - Anushna seetha
Drik (eyesight and eye) -Hraswadrishti
Akriti (body) - Madhyama

Diagnostic Assessment

Diagnosis of Parkinson's disease is based on clinical presentation. As the patient had fulfilled the cardinal features of rigidity dominant Parkinson's disease along with red flag sign (frequent nightmares) and with no specific causative factors, it can be diagnosed as rigidity dominant idiopathic Parkinson's disease. The cardinal features like cogwheel rigidity, pill rolling tremor (mild), bradykinesia, postural instability was present in this case. For the evaluation of the effect of treatment, grading of subjective parameters has been adopted as mentioned in Table 1.

Tabl	e 1:	Grading	of A	Assessm	ıent	Criteria
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Grade	Tremor
0	No tremor
1	Unilateral slight tremor present at rest decreased by action
2	Bilateral tremor
3	Tremor is not violent but involves few body organs
4	Bilateral violent tremor not suppressed or diminished by the desired movement
Grade	Rigidity
0	No rigidity
1	Rigidity is present but vanishes on continuous examination
2	Moderate rigidity was demonstrable and remained throughout the examination
3	Marked rigidity and full range of motion achieved easily
4	Severe rigidity and full ROM achieved with difficulty
Grade	Akinesia/Bradykinesia
0	Can walk without aid
1	Can walk without assistance slowly but with a shuffling gait
2	Can walk with assistance slowly

3	Can walk slowly but need substantial help shuffling with retropulsion/ propulsion lack of associated movements
4	Unable to walk without assistance
Grade	Postural Instability (p)
0	Normal
1	Slightly stooped, not quite erect
2	Moderately stooped
3	Severely stooped with kyphosis can be moderately leaning to one side
4	Marked flexion with extreme abnormality, unable to arise from the chair without help

^{*}The scale is not validated and is being used in the institute to evaluate the efficacy of the formulation in Parkinson's disease.

*Treatment protocol adopted for this patient is given below

Stage 1- Pachana and Anulomana (internal), Rukshana kriya (external)

Stage 2- Samana kashaya, Sodhanapurva achasnehapana (internal)

Stage 3- Sodhana- Vamana

Stage 4- Mrudu brumhana oushadha (internal), Pinda sweda and Prathimarsa nasya (external)

Stage 5 - Sodhana- Virechana

Stage 6- *Brumhana oushadha* (internal), *Pizinju thadaval* (external)

Stage 7- Sodhana- Virechana

Stage 8- Niruhavasti

Stage 9- Brumhana oushadha (internal), Murdhnitaila,

Marsa nasya (external)

Stage 10- Rasayana

Table 2: Internal Medications

Chaga	Chara Calastian of duna Describing					
Stage	Selection of drug	Dose of drug	Duration			
1	Gandarvahastadi kashaya Shaddharana churna Vaiswanara churna	90ml bd, 6 am & 7 pm with Saindava (3g) and Gudam (12g) 3g with Kashaya 12gm with Takra before meals	Initial 3 days			
	Ashtavargam kashaya	90ml bd, 8 am & 5 pm	Next 4 days			
	Vata gajankusha rasa Parasika yavani choorna	100-200mg bd with Guggulutiktaka ghrita (sufficient quantity for properly mix the powder of medicine) 3g with Dadima swarasa/warm water	Next 4 days			
2	Astavargam kashaya Nayopayam Kashaya	90 ml bd 90 ml bd + <i>Dhanwantaram gulika</i> 1-0-1 as <i>Anupana</i>	7 days			
	Sodhanapurva achasnehapana with Sahacharadi taila	Fixed <i>Hrasiyasi matra</i> according to <i>Jaranasakti</i> of the patient and gave <i>Taila</i> till <i>Samyak snigda lakshana</i> seen	7 days			
4	Sahacharadi kashaya Maharasnadi kashaya	90 ml bd given as <i>Panam</i>	5 days			
6	Badradarvadi kashaya Maharasnadi kashaya	90 ml bd 6 am, 6 pm 90 ml bd 11 am, 8 pm	7 days			
9	Vidaryadi kashaya Maharasnadi kashaya Aswagandharishta	90 ml bd + Sahacharadi taila 21(A) 12ml with Kashaya Given as Panam 25ml bd	7 days			
10	Chitraka rasayana	3g with Sahacharadi taila (12 ml) as Anupana	1 month			

Table 3: Procedural Interventions

Stage	Type of Opted Therapy	Drug of Choice	Duration
1	Udvartana	Kolakulathadi churna	First 5 days
	Dhanyamladhara	Dhanyamla	Next 7 days
	Churnapinda sweda (Ruksha)	Kolakulathadi churna + Triphala churna (since patient is diabetic)	Next 7 days
	Utgharshana	Kolakulathadi churna + Kulatha churna (3:1) ratio with medium depending upon the Prakriti and skin type, so used – Dhanyamla	Next 7 days
2	Abhyanga + Ooshma sweda	Chinchadi taila	3 days
3	Vamana	Madanaphaladi yoga	1 day
4	Jambeera pinda sweda	Parinithakeri ksheera taila for Abhyanga	7 days
4	Pratimarsa nasya	Anutaila – 2 Bindu each nostril	7 days
5	Virechana Sukumaraeranda taila dose-30-40ml Anupana-milk time- 8 am		1 day
6	Pizhinju thadaval	Pizhinju thadaval Masha saindava taila	
7	Virechana Sukumaraerandataila dose-30-40ml Anupana-milk time- 8 am		1 day
8	Musthadi rajayapana vasthi Sneha dravya- Ksheerabala taila (Chikkana paka-ml) Saindava - 15g Madhu – 100ml Ghrita- 50ml F 30g Ksheerakashaya- 240ml Mamsarasa - 100ml		8 days (as <i>Yogavasthi</i> pattern)
9	Sirodhara	Maha <mark>n</mark> arayan <mark>a t</mark> aila	7 days
	Dhmana nasya	Nasik <mark>a c</mark> hurna (3 Much <mark>u</mark> di)	3 days
	Brihmana nasya	Maharaja prasaranyadi taila (1ml in each nostril)	4 days

RESULT AND OUTCOME

Patient showed marked improvement after each stage of treatment. The first positive response (rigidity of upper limb reduced to 50%) was obtained after *Vamana*. The pain and heaviness of upper limb reduced after initial stages of *Rukshana* therapy (especially after *Udghrshana* therapy). After *Jambeera pindasweda*, tremor also reduced. The *Koshta* and *Agni* level got improved after each stage of *Virechana*. Bradykinesia got reduced and the quality of life improved after *Brumhana* therapies done to patient. Postural instability improved after *Rasayana* therapy.

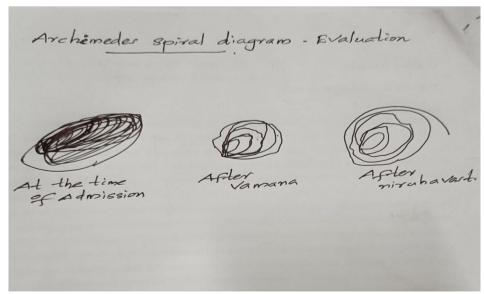


Diagram 1: Showing Archimedes spiral diagram evaluation after Vamana and Niruha vasti done to patient

Table 4: Effect of Treatment on Tremor, Rigidity, Bradykinesia & Postural instability

Location	BT	AT	Relief
Tremor on RUL	2	1	50%
Tremor on LUL	3	1	66%
Tremor on RLL	3	0	100%
Tremor on LLL	3	1	66%
Rigidity on RUL	3	0	100%
Rigidity on LUL	4	1	75%
Rigidity on RLL	3	0	100%
Rigidity on LLL	4	1	75%
Bradykinesia	2	1	50%
Postural instability	2	1	50%

(BT- Before treatment, AT-After treatment)

*RUL- right upper limb, LUL- left upper limb, RLL- right lower limb, LLL- left lower limb

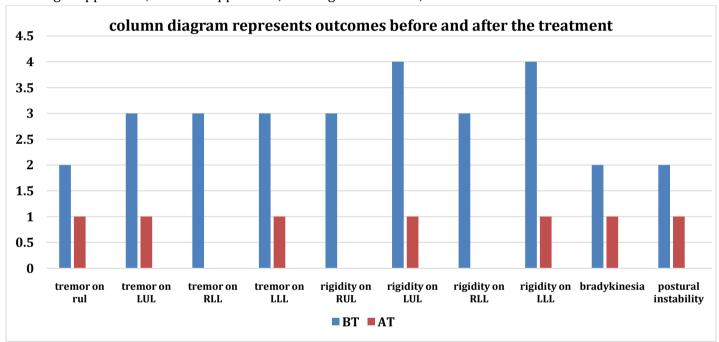


Figure 1: Column diagram represents outcomes before and after treatment

Follow Up

Marked reductions in the symptoms were seen after the treatment. 2 weeks after discharge patient came to OPD for follow up and there was significant improvement in rigidity, tremor, heaviness and difficulty in raising upper limb etc. Sleep pattern also got corrected. Medicines advised were *Vidaryadi ksheerapaka*, *Brahmi drakshadi kashaya* (45ml od) and *Vata gajankusa rasa* (1-0-1).

DISCUSSION

Wide range of Ayurvedic principles can be judiciously administered to different stages of Parkinson's disease according to *Yukti*. Considering the pathologic presentations, *Sthambha*, *Kampa*, *Gatisanga*, *Chestasanga* are the important features of all kind of

Unilateral/bilateral Parkinson's disease. involuntary movements of other body parts, postural instability, bradykinesia, monotonous speech, weight loss, depressed facies, constipated bowels with black stools, abdominal discomfort, flatulence, shoulder pain and heaviness, sleeplessness, drvness of skin. giddiness, mild memory impairment, difficulty to write using pen, olfactory dysfunctions, reduced appetite, indigestion and nightmares etc presented by the patient can be considered as Vata samsrushta lakshanas. Rigidity dominant Parkinson's disease can be considered as a Kapha predominant condition. Hence Ushna-ruksha and Langhana line management will give *Upasaya* for *Sthambha pradhana*

^{*}BT- before treatment, AT- after treatment

^{*}RUL- right upper limb, LUL- left upper limb, RLL- right lower limb, LLL- left lower limb

Parkinson's disease. Sheeta guna is always Anupasaya since it is aggravating Vata kapha doshas. Initially we should consider Agnibala and state of Apana vata. So, the first step in treatment is Vatanulomana, Amapachana and Agni deepana. For that we gave Gandarvahastadi kashaya with Yadarha prakshepa dravyas. Gandarvahastadi kashaya is a formulation having Tikta katu kashaya rasa pradhana and slightly Ushna veerya swabhava^[5]. It is having Apana vata anulomana, Srishtavinmutratva, Deepana and Ruchya properties. It is best to impart Apana vatanulomana and Agni deepana in initial stages. It is given in Apana vayu oushadha kala by conventional practice.

Bradykinesia even affects the peristaltic movement of intestine and patients present with chronic constipation as *Poorvarupa* and *Rupa*. Administration of *Gandarvahastadi kashaya* at *Apanavata oushadhakala* with appropriate *Churna* or *Prakshepa dravyas* will address the complaint more effectively. *Shaddharanam churna* explained in *Vatavyadhi prakarana* is having *Katu tikta rasa pradhana*, *Ushna veerya* and *Kaphaharatva* property^[6]. So, it is best to impart *Amapachanam* in initial phases. *Shaddharana churna* is told as *Mahavatavyadhi prasamana yoga* in *Cakradatta*. So, it can be safely administered to rigidity predominant Parkinson's disease.

After attaining proper Deepana, Pachana and Vatanulomanatva we gave Samana kashayas like Ashtavargam kashaya. Ashtavargam kashaya is commonly using for neurological conditions having Vatakapha origin. Major part of the drugs is Ushnaveerya and Vata kapha samana which in turn acts as Pachana as well as Avaranahara^[7]. Lasuna is one among the content and which exclusively act as Avaranaghna (except for Pitta and Rakta avarana). More over rigidity and bradykinesia can be consider as a Kaphavrita vata state. So, this yoga can be used as a broad spectrum Vatavyadhi drug having neurological origin. Parasika vavani churna is Katu tikta rasa pradhana, Guru ruksha guna, Ushna veerya and Madaka (narcotic) in general. It is having Vedanasthapaka, Soolaprasamana and Nidrajanaka properties[8]. Vatagajankusa rasa is prepared as Antardhuma method explained in *Bhaishajya Ratnavali*. Contents like Gandhaka, Kantha bhasma, Abraka bhasma are having Rasayana property.

Along with first stage of *Deepana pachana kashayas* internally, external *Kriyas* like *Udvartana*, *Utgharshana*, *Dhanyamla dhara* and *Churna pinda sweda* are done. *Udvartana* with *Kolakalathadi churna* is administered, to clear the association of vitiated *Kapha*. *Kolakulatha churna* is a compound drug with specific indication on *Vatavyadhi*[9]. Since the patient is presented with heaviness along with cog wheel rigidity, we did *Udgharshana* with *Kolakulathadi* or

Kulatha churna mixed with Dhanyamla. Dhanyamladhara is a procedure explained in the renowned treatise of traditional Kerala Ayurveda, Chikitsa Manjari which recommends Dhanyamla dhara as a first line therapy in Pakshaghatha^[10]. It is exclusively done for the Vatavyadhis associated with Ama as well as Pitta and Kapha. Churna pinda sweda is a type of Tapa sweda which is administered to clear the Avarana of Kapha. Kolakulathadi churna is the drug of choice opted for Churna pinda sweda.

The duration of *Bahya rukshana* can be modified to below 7 days if the patient attains the following changes namely improvement in appetite, proper evacuation of bowel, reduction in symptoms like heaviness, swelling, flaccidity, burning sensation, numbness and body pain. *Abyanga* with *Taila* having more *Kapha vata samana* property can be used from the second stage onwards. Hence, we selected *Chinchadi taila*, which is processed with *Amla lavana pradhana dravyas* and *Ushna, Teekshna* and *Kapha vatahara* in nature. *Ooshma sweda* also helps to bring back *Leena utklishta doshas* from *Uthamanga marma* to *Koshta* for elimination.

Sodhanapurva achasnehapanam is advised for imparting Dosa utklesa which is necessary prior to Sodhana procedures. For that purpose, we selected Taila preparations since the patient is having Vatakapha prakriti. The first phase of Sodhana should be Vamana because Vamana is found to be more effective in rigidity dominant Parkinson's disease. It is best to impart Uthamanga dosha nirharanatva, Kapha avaranagna and it is clinically experienced by expertise the improvement of rigidity (by drawing Archimedes spiral diagram before and after the procedure of Vamana) after first sitting of Vamana itself. After Vamana, Sodhana procedures like Virechana, Sodhana nasya and Sodhana vasti can be done. Nithyaanulomana is also recommended for this of patients because abnormal accumulation and aggregation is one of the strong causative factors for the pathology neurodegenerative disorder. So, the disease itself demands Nityaanulomana and Erandataila is the best drug of choice for imparting Mridu sodhana for longterm purpose. Eranda taila is a Snigda virechana oushada with Ushna teeksha sukshma and Sara qunatva. So, it is best to explicit its function as Srotosodhana and Amaharatva.

After the initial *Sodhana*, *Taila* and *Ghrita* can be used along with above said *Kashayas* as *Anupana dravyas*. As *Taila* and *Ghrita* are lipid medium, it can cross blood brain barrier more easily and can results in reducing the rate of progression of the disease. After first line of *Sneha sweda* therapy, in the second stage *Pinda sweda* is opted. *Pinda sweda* is a type of *Ooshmasweda* and is *Kapha vata samana* in nature.

Jambeera pinda sweda offers a better Kapha samanatva and Avaranaharatva due to its presence of Ushnaveerya and Kaphahara drugs. It is useful for curing stiffness or rigidity in limbs. Jambeera pinda sweda is Kaphavata samana and found to be more efficient in reducing the rigidity and bradykinesia. Samana kashayas are then replaced with Brimhana kashayas rather than giving Ruksha kashayas we can add Taila or Ghrita as Prakshepa along with Kashayas if the patient's condition is satisfactory.

In the next stage we continued *Brihmana kashayas* internally and *Snigda* procedures externally. For that purpose, we administered *Pizhinj thadaval*. Considering the involvement of *Upadhatus* like *Sira snayu kandara*, *Masha saindavadi taila* is a drug of choice for *Pizhinj thadaval*. Since the patient presenting more aching pain as clinical symptom *Mahanarayana taila*^[1] is selected for *Sirodhara*.

Being a Vatavyadhi, Vasti has got the prime role. The main aim of *Niruha vasti* is to impart *Dosha* samanatva. So that, Mustadi rajayapana vasti is selected. *Musthadi rajayapana vasthi*^[12] has an immense role in all three types of Parkinson's disease since it is a Yapana vasthi. Its Vrishva nature has the Prabhava to cure Dhatukshava especially in Majja (Masthulunga majja). dhatukshaya All Vrishva oushadhas has direct action in limbic system. Extra pyramidal structures have close relation with limbic system as they can increase the dopamine surge. Therefore, *Vrishyavasthis*^[13] are also commonly practicing in the management of Parkinson's disease. Nasya always has excellent results in neurological disorders since it is the easiest way to impart active principles to site of pathology than other *Panchakarma* modalities. Pratimarsa nasva is done throughout the treatment course after initial Kayasodhana since it is the procedure explained in our classics as "Aajanma maranam sastam".[14] At the later stage of the treatment, the patient's condition became Vatika avasta or free of Avarana state, so we did Marsa nasva with Taila after 3 days of Teekshna nasya (Dhmana nasya).

Rasayana chikitsa is very important in the treatment and prevention of Parkinson's disease. Since it is a rigidity dominant case, we selected *Chitraka rasayana* with *Sahacharadi taila* as *Anupana*. The total duration and number of days opted for both *Antaparimarjana* and *Bahiparimarjana chikitsa* is based on the *Upasaya* and *Anupasaya* of the *Chikitsa*.

CONCLUSION

Movement disorders are extremely common in clinical practice and account for a considerable proportion of neuro morbidity. Parkinson's disease is the second most common neurodegenerative disease after Alzheimer's disease. Direct reference of parkinsonian movement disorders in ancient

Ayurvedic literature is sparse and refers only to related symptoms such as *Kampa*, *Sthambha*, *Cheshtasanga*, *Gatisanga*. There so many varieties of *Chikitsa* explained in our classics and they can be logically applied for accurate condition in *Rogi* based on *Vaidya's yukti*. In clinical practices also we can see so many variations according to practitioners. Here the patient got considerable relief from his symptoms and his quality of life also got improved. Marked improvement in rigidity and heaviness of upper limb achieved. He was very much satisfied and trusted after the treatment. So, this successfully managed case and the treatment protocol will be a stepping stone to practitioners.

Declaration of Patient Consent

Authors certify that they have obtained patient consent form, where the patient has given his consent for reporting the case along with the images and other clinical information in the journal. The patient understands that his name and initials will not be published and due efforts will be made to conceal his identity, but anonymity cannot be guaranteed.

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*Address for correspondence Dr. Jesny V Jose

Post Graduate Scholar, Department of Kayachikitsa, Government Ayurveda College Tripunithura, Kerala, India.

Email: jesnyvjose@gmail.com

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