

# AYUHOM

A Peer Reviewed Bi-annual Research Journal of Ayurveda & Homoeopathy

Vol. 6 Issue 1 (Jan-June, 2019)



Published by

NORTH EASTERN INSTITUTE OF AYURVEDA & HOMOEOPATHY (NEIAH)
MAWDIANGDIANG, SHILLONG, MEGHALAYA -793018 (INDIA)

ISSN: 2349-2422

# AYUHOM

A Peer Reviewed Bi-annual Research Journal of Ayurveda & Homoeopathy Vol. 6, Issue 1 (Jan - June, 2019)



# Published By

# North Eastern Institute of Ayurveda & Homoeopathy (NEIAH)

Mawdiangdiang, Shillong, Meghalaya -793018 (INDIA)
(An autonomous institute under the Ministry of AYUSH, Government of India)
E-mail: ayuhom.neiah@gmail.com / neiahshillong@gmail.com / dir-neiah@nic.in
Telephone: +91-364-2538134; Website: www.neiah.nic.in

# **EDITORIAL BOARD**

## **CHIEF EDITOR**

Prof. (Dr.) P.K. Goswami, MD (Ay), Ph.D

Director

North Eastern Institute of Ayurveda & Homoeopathy (NEIAH)

Shillong, Meghalaya

## **EXECUTIVE EDITOR**

Dr. Bishnu Choudhury, MD (Ay), Ph.D

Lecturer (Ayurveda - Kayachikitsa)

North Eastern Institute of Ayurveda & Homoeopathy (NEIAH)

Shillong, Meghalaya

#### **ASSOCIATE EDITORS**

Dr. Abhishek Bhattacharjee, MD (Ay)

Lecturer (Ayurveda – Panchakarma)

NEIAH, Shillong

Dr. Himangshu Baruah, MD (Ay)

Lecturer (Ayurveda - Rasashastra)

NEIAH, Shillong

Dr. B.P. Chyne, MD (Hom)

Lecturer (Homoeopathy - Organon of Medicine)

NEIAH, Shillong

#### **ASSISTANT EDITORS**

Dr. Vijay Kumar, MD (Ay)

Associate Professor (Ayurveda - Swasthavritta & Yoga)

NEIAH, Shillong

Dr. Tapan Nath, MD (Hom)

Lecturer (Homoeopathy - Repertory)

NEIAH, Shillong

Dr. O.P. Patel, MD (Hom)

Lecturer (Homoeopathic Pharmacy)

NEIAH, Shillong

Dr. Monica Gupta, MD (Hom)

Lecturer (Homoeopathy)

NEIAH, Shillong

Dr. Himadri Bhaumik, MD (Hom)

Lecturer (Homoeopathy - Materia Medica)

NEIAH, Shillong

Dr Sikha Lekharu MD (Ay)

Lecturer (Ayurveda - Samhita & Sidhanta)

NEIAH, Shillong

## **ADVISORY MEMBERS**

Vaidya Rajesh Kotecha Secretary, Ministry of AYUSH

Govt. of India, New Delhi

Prof. (Dr) Dipika Deka

Vice Chancellor

Srimanta Sankaradeva University of Health Sciences, Guwahati

Prof. (Dr.) Abhimanyu Kumar

Vice Chancellor, Dr. Sarvepalli Radhakrishan Rajasthan

Ayurved University, Jodhpur

Prof (Dr.) K.S. Dhiman

Director General, CCRAS, New Delhi

Prof. (Dr.) P. Bhattacharya

Registrar (Academic) cum HOD, Anesthesiology

NEIGRIHMS, Shillong

Prof. (Dr.) A. C. Phukan

Dean (Academic) cum HOD, Microbiology,

NEIGRIHMS, Shillong

Prof (Dr.) R.K. Manchanda

Ex - Director CCRH, New Delhi

Dr. Manoj Nesari

Advisor (Ayu.), Ministry of AYUSH, Govt. of India, New Delhi

Dr. D.C. Katoch

Advisor (Ayu.), Ministry of AYUSH, Govt. of India, New Delhi

Prof. (Dr) Tanuja Nesari

Director, All India Institute of Ayurveda, New Delhi

Prof. (Dr.) Sanjeev Sarma

Director, National Institute of Ayurveda, Jaipur, Rajasthan

Prof. (Dr.) Anup B. Thakar

Director, IPGT & RA, Jamnagar

Prof. (Dr.) Subhash Singh

Director, National Institute of Homoeopathy

Salt Lake, Kolkata

Prof. (Dr) P.K. Prajapati

Head of Dept of Rasa Shatra, All India Institute of Ayurveda, New Delhi

Cum Managing Director, IMPCL Uttarakhand

Prof. (Dr.) B.K. Dwibedy

Head, Deptt. of Sidhanta Darsan, Faculty of Ayurveda

IMS, BHU, Varanasi, Uttar Pradesh

Prof. (Dr.) Anand Chaudhury

Head, Deptt. of Rasa Shastra,

Faculty of Ayurveda, IMS, BHU, Varanasi

Prof. S.R. Joshi

Deptt. of Biotechnology and Bioinformatics

North Eastern Hill University (NEHU), Shillong

# **CONTENTS**

Editorial	
Good Mentoring - Prof. (Dr) P.K. Goswami	1
Review Article	
Hyperactivity of Catecholamines - A possible pathophysiological background of	3
Homoeopathic pathogenesis of Aconitum napellus - Kisor Kumar Naskar, Ompriya Mishra	
Research Article	
Herbo-Mineral Combination Drug as Matravasti in Katigraha w.s.r to Inter Vertebral Disc	11
Prolapse - Divya Bendi, Perugu Srikanthbabu	
Case Reports	
Effectiveness of Ayurveda modalities in the management of Gridhrasi (Prolapsed Inter	19
Vertebral Disc)	
- Ritu Kumari, N.R.Singh, Jitendra Varsakiya	
Ayurvedic management of Sub-Acute Pyelonephritis - Alok Kumar	24
An Ayurveda Approach to Anidra w.s.r. to Anxiety induced Insomnia - Gopesh Mangal, Nidhi Gupta, Pravesh Srivastava	27
Verruca Vulgaris of scalp annihilated by Homoeopathic Medicine - Partha Pratim Pal	32
Homoeopathic approach to common viral skin diseases - Sangita Saha, Koushik Bhar, Mahakas Mandal	38



# NORTH EASTERN INSTITUTE OF AYURVEDA & HOMOEOAPTHY (NEIAH)

सत्यमेव जयते

(An autonomous institute under the Ministry of AYUSH, Government of India)
Mawdiangdiang, Shillong, Meghalaya-793018



Prof. (Dr.) P.K. Goswami Director

Ph.0364-2538134 Mob. 9415385128 Website www.neiah.nic.in

Email: neiahshillong@gmail.com pkgoswamibhu@gmail.com

ISSN: 2349-2422

# **EDITORIAL**

# **Good Mentoring**

The highly essential components for a good academic institution are quality teachers, quality students, up to date curriculum and necessary basic required infrastructure. Out of these four components, infrastructure though given visible identity of the institute, it is not necessarily basic pillar. It can be added, uplifted and can be used up to maximum utilisation as per availability. Curriculum and syllabus are implemented and governed by university and respective governing councils and it is observed that continuous process of revision, up gradation are undertaken by the concerned agencies time to time as per the policy of Govt. of India. So, these two are relatives and not directly under the control of college administration. On the other hand, for qualitative outcome of human resource from these institutes depends on quality intake of students and quality faculty members is highly essential. It is praiseworthy initiative has been taken by the Government of India by streamlining the admission policy to different stream of medical science through NEET. Though initially there was a minor difficulty, now majority of the institutes all over the country, student society, it is well accepted. Still there are some efforts seen in certain quarters of AYUSH system to smoothly accept NEET entrance policy. Need of hour is strict and uniform implementation of entrance policy through NEET. There should not be any other thoughts in this regard.

Next important matter is to select and recruit quality teachers (faculty members) in AYUSH colleges. It is the need of the hour to administratively streamlining teachers quality and their delivering of quality teaching activities, which does not effect and jeopardise the academic features, professional aptitude. Different agencies namely Government of India, different governing councils, state government, universities and management of private institutes should take very serious effort and strong determination to recruit quality teaching faculties to properly motivate young tender minds of students, admitted through NEET, in different AYUSH system of colleges. Unfortunately it is observed that majority of private institutes are not giving concern in this regard. It is encouraging that Ministry of AYUSH, GOI has taken serious views for streamlining quality teaching, for recruitment of efficient and quality teachers. The Ministry has notified advisory in this regard. But, recruitment of quality faculty in AYUSH College is a big challenge and complicated matter in totality. There is ambiguity, divergent problematic issues to proper recruitment of quality teachers. Mushrooming of large number of colleges in AYUSH systems and simultaneously opening of post graduate teachings and producing a large number of post graduate students in different discipline of different systems is not up to the mark. A very few notable institutes/colleges are countable in finger tips in terms of quality of their product with post graduate qualifications. This certification of qualification without quality forced. These teachers to impart sub-standard education to their students or indulge in undesired shameful practices from legal point of view and morality. These are core issues in AYUSH system for uniformly imparting quality UG and PG education and this is a very serious issue which push the whole system of AYUSH education in a critical juncture. Students are not trained properly in their own system forcing them to opt out for practice of other system of medicine (allopathy) for earning their livelihood and ultimately bringing bad reputation to their own system. It serves the vested interested groups, education mafias, pharmaceutical industry and also promotes quackery in legal sense.

So, it is highly essential that they recruit quality faculty in AYUSH colleges, only having PG degree in concerned subject to be adopted. Allied or inter-disciplinary appointment should be discouraged. Recruitment rules should be properly framed and to be monitored and modified from time to time as per guidelines of UGC. Even during determination of MSR, governing council rules should be in reference to basic criteria of higher education like UGC structure of pyramidal shape of each department such that minimum one Professor, two Associate Professors, four Assistant Professors should be composed for one department in any system. There may be some prospective positive signs but it may not be pleasing to all. Same thing is also applicable in cases of opening of new colleges and permission to open PG courses. So, after admitting quality students, it is to be expected smooth functioning, desired outcome, scientific rationale to accept by the society all systems of AYUSH in true spirit. It is matter of great need of introspection by all stake- holders of AYUSH system. Hopefully under the leadership of dynamic administration in AYUSH Ministry, we can see a ray of hope, that quality teachers will be available in colleges of AYUSH system and really capable teachers who can deliver to ignite the quest of knowledge in the mind of the young students, those who aspire quality education in AYUSH system to be better professional in future.

Young energetic and vibrant group of students need mentor to guide in order to excel in their own professional career within their own stream. Mentors who can help tender minds to understand holistic concepts of Ayurveda and create new knowledge with scientific temperament. Teachers not only to be preachers and mentors, they need to become good mentor as well as philosopher and guide to young minds and attract respect as ideal teachers.

Place: NEIAH, Shillong

Date: 30/12/2019

Prof. (Dr.) P.K. Goswami Director, NEIAH, Shillong Chief Editor - AYUHOM

#### **Review Article**

# Hyperactivity of Catecholamines - A possible pathophysiological background of Homoeopathic pathogenesis of Aconitum napellus

<sup>1</sup>Kisor Kumar Naskar, <sup>2</sup>Ompriya Mishra

<sup>1</sup>H.O.D., Department of Materia Medica, D. N. De Homoeopathic Medical Colleges & Hospital, Kolkata <sup>2</sup>H.O.D., Department of Obstetrics & Gynaecology, National Institute of Homoeopathy, Kolkata

Manuscript Received on 15/10/2019

Reviewed on 16/11/2019

Accepted on

21/11/2019

# **Abstract**

Background - Aconitum napellus (Aconite) is one of the most popular well proved Homoeopathic medicines. It is frequently prescribed at the beginning of acute conditions related to inflammation or functional abnormalities in different systems and organs, as in cardiovascular, respiratory, gastrointestinal, neurological, endocrinal disturbances associated with a state of anxiety and restlessness. These are often caused by fear or fright, or exposure to dry and cold weather. In proper application, Aconite can cut short the career of these conditions. Catecholamines are the secretary products of the adrenal medulla and sympathetic nervous system, which are important for defensive reaction to potentiate the organism's chances of survival by fight-or-flight response. Objective - In this study it has been tried to understand the possibility of influence of catecholamines which are directly or indirectly involved in the patho-physiological background of the Homoeopathic pathogeneses of Aconite. Method – Correlation between Homoeopathic pathogeneses of Aconite collected from the experiences of the pioneers with the actions of catecholamines related to these pathogeneses from different research works (in animal and human models) and their explanations. Result - Expression of psycho-neuro-endocrinal activities of catecholamines, especially noradrenaline have similar effects as that of essence of Homoeopathic pathogenesis of Aconite. Conclusion - Hyperactivity of catecholamines, especially noradrenalin (norepinephrine) might have a major role in the patho-physiological background of the pathogeneses of Aconite. So, it might be the possible background of producing pathology when Aconite seems to be indicated to prescribe as per Homoeopathic principle.

**Key-words:** Aconitum napellus (Aconite), Catecholamines, Homoeopathic pathogenesis, Noradrenalin (Norepinephrine).

# Introduction

Aconite roots are traditionally used in Chinese and Japanese medicine for analgesic, anti-rheumatic and neurological indications <sup>1</sup>. There are several alkaloids in aconite species. Among these alkaloids, Aconitine, 3-acetylaconitine and hypaconitine have been reported for acute toxicity <sup>2, 3</sup>.

It is also known that roots of Aconite contain physiologically active catecholamine analogs <sup>4</sup>. It has direct effect on adrenal and in release of catecholamines <sup>5</sup>. Aconitine has also an ability to increase plasma corticosterone level <sup>6</sup>.

In the Homoeopathic pathogenesis, Aconite is often used for acute inflammatory conditions in different systems or organs; fever; vertigo, fainting, palpitations as are from acute cardiac affections; cessation of menstruation, miscarriage related to female endocrinal disorders; neuralgias; acute haemorrhages etc. associated with mental anxiety, fear, restlessness, and increased thirst. These conditions are found to be caused by fear or fright, or from exposure to dry, cold weather 7-15.

Catecholamines, which are secreted from adrenal medulla comprises 75-80% adrenaline and 20-25% noradrenaline. Other than that neuronal sympathetic release of catecholamines comprises 80-90% is noradrenaline and 10-20% dopamine <sup>16</sup>. The Locus Coeruleus (LC), locaed adjacent to fourth ventricle in pontine brainstem is the largest group of noradrenergic neurones in Central Nervous System (CNS) <sup>17, 18</sup>. Besides that, phagocytes (including macrophages, neutrophils and blood polymorphonuclear cells) also synthesize and release catecholamines <sup>19</sup>. On the other side, catecholamines, particularly noradrenalin has direct excitatory action on hypothalamus to increase secretion of corticotrophin releasing hormone (CRH) and so to elevate circulatory adrenocorticotrophic hormone (ACTH) <sup>20-22</sup>. Secretion of catecholamines is stimulated in response to number of stressful events, related to fight-or-flight responses, exercise, hypoglycaemia, cold, haemorrhage and hypotension. And increased secretion may accompany different emotional reactions, like fear, anger, pain

Corresponding Author: Dr. Kisor Kumar Naskar, H.O.D., Department of Materia Medica, D. N. De Homoeopathic Medical Colleges & Hospital; 12, Gobinda Khatick Road; Kolkata – 700046, WB, India, E-mail: drkisorkumarnaskar@gmail.com

**How to cite this article:** Naskar KK, Mishra O: Hyperactivity of Catecholamines - a possible pathophysiological background of Homoeopathic pathogenesis of Aconitum napellus; AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 3 - 10

and sexual arousal <sup>23</sup>. Short term danger like fear factor activates the amygdale. The amygdale responds to this fear or danger by initiating an immediate sympathetic response through neuroendocrine system to restore homoeostasis, and in initial stage, signals the brain stem to release sympathetic catecholamine - norepinephrine. Once release in blood flow it increases heart rate, blood pressure and respiration, vasoconstriction of arterioles, stimulate sweat secretion and papillary dilatation. Importantly, this short term sympathetic response is proinflammatory, functioning to destroy antigen, pathogens or foreign invaders <sup>24</sup>.

In respect of the duration of course regarding stress related altered immune function, chronic stressors are deleterious. But, the short stressors (last less than two hours) also enhance some aspects of immune function as in sudden unpredictable or uncontrollable stress may cause more persistent immune dysregulation <sup>25</sup>.

Cold exposure stimulates catecholamine secretion. In whole body cold exposure, exposure of acute cold air, eating ice cubes, short time cold exposure even in cold season, nor-epinephrine level increases significantly. It was also found that, in very hot environment (in hyperthermia fatalities) level of noradrenalin was high with low adrenalin <sup>23, 26</sup>.

In the circadian clock, catecholamines secretion level peaked in late morning and become low at night during sleep but on the other hand, norepinephrine was greater in older subjects by 28% during the day and 75% at night <sup>27</sup>.

Similar to the characteristic Homoeopathic pathogeneses of Aconite, hyperactivity of catecholamines also cause panic disorders <sup>28-35</sup>, acute inflammatory responses <sup>36-53</sup>, fever <sup>18,54,55</sup>, increased susceptible to various infections <sup>56-79</sup>, acute heart affections even cardiogenic shock <sup>80-86</sup>, haemorrhage <sup>87-90</sup>, disturbed endocrinal activities related to secondary amenorrhoea or spontaneous preterm delivery <sup>91-96</sup>, neuralgia <sup>97,98</sup>, retinal vasculitis 99, periodontal diseases <sup>100-102</sup>, increased thirst <sup>103,104</sup> etc.

# Fundamentals of Homoeopathic Pathogenesis of Aconite from the experiences of Pioneers 7-15

- Aconite causes general functional disturbances, no evidence that it can produce tissue change its action is brief and shows no periodicity. Its sphere is in the beginning of an acute disease and not to be continued after pathological change comes <sup>7</sup>. It raises an arterial and nervous storm, and, though in fatal cases its fury may be great enough to induce chaos, that is, death, yet it does not localize itself in organic changes <sup>8</sup>. It seems to be useful only in acute diseases, yet it is an indispensable accessory remedy in even the most obstinate chronic affections, when the system requires a diminution of so called tension of the blood vessels <sup>9</sup>.
- Chill, fright, or surgical operation the effect of these will be met in large majority of cases by Aconite <sup>10</sup>. Ailments from fright; afraid in dark; vertigo; faintness; trembling; cardiac weakness; threatened miscarriage; impending cessation of menstrual flow; burning in stomach <sup>11</sup>. Complaints and tension caused by exposure to dry, cold weather, drought of cold air, checked perspiration, also complaints from very hot weather, especially gastro-intestinal disturbance, etc. <sup>7</sup>.
- A state of fear, anxiety; anguish of mind and body. Physical and mental restlessness, fright, is the most characteristic manifestations of Aconite <sup>9,7</sup>. In addition to thirst and quick pulse, present anxious impatience, an unappeasable mental agitation, and agonizing tossing about in the morbid state of acute inflammations of any part to those seen in persons who have had a fright combined with vexation, is the surest and quickest remedy for them <sup>9</sup>.
- Acute, sudden and violent invasion, with fever. Fever in which Aconite is specific is neurotic, not toxaemic
  or sympathetic in nature. It is "synocha" of old authors, the pure inflammatory fever with full and bounding
  pulse, great heat and restlessness, unquenchable thirst for large quantities of water, with extreme nervous
  excitability and tossing about in agony, worse towards evening and after getting in bed <sup>7, 12, 13</sup>.
- It may be employed in meningitis, ophthalmia, tonsillitis, croup, bronchial catarrh, pneumonia, pulmonary congestion and haemoptysis, pleuritis, pericarditis and endocarditis, gastritis, peritonitis, acute rheumatism, neuralgia supra-orbitalis but only when moral symptoms named by Hahnemann are present as characteristic indications of Aconite, "the anguish of mind and body, the restlessness, the disquiet not to be allayed" <sup>8</sup>. In inflammation of the bladder, in suppressed urine, or menses, and endless other conditions resulting from or accompanied by chill, shock, fright, fear <sup>14</sup>. Toothache caused by dry, cold wind; gastric catarrh from chilling stomach with ice water, especially when overheated. Pain intolerable; more so at night <sup>11</sup>.
- Plethora; active capillary congestions <sup>11</sup>. Active haemorrhage in stout, plethoric people. Passes almost pure blood by stool. In haemoptysis the blood comes up with great ease by hemming and coughing, bright red in large quantities, from cold, dry winds, with great fear, anxiety and palpitations <sup>10</sup>.
- It is very useful in all diseases of the heart characterised by increased action <sup>12</sup>. It has wonderful soothing effect in heart disease, where an acute condition has supervened, with palpitations, anguish, and great distress <sup>14</sup>. On rising from a recumbent position the red face become deathly pale, or he becomes faint or giddy and falls, and he fears to raise again; often accompanied by vanishing of sight and unconsciousness <sup>15</sup>
- In other conditions Urethral fever ensues upon catheterism <sup>12</sup>. Facial paralysis from exposure to cold and

dry winds <sup>10</sup>. Almost specific in facial neuralgia, especially of congestive form. Incipient inflammation of the eye after mechanical injury, whether accidental or operative <sup>12</sup>.

# Action of Catecholamines -

## **Emotional disturbances – Panic disorders:**

Along with other systemic affections, massive secretion of catecholamines (due to uncontrolled stress response) may also responsible for fear, anxiety in the level of panic disorders. In this respect, noradrenergic over activity due to increased and altered regulations of brain noradrenergic functions, related to elevated LC neurones firing, is often associated with anxiety and fear responses as in panic disorders <sup>28-34</sup>.

It has been reported that Norepinephrine Bitartrate has adverse side effect to manifest fear of death 35.

# Immune response – Pro/anti-inflammatory activity:

The CNS, endocrine system and immune system are complex systems and they interact with each other. In between CNS and immune system, there is reciprocal regulation exists. CNS signals immune system via hormonal and neuronal pathway, and immune system signals the CNS through cytokines <sup>36</sup>.

Short term stressors increase natural killer cells activity, increase the number of some types of leukocytes. Stress hormones modulate immune cell function by binding its receptors at the surface of cells within lymphoid organs. Norepinephrine is released from the sympathetic nerve terminals and the target immune cells express adrenoreceptors. Stimulation of these receptors, locally released norepinephrine, or circulating catecholamine affect lymphocyte traffic, circulation, and proliferation as well as modulate cytokine production and the functional activity of different lymphoid cells <sup>37-44</sup>. In a study, norepinephrine has found to modulate inflammatory and proliferative phases of wound healing by increasing recruitment of innate immune cells and expediting wound closure <sup>45</sup>. It was also observed that not only pro-inflammatory, but role of norepinephrine has been found in both pro- as well as anti-inflammatory under stressful conditions, depends on the duration, intensity, perception, acute or chronic in nature, time point - whether before outbreak or in later phases of the symptomatic disease in relation to the outbreak of a chronic inflammatory disease. In this respect in a study it has been found that since application of noradrenalin at high concentrations (10-<sup>6</sup> M) via 2-adrenoceptors is anti-inflammatory by strongly inhibiting Tumour-necrosis factor (TNF) and the other side, the loss of sympathetic nerve fibers in inflamed tissue of rheumatoid arthritis patients leads to a pro-inflammatory situation <sup>46</sup>.

On the other hand, IL-6, a type of cytokines, produced by T cells, B cells, monocytes and several non-lymphoid cell types has pro-inflammatory action. It has an important role in acute phase response <sup>47</sup>. Both physical and psychological stressors can provoke transient increase of this pro-inflammatory cytokines <sup>48, 49</sup>. It has been demonstrated that norepinephrine stimulated IL-6 production <sup>50, 51</sup>. IL-6 is also an important inducer of C-reactive protein (CRP) from the liver. Combination of IL-6 and CRP may cause development of cardiovascular disease also <sup>47, 52</sup>. Besides that, it has also link with other various age related diseases like osteoporosis, arthritis, type 2 diabetes, frailty and functional decline even certain cancer such as chronic lymphocytic leukaemia <sup>53</sup>.

In another study, it has been observed that in acute condition phagocytes release catecholamines which directly enhance inflammatory response <sup>19</sup>.

# Fever:

In response to systemic inflammation brain plays a central role and causes fever. Febrigenic peripheral signals including blood born cytokines may gain access to brain through different afferent pathways and increase the level of Prostaglandin E2 (PGE2). PGE2 triggers thermoeffector response and increases body core temperature. In this respect involvement of brain noradrenalin has been observed to release of PGE2 <sup>18, 54</sup>. LC, the major noradrenergic nucleus in brain, is specifically activated and ultimately increases the level of PGE2 to activate thermogenesis <sup>55</sup>.

# Increased susceptibility to different pathogenetic organisms:

Elevated levels of catecholamines activity may increases susceptibility to diseases <sup>56</sup>. Norepinephrine acts as a growth stimulating factor of the pathogens <sup>57-62</sup>. It induces not only adhesion to the host cells but also supplies iron for the growth of bacteria <sup>63-66</sup>. As an example it has been found in Escherichia Coli, catecholamines can increase adhesion to host cells by up-regulating adherence association genes and alter host susceptibility to enteric infection <sup>67, 68</sup>. Bordetella cells take benefit from catecholamines to grow on respiratory epithelium <sup>69</sup>.

In catheter-related urinary tract infection, the common pathogenic bacteria are Pseudomonas aeruginosa, Klebsiella pneumoniae, Proteus mirabilis, E. coli, coagulase-negetive staphylococci etc. <sup>70-72</sup>. Different research works established that catecholamines, especially norepinephrine acts as a growth stimulating factor for these pathogenic microorganisms <sup>57-62</sup>.

Intestinal pathogenic micro-organisms like Shigella, Salmonella, E. coli etc. increase their number when concentration of catecholamine increased especially noradrenalin and cause intestinal disorders like diarrhoea, dysentery or vomiting <sup>61, 62, 67, 68, 73, 74</sup>.

Herpes simplex virus (HSV) is a natural human pathogen which is the most important causative organism for facial paralysis <sup>105-107</sup>. Stress not only increases the development and severity of HSV infection in both the peripheral nervous system and CNS, but also suppresses components of primary cytotoxic T lymphocyte response to HSV infection 75-79, 39. Stress related neuro-hormones, norepinephrine and epinephrine modify the cytolytic activity of macrophages against virus infected cells and suggest a possibleneuroendocrine immunologic basis for the recurrence of HSV infection 108. Regarding teeth and gum affections, stress induced changes in local catecholamines level and plays a significant role in the aetiology and pathogenesis of periodontal diseases. It is observed that autoinducer mechanisms may play an important role in the response of oral microorganisms to these stress hormones; thereby contributing to the clinical course of stress associated periodontal diseases 100-102.

# **Heart:**

Aconite has direct action on myocardium. After consumption of its roots, it may cause life threatening ventricular tachycardia 80. It has been described that, Aconite induced arrhythmia does not seem to be the effect of neural cause but due to centrally evoked release of endogenous catecholamines 81.

In this respect, over stimulated sympathetic nerve activity which may be induced by violent psychological stress like anxiety or any other means is resulting excessive concentrations of catecholamines capable of producing myocardial necrosis, even in the non-ischemic heart. Not only that, it may also responsible for the pathogenesis of ventricular fibrillation in early ischemia 82.

Besides that, it has been observed noradrenergic over activity in the LC coinciding with the development of the hypertension 83. Systemic surges in catecholamines cause coronary vasospasm and severe acute hypertension due to acute peripheral vasospasm which may be followed by peripheral vasodilatation and hypotension. So, clinically profound hypotension and cardiogenic shock are may be common complications which are not in extent of left ventricular impairment 84.

# Dizziness and fainting - related to Cardiogenic shock/ Postural or orthostatic intolerance:

Increased secretion of catecholamines is one of the causes of these conditions. Acute coronary as well as peripheral vasospasms are the resulting manifestations of systemic surges in catecholamines which may be followed by vasodilatations, clinically found as hypotension and cardiogenic shock 84.

Sudden raise of plasma norepinephrine levels is related to postural or orthostatic intolerance. It is characterised by lightheadedness, dizziness, palpitations, sweating, nausea and syncope. It primarily occurs not only with upright posture but sudden rising from a recumbent position are the precipitating circumstances and relieved by lying down 85,86.

# Capillary vessels and Haemorrhage:

Sudden and sustained increases in systemic catecholamines potentiate the activation of endothelium causing vasospasm 87. Catecholamines of peripheral systemic nerve endings lead to peripheral veso-constriction which subsequently increases systemic vascular resistance and systemic blood pressure. Moreover, it increases central blood volume along with reduction in the compliance of left ventricle. All of these changes are followed by increase in pulmonary capillary hydrostatic pressure and damage to alveolar wall leads to leakage of fluid into the interstitium as well as intra alveolar space and resulting haemorrhage - the typical picture of neurogenic pulmonary oedema 88.

It is also observed that association between presences of enlarged vessels at rhinoscopy with history of epistaxis in hypertensive patients 89. An association between duration of hypertension and left ventricular hypertrophy and nasal artery enlargement determined by rhinoscopy might contribute some relation between these 90.

## Secondary amenorrhoea – Pregnancy - Spontaneous Preterm delivery:

In case of secondary amenorrhoea, there is no such direct role of catecholamine in this regard. But stress induced catecholamine especially noradrenalin can directly excites upon CRH secretion and CRH may directly inhibit gonadotropin secretion which causes functional secondary amenorrhoea 91-93.

High catecholamines level in mid-pregnancy may be indicative of excess stressors and/or predisposition to elevated sympathetic activation related to the risk of spontaneous Pre-term delivery 94. In another aspect, increased catecholamines cause uterine contractions by interrelationship action of alpha- and beta-adrenergic stimulation and the prostaglandin system 95. Norepinephrine increases coordinate uterine activity and a presser response 96.

It has been found that sympathetically maintained pain (neuralgia) become painfully hypersensitive to norepinephrine

#### Retinal vasculitis:

Retinal vasculitis has been observed as an adverse side effect of noradrenaline 99.

# Thirst:

Involvement of catecholaminergic activity is stronger in induced drinking mechanism. In an animal (rat) study intracranial administration of noradrenaline frequently causes a burst of drinking before it starts to eat 103. Dopaminergic neural system involves influencing polydipsia in humans. Elevated dopamine levels stimulate thirst centre 104.

On the other hand dry tongue as well as dry mouth is usual phenomena in panic attack of anxiety and it may causes burning or unquenchable thirst.

# **Conclusion**

This study is intended to find out the area of altered physiological functions and developing internal pathologies of Aconite responsible for its pathogeneses. It has been found that fundamentals of Homoeopathic pathogeneses of Aconite are similar to the manifestations of patho-physiological changes resulting from hyperactivity of catecholamines especially noradrenalin (norepinephrine) in different systems as well as organs through neuro-endocrinal pathway. So, it might be concluded that hyperactivity of catecholamines especially noradrenalin is the possible initiating responsible pathological essence to develop pathogeneses of Aconite and similarly possible background of producing pathology when Aconite seems to be indicated to prescribe as per Homoeopathic principle.

#### References

- 1. Ameri A; The effects of Aconitum alkaloids on the central nervous system; Prog Neurobiol. 1998 Oct;56(2):211-35
- 2. Gutser U.T., Friese J., Heubach J. F., Matthiesen T., Selve N., Wilffert B., Gleitz J.; Mode of antinociceptive and toxic action of alkaloids of Aconitum spec.; Naunyn-Schmiedeberg's Archives of Pharmacology; 1997 Dec, 357(1), 39–48
- 3. Chan Thomas Y.K.; Aconite poisoning; Clinical Toxicology; 2009, 47(4), 279-285
- 4. SenthilkumaranS, MeenakshisundaramR, Thirumalaikolundusubramanian P; Plant Toxins and the Heart (Chapter 5). In Heart and Toxins, Meenakshisundaram R; Academic Press Elsevier; 2015, P-157;
- 5. Afolabi Clement Akinmoladum, Mary Tolulope Olabye and Ebenezer Olatunde Farombi; Cardiotoxicity and Cardioprotective Effects of African Medicinal Plants; In Toxicological Survey of African Medicinal Plants, Victor Kuete; Elsevier; 2014; p-403.
- Kimura I., Makino M., Honda R., Ma J. and Kimura M.- Expression of major histocompatibility complex in mouse peritoneal macrophages increasingly depends on plasma corticosterone levels; stimulated by aconite.- Biol Pharm Bull. – 1995; 18(11): 1504-1508, 8593467
- 7. Boericke W, Pocket manual of Homoeopathic Materia Medica, 9th edition, New Delhi, B. Jain Pub. Pvt. Ltd, 1994 (Reprint), pp 7-11
- 8. Dunham C, Lectures on Materia Medica, 5th edition, New Delhi, B. Jain Pub. Pvt. Ltd, Vol-1, 2004 (Reprint), pp 66-88
- 9. Hahnemann S, Materia Medica Pura, New Delhi, B. Jain Pub. Pvt. Ltd, Vol-1, 1999 (Reprint), pp 25-46
- Clarke JH, A Dictionary of Practical Materia Medica, New Delhi, B. Jain Pub. Pvt. Ltd, vol-1, 1999 (Reprint), Vol.1, pp-15-25
- 11. Hering C, The Guiding Symptoms of our Materia Medica, New Delhi, B. Jain Pub. Pvt. Ltd, Vol-1, 2000 (Reprint), pp 28-55
- 12. Hughes R, A manual of Pharmacodynamics, 6th edition, New Delhi, B. Jain Pub. Pvt. Ltd, 1999 (Reprint), pp 147-166
- 13. Blackwood AL, A manual of Materia Medica Therapeutics and Pharmacology, 2nd edition, New Delhi, B. Jain Pub. Pvt. Ltd, 1999 (Reprint), pp 83-88
- 14. Tyler ML, Homoeopathic Drug Pictures, New Delhi, B. Jain Pub. Pvt. Ltd, 2000 (Reprint), pp 4-11
- 15. Allen H C, Keynotes and characteristics with comparisons of some of the leading remedies of the Materia Medica with Bowel Nosodes, 8th edition, New Delhi, B. Jain Pub. Pvt. Ltd., 2002 (Reprint), p 14
- Lechin Fuad, Dijs Bertha van der, Lechin Alex E.; Circulating Serotonin, Catecholamines, and Central Nervous System Circuitry Related to Some Cardiorespiratory, Vascular, and Hematological Disorders; The Journal of Applied Research, Vol. 5, No. 4, 2005; 605-21
- 17. Berridge CW, Waterhouse BD; The locus coeruleus-noradrenergic system: modulation of behavioral state and state-dependent cognitive processes; Brain Res Brain Res Rev. 2003 Apr;42(1):33-84
- 18. Almeida Maria C, Steiner Alexandre A, Coimbra Norberto C, and Branco Luiz G S; Thermoeffector neuronal pathways in fever: a study in rats showing a new role of the locus coeruleus; J Physiol. 2004 Jul 1; 558(Pt 1): 283–294
- 19. Flierl MA, Rittirsch D, Nadeau BA, Chen AJ, Sarma JV, Zetoune FS et al.; Phagocyte-derived catecholamines enhance acute inflammatory injury; Nature. 2007 Oct 11;449(7163):721-5
- 20. Calogero AE, Gallucci WT, Chrousos GP, Gold PW. Catecholamine effects upon rat hypothalamic corticotropin-releasing hormone secretion in vitro. Journal of Clinical Investigation. 1988;82(3):839-846.
- 21. Itoi K, Suda T, Tozawa F, Dobashi I, Ohmori N, Sakai Y et al; Micro injection of norepinephrine into the paraventricular nucleus of the hypothalamus stimulates corticotrophin-releasing factor gene expression in conscious rats; Endocrinology; 1994, Nov.,
- 22. Al-Damluji S, Rees LH. Effects of catecholamines on secretion of adrenocorticotrophic hormone (ACTH) in man. Journal of Clinical Pathology. 1987;40(9):1098-1107
- 23. Pääkkönen Tiina, Leppäluoto Juhani; Cold exposure and hormonal secretion: A review; International journal of circumpolar health September 2002
- 24. Hannibal Kara E., Bishop Mark D.; Chronic Stress, Cortisol Dysfunction, and Pain: A Psychoneuroendocrine Rationale for Stress Management in Pain Rehabilitation; Phys Ther. 2014 Dec; 94(12): 1816–1825
- 25. Ironson Gail, Wynings Christina, Schneiderman Neil, Baum Andrew, Rodriguez Mario, Greenwood Debra et al; Posttraumatic Stress Symptoms, Intrusive Thoughts, Loss, and Immune Function After Hurricane Andrew; Psychosomatic Medicine 59:128-141 (1997)
- 26. Kortelainen ML, Huttunen P, Lapinlampi T.; Urinary catecholamines in hyperthermia-related deaths; Forensic Sci Int. 1990 Nov;48(1):103-10
- 27. Prinz PN, Halter J, Benedetti C, Raskind M.; Circadian variation of plasma catecholamines in young and old men: relation to rapid eye movement and slow wave sleep; J Clin Endocrinol Metab. 1979 Aug;49(2):300-4
- 28. Wilkinson DJC, Thompson JM, Lambert GW, Jennings GL, Schwarz RG, Jefferys D et al.; Sympathetic Activity in

- Patients With Panic Disorder at Rest, Under Laboratory Mental Stress, and During Panic Attacks. Arch Gen Psychiatry. 1998;55(6):511-520.
- 29. Charney DS, Woods SW, Goodman WK, Heninger GR; Neurobiological mechanisms of panic anxiety: biochemical and behavioral correlates of yohimbine-induced panic attacks; Am J Psychiatry; 1987 Aug; 144(8); 1030-6
- 30. Charney DS, Woods SW, Nagy LM, Southwick SM, Krystal JH, Heninger GR; Noradrenergic function in panic disorder; J. Clin. Psychiatry; 1990; 51(Suppl A):5-11
- 31. Uhde TW, Boulenger JP, Post RM, Siever LJ, Vittone BJ, Jimerson DC et al; Fear and anxiety: relationship to noradrenergic function; Psychopathology; 1984; 17(Suppl 3):8-23
- 32. Lee YJ, Hohoff C, Domschke K, Sand P, Kuhlenbäumer G, Schirmacher A et al. Norepinephrine transporter (NET) promoter and 5'-UTR polymorphisms: association analysis in panic disorder. Neurosci Lett. 2005;377:40–43
- 33. Buttenschon HN, Kristensen AS, Buch HN, Andersen JH, Bonde JP, Grynderup M et al. The norepinephrine transporter gene is a candidate gene for panic disorder. J Neural Transm (Vienna) 2011;118:969–976
- 34. Park HJ, Kim SK, Kang WS, Kim YJ, Cho AR, Park JK. Potential involvement of NET polymorphism in serotonin/ norepinephrine reuptake inhibitor response in panic disorder. Nord J Psychiatry. 2016;70:314–317.
- 35. FactMed; Is fear of death a side effect of Norepinephrine Bitartrate?; factmed.com/study-NOREPINEPHRINE%20 BITARTRATE-causing-FEAR%20OF%2...; 26/02/2017
- 36. Webster JI, Tonelli L, Sternberg EM; Neuroendocrine regulation of immunity; Annu Rev Immunol. 2002;20:125-63. Epub 2001 Oct 4.
- 37. Segerstrom Suzanne C. and Miller Gregory E.; Psychological Stress and the Human Immune System: A Meta-Analytic Study of 30 Years of Inquiry; Psychol Bull. 2004 Jul; 130(4): 601–630.
- 38. Altemus M, Rao B, Dhabhar FS, Ding W, Granstein RD.; Stress-induced changes in skin barrier function in healthy women; J Invest Dermatol. 2001 Aug;117(2):309-17
- 39. Glaser Ronald and Glaser Janice K. Kiecolt; Stress-induced immune dysfunction: implications for health; Immunology; Volume 5, March 2005; 243-51
- 40. Srinivasan Thyaga Rajan, Hannah Priyanka and Uday Pundir; Aging Alters Sympathetic Noradrenergic Innervation and Immune Reactivity in the Lymphoid Organs: Strategies to Reverse Neuro-Immune Senescence; Brain Immune Trends; http://brainimmune.com/aging-alters-sympathetic-noradrenergic-innervation-and-immune-reactivity-in-the-lymphoidorgans-strategies-to-reverse-neuro-immune-senescence/; Posted on July 29, 2012
- 41. Elenkov Ilia J., Wilder Ronald L., Chrousos George P., and Vizi E. Sylvester; The sympathetic Nerve—An Integrative Interface between two supersystems: The Brain and the Immune System; Pharmacol Rev 52:595–638, 2000
- 42. Hübner G, Brauchle M, Smola H, Madlener M, Fässler R, Werner S.; Differential regulation of pro-inflammatory cytokines during wound healing in normal and glucocorticoid-treated mice; Cytokine. 1996 Jul;8(7):548-56
- 43. Padgett DA, Marucha PT, Sheridan JF; Restraint stress slows cutaneous wound healing in mice; Brain Behav Immun. 1998 Mar;12(1):64-73
- 44. Glaser R, Kiecolt-Glaser JK, Marucha PT, MacCallum RC, Laskowski BF, Malarkey WB; Stress-related changes in proinflammatory cytokine production in wounds; Arch Gen Psychiatry. 1999 May;56(5):450-6
- 45. Gosain A, Jones SB, Shankar R, Gamelli RL, DiPietro LA; Norepinephrine modulates the inflammatory and proliferative phases of wound healing; J Trauma. 2006 Apr;60(4):736-44
- 46. Rainer H Straub and Joachim R Kalden; Stress of different types increases the proinflammatory load in rheumatoid arthritis; Arthritis Res Ther. 2009; 11(3): 114
- 47. Black PH; The inflammatory response is an integral part of the stress response: Implications for atherosclerosis, insulin resistance, type II diabetes and metabolic syndrome X; Brain Behav Immun. 2003 Oct;17(5):350-64
- 48. DeRijk R, Michelson D, Karp B, Petrides J, Galliven E, Deuster P et al.; Exercise and circadian rhythm-induced variations in plasma cortisol differentially regulate interleukin-1 beta (IL-1 beta), IL-6, and tumor necrosis factor-alpha (TNF alpha) production in humans: high sensitivity of TNF alpha and resistance of IL-6; J Clin Endocrinol Metab. 1997 Jul;82(7):2182-91
- 49. Zhou D, Kusnecov AW, Shurin MR, DePaoli M, Rabin BS; Exposure to physical and psychological stressors elevates plasma interleukin 6: relationship to the activation of hypothalamic-pituitary-adrenal axis; Endocrinology. 1993 Dec:133(6):2523-30
- 50. von Patay B, Loppnow H, Feindt J, Kurz B, Mentlein R; Catecholamines and lipopolysaccharide synergistically induce the release of interleukin-6 from thymic epithelial cells; J Neuroimmunol. 1998 Jun 15;86(2):182-9
- 51. von Patay B, Kurz B, Mentlein R; Effect of transmitters and co-transmitters of the sympathetic nervous system on interleukin-6 synthesis in thymic epithelial cells; Neuroimmunomodulation. 1999 Jan-Apr;6(1-2):45-50
- 52. Black PH; Stress and the inflammatory response: a review of neurogenic inflammation; Brain Behav Immun. 2002
- 53. Harris Tamara B., Ferrucci Luigi, Tracy Russell P., Corti M. Chiara, Wacholder Sholom, et al.; Associations of Elevated Interleukin-6 and C-Reactive Protein Levels with Mortality in the Elderly; The American Journal of Medicine; Volume 106; May 1999; pp-506-12
- 54. Blatteis Clark M. and Sehic Elmir; Fever: How May Circulating Pyrogens Signal the Brain?; News Physiol. Sci. I Volume 12 I February 1997
- 55. Cannon B, Houstek J, Nedergaard J.; Brown adipose tissue. More than an effector of thermogenesis?; Ann N Y Acad Sci. 1998 Sep 29;856:171-87.
- 56. Gruchow HW: Catecholamine activity and reported morbidity. Journal of Chronic Disease, 29:773-783, 1976; http://www.

- elsevier.com/
- 57. Wang Li, Mark Lyte, Primrose P. Freestone, Aziba Ajmal, Jane A. Colmer-Hamood, Abdul N. Hamood.; Norepinephrine represses the expression of toxA and the siderophore genes in Pseudomonas aeruginosa; FEMS Microbiol Lett., 2009 Oct; 299(1):100-109
- 58. Kinney KS, Austin CE, Morton DS, Sonnenfeld G.; Norepinephrine as a growth stimulating factor in bacteria--mechanistic studies; Life Sci. 2000 Nov 10;67(25):3075-85.
- 59. Belay T, Sonnenfeld G.; Differential effects of catecholamines on in vitro growth of pathogenic bacteria; Life Sci. 2002 Jun 14:71(4):447-56
- 60. Mark Lyte, Primrose PE Freestone, Christopher P Neal, Barton A Olson, Richard D Haigh, Roger Bayston et al.; Stimulation of Staphylococcus epidermidis growth and biofilm formation by catecholamine inotropes; The Lancet; Vol.361, No.9352, p130-135, Jan 2003.
- Freestone P.P.E, Lyte M., Haigh R.D., Williams P.H., Stimulation of bacterial growth by heat-stable norepinephrineinduced autoinducers, FEMS Micro. Letts., 1999, 172, 53-60
- 62. Sharaff Fathima, Freestone Primrose; Microbial Endocrinology; Cent. Eur. J. Biol., 6(5); 2011; 685-694
- 63. Lu Li, Zhuofei Xu, Yang Zhou, Lili Sun, Ziduo Liu, Huanchun Chen et al.; Global Effects of Catecholamines on Actinobacillus pleuropneumoniae Gene Expression; PLOS; February 8, 2012
- 64. Freestone PPE, Lyte M, Neal CP, Maggs AF, Haigh RD, Williams PH. The Mammalian Neuroendocrine Hormone Norepinephrine Supplies Iron for Bacterial Growth in the Presence of Transferrin or Lactoferrin. Journal of Bacteriology. 2000;182(21):6091-6098.
- 65. Burton Claire L., Chhabra Siri Ram, Swift Simon, Baldwin Tom J., Withers Helen, Hill Stephen J., et al. The Growth Response of Escherichia coli to Neurotransmitters and Related Catecholamine Drugs Requires a Functional Enterobactin Biosynthesis and Uptake System. Infection and Immunity. 2002;70(11):5913-5923. doi:10.1128/IAI.70.11.5913-5923.2002.
- Anderson MT, Armstrong SK. Norepinephrine Mediates Acquisition of Transferrin-Iron in Bordetella bronchiseptica. Journal of Bacteriology. 2008;190(11):3940-3947. doi:10.1128/JB.00086-08.
- 67. Vlisidou I, Lyte M, van Diemen PM, Hawes P, Monaghan P, Wallis, TS et al.; The Neuroendocrine Stress Hormone Norepinephrine Augments Escherichia coli O157:H7-Induced Enteritis and Adherence in a Bovine Ligated Ileal Loop Model of Infection; Infection and Immunity, 2004 Sep; 72(9), 5446–5451.
- 68. Chunsheng Chen, Mark Lyte, Mark P. Stevens, Lucy Vulchanova, and David R. Brown; Mucosally-directed adrenergic nerves and sympathomimetic drugs enhance non-intimate adherence of Escherichia coli O157:H7 to porcine cecum and colon; Eur J Pharmacol. 2006 Jun 6; 539(0): 116–124
- 69. Anderson MT, Armstrong SK. The Bordetella Bfe System: Growth and Transcriptional Response to Siderophores, Catechols, and Neuroendocrine Catecholamines. Journal of Bacteriology. 2006;188(16):5731-5740. doi:10.1128/JB.00495-06.
- John L Brusch; Catheter-Related Urinary Tract Infection; Medscape; emedicine.medscape.com/article/2040035overview; Updated: Aug 18, 2015
- 71. Nicolle LE. Catheter associated urinary tract infections. Antimicrobial Resistance and Infection Control. 2014;3:23. doi:10.1186/2047-2994-3-23.
- 72. Gahlot R, Nigam C, Kumar V, Yadav G, Anupurba S. Catheter-related bloodstream infections. International Journal of Critical Illness and Injury Science. 2014;4(2):162-167. doi:10.4103/2229-5151.134184.
- 73. Walker C.E. and Drouillard J.S.; Effects of Catecholamines on Gut Microflora and Potential for Beta-Adrenergic Agonists to Impact Ruminal Fermentation; The Open Agriculture Journal, 2012, 6, 57-66
- 74. O'Donnell Phyllis M., Aviles Hernan, Lyte Mark, and Sonnenfeld Gerald; Enhancement of In Vitro Growth of Pathogenic Bacteria by Norepinephrine: Importance of Inoculum Density and Role of Transferrin; Appl Environ Microbiol. 2006 Jul; 72(7): 5097–5099
- 75. Kusnecov AV., Grota LJ, Schmidt SG, Bonneau RH, Sheridan JF, Glaser R et al.; Decreased herpes simplex viral immunity and enhanced pathogenesis following stressor administration in mice. J. Neuroimmunol. 38, 129–137(1992)
- 76. Wonnacott KM, Bonneau RH; The effects of stress on memory cytotoxic T lymphocyte-mediated protection against herpes simplex virus infection at mucosal sites; Brain Behav Immun. 2002 Apr;16(2):104-17
- 77. Cao, L., Martin, A., Polakos, N. & Moynihan, J. A.; Stress causes a further decrease in immunity to herpes simplex virus-1 in immunocompromised hosts; J. euroimmunol.;156, 21–30 (2004)
- 78. Bonneau R H., Sheridan J F., Feng N.; Stress-induced modulation of the primary cellular immune response to herpes simplex virus infection is mediated by both adrenal-dependent and independent mechanisms; J. Neuroimmunol. 42, 167–176 (1993)
- 79. Leo NA, Bonneau RH; Mechanisms underlying chemical sympathectomy-induced suppression of herpes simplex virusspecific cytotoxic T lymphocyte activation and function; J Neuroimmunol. 2000 Oct 2;110(1-2):45-56
- 80. Weijters B.J., Verbunt R.J.A.M., Hoogsteen J., and Visser R.F.; Salade malade: malignant ventricular arrhythmias due to an accidental intoxication with Aconitum napellus; Neth Heart J. 2008 Mar; 16(3): 96–99
- 81. Bhargava K. P., Kohli R. P., Sinha J. N. and Tayal G.; Role of catecholamines in centrogenic cardiac arrhythmia induced by aconitine; Br. J. Pharmac. (1969), 36, 240-252.
- 82. Schömig A; Catecholamines in myocardial ischemia. Systemic and cardiac release; Circulation. 1990 Sep;82(3 Suppl):II13-22.
- 83. Koulu M., Saavedra J.M., Niwa M., Linnoila M.; Increased catecholamine metabolism in the locus coeruleus of young spontaneously hypertensive rats; Brain Research; Volume 369, Issues 1–2, 26 March 1986, Pages 361–364
- 84. Akashi Yoshihiro J., Nef Holger M., Lyon Alexander R.; Epidemiology and Pathophysiology of Takotsubo Syndrome; Nat Rev Cardiol; 12,387–397(2015)
- 85. Robertson D, Flattem N, Tellioglu T, Carson R, Garland E, Shannon JR et al; Familial orthostatic tachycardia due to

- norepinephrine transporter deficiency; Ann NY Acad Sci. 2001 Jun; 940:527-43
- 86. Miklos Szathmari; Syncope and Shock; Semmelwel University First department of Medicine; 05 Nov 2013; www.bel1. semmelweis.hu
- 87. Chen Sheng, Li Qian, Wu Haijian, Krafft Paul R., Wang Zhen and Zhang John H.; The Harmful Effects of Subarachnoid Hemorrhage on Extracerebral Organs; BioMed Research International; Volume 2014 (2014)
- 88. Šedý J., Zicha J., , J. Kuneš J., Jendelová P., Syková E.; Mechanisms of Neurogenic Pulmonary Edema Development; Physiol. Res. 57: 499-506, 2008
- 89. Lubianca Neto JF, Fuchs FD, Facco SR, Gus M, Fasolo L, Mafessoni R et al.; Is epistaxis evidence of end-organ damage in patients with hypertension?; Laryngoscope. 1999 Jul;109(7 Pt 1):1111-5.
- 90. Sarhana Nabil Abdulghany and Algamal Abdulsalam Mahmoud; Relationship between epistaxis and hypertension: A cause and effect or coincidence?; J Saudi Heart Assoc. 2015 Apr; 27(2): 79–84
- 91. Meczekalski B, Katulski K, Czyzyk A, Podfigurna-Stopa A, Maciejewska-Jeske M. Functional hypothalamic amenorrhea and its influence on women's health. Journal of Endocrinological Investigation. 2014;37(11):1049-1056. doi:10.1007/s40618-014-0169-3.
- 92. El bieta Sowi ska-Przepiera, El bieta Andrysiak-Mamos, Gra yna Jarz bek-Bielecka, Aleksandra Walkowiak, Lilianna Osowicz-Korolonek, Małgorzata Syrenicz et al.; Functional hypothalamic amenorrhoea diagnostic challenges, monitoring, and treatment; Endokrynol Pol 2015; 66 (3): 252-268
- 93. Fourman LT, Fazeli PK; Neuroendocrine Causes of Amenorrhea—An Update. The Journal of Clinical Endocrinology and Metabolism. 2015;100(3):812-824. doi:10.1210/jc.2014-3344.
- 94. Holzman Claudia, Senagore Patricia, Tian Yan, Bullen Bertha, DeVos Eric, Leece Cheryl et al. Maternal Catecholamine Levels in Midpregnancy and Risk of Preterm Delivery. American Journal of Epidemiology. 2009;170(8):1014-1024. doi:10.1093/aje/kwp218.
- 95. Quaas L and Zahradnik HP; The effects of alpha- and beta-adrenergic stimulation on contractility and prostaglandin (prostaglandins E2 and F2 alpha and 6-keto-prostaglandin F1 alpha) production of pregnant human myometrial strips. Am J Obstet Gynecol. 1985;152(7 pt 1):852–856.
- 96. Zuspan Frederick P, Cibils Luis A, Pose Serafin V; Myometrial and cardiovascular responses to alterations in plasma epinephrine and norepinephrine; Am J Obstet Gynecol. 1962;84:841–851.
- 97. Torebjörk E, Wahren L, Wallin G, Hallin R, Koltzenburg M; Noradrenaline-evoked pain in neuralgia; Pain. 1995 Oct;63(1):11-20.
- 98. Xanthos Dimitris N, Bennett Gary J and Coderre Terence J; Norepinephrine-induced nociception and vasoconstrictor hypersensitivity in rats with chronic post-ischemia pain; Pain. 2008 Jul 31; 137(3): 640–651.
- FactMed; NORADRENALINE patients who developed RETINAL VASCULITIS About this FactMed analysis covering adverse side effect reports; factmed.com/report-NORADRENALINE-causing-RETINAL%20VASCULITIS.php; - 19th July 2017
- 100. Robert A, Matthews JB, Socransky SS, Freestone PP, Williams PH, Chapple IL; Stress and the periodontal diseases: effects of catecholamines on the growth of periodontal bacteria in vitro; Oral Microbiol Immunol; 2002 Oct; 17(5):296-303
- 101. Robert A, Matthews JB, Socransky SS, Freestone PP, Williams PH, Chapple IL; Stress and the periodontal diseases: growth responses of periodontal bacteria to Escherichia coli stress-associated autoinducer and exogenous Fe; Oral Microbiol Immunol; 2005 Jan; 20(3):147-53
- 102. Robert A; The Role of Microbial Endocrinology in Periodontal Disease. In: Lyte M., Freestone P. (eds) -Microbial Endocrinology; Springer, New York, NY, 2010, pp135-150
- 103. James T. Fitzsimons; The Physiological basis of Thirst; Kidney International; vol.10; 1976; pp 8-9
- 104. Mittleman G, Rosner AL, Schaub CL; Polydipsia and dopamine: behavioral effects of dopamine D1 and D2 receptor agonists and antagonists; J Pharmacol Exp Ther. 1994 Nov;271(2):638-50.
- 105. Schirm J, Mulkens PS; Bell's palsy and herpes simplex virus; APMIS. 1997 Nov;105(11):815-23
- 106. Paulo Roberto Lazarini, Melissa Ferreira Vianna, Mônica Porto Alves Alcantara, Rodolfo Alexander Scalia, Hélio Hehl Caiaffa Filho; Herpes Simplex Virus in the saliva of peripheral Bell's palsy patients; Rev. Bras. Otorrinolaringol. vol.72 no.1 São Paulo Jan./Feb. 2006
- 107. Murakami S, Mizobuchi M, Nakashiro Y, Doi T, Hato N, Yanagihara N; Bell Palsy and Herpes Simplex Virus: Identification of Viral DNA in Endoneurial Fluid and Muscle; Ann Intern Med. 1996;124(1):27-30
- 108. Koff WC and Dunegan MA., Neuroendocrine hormones suppress macrophage-mediated lysis of herpes simplex virus-infected cells, J. immunol., 1986 Jan; 136(2):705-9

#### **Research Article**

# Herbo-Mineral Combination Drug as Matravasti in Katigraha w.s.r to Inter Vertebral Disc Prolapse

- A NOVEL CONCEPT

# <sup>1</sup>Divya Bendi, <sup>2</sup>Perugu Srikanthbabu

<sup>1</sup>Asstt. Professor, Dept of Kayachikitsa, Dr.NRS Govt. Ayurvedic college Vijayawada, A.P <sup>2</sup>Professor & HOD, Dept of Kayachikitsa, Dr. BRKR Govt. Ayurvedic College, Hyderabad, Telangana.

Manuscript Received on 04/10/2019

Reviewed on 07/11/2019

Accepted on 27/11/2019

# Abstract

Low Back Pain is also said to be the price man had to pay for the evolution from four-legged animal to bipedal being. It was ranked as 1st in leading cause for disability according to Global Burden of Disease Study 2017. Hence, effort has been made to search for a safe, effective and economical medicine. Based on patho physiology Inter Vertebral Disc Prolapse can be correlated with katigraha. Katigraha is a condition, where kati ashrita saama vayu causes ruja and graha in kati. Katigraha was first explained by Shodhala in Gadanigraha. As Vasti is known as Ardha chikitsa we have chosen Matravasti for Katigraha. Herbo-mineral formulation by name Mahamasha tailam 60ml and 1gm Pravala Bhasma for matravasti and Rasona Ksheerapaka 50ml as oral medicine have been chosen for the study. By performing Vasti with Mahamasha tailam and Pravala Bhasma we can nourish the disc with guru property of Masha and snigdha of Tailam. Pravala is known as best asthi rasayana. According to Dalhana, "Yaeva kala Purishadharasaaeva Asthidharaiti" Based on this quote pureeshadhara as asthidhara kala, we assume this as GIT contain more Na-Ca channels in its Auerbach plexus of muscle layers. When we give pravala in rectal route, large amounts of ionic calcium will be absorbed and action potential will be generated in short time. This will increase intestinal muscle contractions. When contractions increase, absorption of nutrients also will be more. So dhatu kshaya can be corrected. Through deepana, pachanaguna of pravala we can correct aama in shorter duration. Hence we have added Pravala Bhasma in vasti. A prospective randomized open label clinical study was carried out at Dr. BRKR Govt Ayurvedic College, Hyderabad. Total 30 patients were taken from OPD and IPD. They were treated for 72 days in three spells with an interval of 16 days. At the end of the treatment significant change was observed in subjective and objective parameters.

**Keywords:** Calcium Sodium channels, IVDP, *Katigraha, Matra Vasti, Maha Masha Tailam, Pravala Bhasma, Rasona ksheerapaka*.

# Introduction

Lumbago is also said to be the price man had to pay for the evolution from four-legged animal to bipedal being. It was ranked as 1<sup>st</sup> in leading cause for disability according to Global Burden of Disease Study 2017<sup>1</sup>. Busy and sedentary life style, improper sitting postures, lack of exercise, human spine is exposed to new patterns of force in distribution of weight and muscular tensions lead to herniation of nucleus pulposus and pressure at Posterior Longitudinal ligament, and finally Inter Vertebral Disc gets prolapsed.

In *Kati graha patho-physiology agnimandya* leads to *aama* which causes *margavarodha* and *vatavriddhi* which finally results in *asthikshaya* (degeneration of vertebrae). According to *Charaka* in *asthikshaya*, *lakshanas* like *asthishoola*, *asthibheda*, *sandhi saithilya* and *medakshaya* were observed. Here in IVDP also degeneration of vertebrae which leads to pain, fragility of vertebrae and prolapse can be observed. Symptomatically it can be appraised as pain and stiffness in the initial stage and paraesthesia in later stages. Based on the patho physiology, Inter Vertebral Disc Prolapse is contributing majority to *Kati Graha*. In Ayurveda, *Katigraha* was first explained by *Acharya Shodala* in *Gada Nigraha*. *Kati Graha* is a condition, caused by *kati ashrita saamavayu*, which causes *ruja* and *graha*<sup>2</sup> in *kati pradesha*. *Sarangadhara* mentioned it as one of the *Vataja nanatmaja vyadhi*.

In spite of the high rates of prevalence all over the world and tremendous advancement of modern system of medicine i.e. NSAIDs, physiotherapy, and surgical intervention, an ideal treatment is not yet available for IVDP. Because of the varied etiological factors, greater recovery period, disc prolapse has become clinician's challenge. The treatments are conflicting and often un-rewarding as well. Treatments in modern medicine provide

**Corresponding Author:** Dr. Divya Bendi, Asst. Professor, Dept of Kayachikitsa, Dr.NRS Govt. Ayurvedic college Vijayawada, A.P, India, Email: drdivyabendi@gmail.com

**How to cite this article:** Bendi D, Srikanthbabu P: Herbo-mineral combination drug as Matravasti in Katigraha w.s.r to Inter Vertebral Disc Prolapse - A novel concept; AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 11 - 18

symptomatic relief and are also very expensive. Symptomatic management of a disease with muscle relaxants is always incomplete and cannot break up the chain of pathogenesis. Surgical intervention is not well accepted by the society as it involves some surgical risks. So, effort has been made here to search for a safe & effective medicine, without any side effect.

Keeping this in view to combat the disease, Herbo mineral formulation by name *Mahamasha tailam* and *Pravala Bhasma* for *matravasti* and *Rasona Ksheerapaka* as oral medicine have been chosen for the study. All ingredients in the formulation exclusively possess *vata* pacifying property.

# **Material and Methods**

30 patients were randomly selected from OPD and IPD of Dr. BRKR Govt. Ayurvedic College and hospital, Hyderabad.

- 1. Inclusion criteria
  - Patients of the age of 25 50 years.
  - Patients who are with signs and symptoms of katigraha.
  - · Patients who were already diagnosed as IVDP.
  - Patients who are willing to participate in the study.

## 2. Exclusion criteria

- Patients who are below the age of 25 and above the age of 50 years.
- · Patients in whom vasti is contraindicated.
- History of hypersensitivity to the trial drug or any of its ingredients.
- Patients having infective conditions of spine like T.B of spine etc.
- Patients having Ankylosing spondylitis and Spina bifida.
- · Patients having abhighata janya katigraha.
- · Patients with evidence of malignancy.
- Patients with uncontrolled Diabetes Mellitus, unstable cardio vascular disease.
- Patients suffering from other major systemic illness necessitating long term treatment.
- · Patients with concurrent serious Hepatic disorder or Renal disorder.
- · Smokers/ alcoholics/drug abusers.
- · Pregnant or lactating women

# Methodology

1. Study type Interventional 2. Purpose Treatment 3. Masking Open label 4. Control Uncontrolled Prospective 5. Timing 6. End point Efficacy 7. No.of groups: One 8. Sample size: 30.

# Treatment plan:



Follow up: 25th, 49th, 73rd days

# **INVESTIGATIONS:**

- Complete Blood Picture
- Fasting Blood Sugar. Post Prandial Blood Sugar HbA1C
- Liver Function Test
- Renal Function Test
- · X-Ray of lumbo sacral Spine Postero-Anterior view and Lateral view
- CT Scan /MRI of Lumbar Spine.

# Methods of assessment of treatment:

- The effect of the therapy was assessed pertaining to improvement recorded in clinical findings.
- Changes observed in signs and symptoms were assessed by adopting suitable scoring methods and objective signs by using appropriate clinical tools.
- Both subjective and objective assessments were done in all patients before and after follow up period.

# **Assessment Parameters with Grading:**

# Table No. I - Grading according to severity of Symptoms

Grade	Symptom
0	Complete relief or no symptom
1	Presence of mild symptoms
2	Presence of moderate symptoms
3	Presence of severe symptoms

# **Table No. II - Grading of Subjective Parameters**

S.No	PARAMETER	Grade-0	Grade-1	Grade-II	Grade-III
1	Pain on rest (Lying)	No complaint	Reveals on enquiry	Complaints frequently when moves joints	continuous pain
2	Pain on rest (On sitting):	No complaint	Reveals on enquiry	Complaints frequently when moves joints	continuous pain
3	Pain on movement (on walk) on plain surface	Can able to walk more than 10 mtrs	Can able to walk only Up to 5 mtrs	Can able to walk only Up to 3 mtrs	Cannot walk due to pain

Table No. III - Objective Parameters with Grading

S.No	PARAMETER	Grade-0	Grade-1	Grade-II	Grade-III	
1	Tenderness	No tenderness	Pain on touch and winces	Withdraws the part	Not allow to touch the part	
2	Lumbar flexion	Able to touch the ground	able to go upto ankle	able to go just below knee	Not upto knee	
3	Lumbar extension	Able to do without difficulty	Able to do with some difficulty	Able to do upto shoulder	Can not able to do	
4	Left Lateral Movement	Able to move hand below knee without difficulty	Able to move hand below knee with pain	Can not go below knee	No movement	
5	Right Lateral Movement	Able to move hand below knee without difficulty	Able to move hand below knee with pain	Cannot go below knee	No movement	
6	Rotation	Can rotate easily	Rotate with difficulty	Rotate to one side	Cannot rotate	
7	Walking Time (Taken for20 steps)	Upto 30 sec	31-40 sec	41-50 sec	51-60 sec	
8	Straight Leg Raising Test	Negative	>70º	35 º -70 º	0-35 º	
9	VAS Score	0 (No Pain)	1-3 (Mild)	4-7 (Moderate)	8-10 (Severe)	
10	10 Sugar Baker & 24 Barofsy score		15-23	8-15	0-8	
11	Groopough &		50-64	30-49	0-29	

# Table No. IV - Drug Intervention

Drug intervention:

SI.No	Name of the Drug	Route of administration	Time of administration	Dose	Period
1	Maha Masha Tailam and Pravala bhasma	Rectal	After Food	60ml and 1gm	72 days in three spells with interval of 16 days
2	Rasona Ksheerapaka	Oral	Empty stomach	50ml	72 days in three spells with interval of 16 days

Table No. V - Rasa Panchaka of Pravala Bhasma<sup>3</sup>

S.No	Sanskrit Name	English name	Latin/ Chemical Name	Rasa	Guna	Veerya	Vipaka	Karma
1	Suddha Pravala	Coral	Anthozoa polypus / Calcium oxide	Madhura, Amla	Lahu, Snigdha	Madhura	Sheeta	Kapha, vatasaamaka, Asthiposhaka

# Rasona Ksheera Paka:

Rasona:

It is also called as Lasuna.

Table No. VI - Rasa Panchaka of Lasuna

S. No	Sanskrit Name	Latin Name	Family	Part Used	Rasa	Guna	Veerya	Vipaka	Karma
1	Lasuna	Allium sativum	Liliaceae	Bulb	Madhura, Lavana, Katu, Tikta, Kashaya	Guru, pichila Snigdha	Ushna	Katu	Asthimamsasandhanakara

#### KSHEERA:

Ksheera is the best dravya among Jeevaniya dravyas. As 10 gunas of milk and ojas are similar, milk acts as ojovriddhikara5. Table No.VII Rasa Panchaka of Ksheera<sup>5</sup>

S.No	Sanskrit Name	Rasa	Guna	Virya	Vipaka	Karma
1	Ksheera	Madhura	Swadu, shita, Mridu, Snigdha, Gurupichila, Bahala, Slakshna	Sheeta	Madhura	Jivaniya, Manasakar, Rasayana

# KSHEERA PAKA:

Medicated milk which is prepared by boiling the milk with drug and water until only milk remains is called *Ksheerapaka*. The ratio of drug, milk and water differ according to different *Acharyas*.

Table No. VIII - Ratio of Milk, Water and Drug according to different acharyas

S.No	Authors	Drug	Milk	Water	Reduction		
1	Charaka <sup>6</sup> and Chakrapani <sup>7</sup>	1 part	8 part	8Part	Till milk part remains		
2	Sharangadhara <sup>8</sup>	1 part	8 part	32 part	Till milk part remains		
3	YadavjiTrikamji <sup>9</sup>	1 Part	15 part	15 part	Till milk part remains		
4	Vagbhata <sup>10</sup>	Kashaya should be prepared as per the procedure, after that, equal quantity of milk is to be added, again re-heated till only milk part remains.					

Rasona ksheerapaka was mentioned in Charaka chikitsa, Gulma chikitsa adhyaya. Ksheerapaka vidhi is advised for the drugs which are teekshna, ushna, and kashaya rasa. As Rasona is vatahara, Asthisandhanakara, and ksheera is asthiposhaka and jeevaniya, rasona ksheerapaka was used orally in the management of Katigraha. We have followed the method said by Charaka and Chakrapani. 6gms of lashuna, 50ml of water and 50ml of milk were

taken. Total drug reduced to 50ml and ksheerapaka was attained.









Pravala

Pravala bhasma

Rasona

Ksheera

# **Results**

- The assessment of the overall effect of the treatment revealed that 13.3% of the patients recorded excellent result while 83.3% of the patients showed good and 3.33% of the patients had moderate result.
- Most of these are found to be statistically highly significant as per the Wilcoxon signed rank test (subjective parameters) and Paired 't' test (objective parameters).

Table No. IX Overall assessment of results

RESULT	NO OF PATIENTS	PERCENTAGE
Excellent	4	13.33
Good	25	83.33
Moderate	1	3.33
Mild	0	0

# Statistical analysis:

Table No.X Statistical analysis of subjective parameters

S.No	Lakshana	Ме	ean	% of relief	9 9 1		S.E		t-value	p-value	s
		BT	AT		BT	AT	BT	AT			
1	Pain on Sitting	2.33	0.67	71.24	0.48	0.61	0.09	0.11	15.05	<0.0001	HS
2	Pain on Lying	2.30	0.56	75.65	0.47	0.57	0.09	0.10	21.079	<0.0001	HS
3	Pain on Walking	2.33	0.70	69.95	0.48	0.53	0.07	0.09	18.25	<0.0001	HS
4	Graha	2.30	0.77	66.52	0.47	0.57	0.08	0.09	14.69	<0.0001	HS

Table No. XI - Statistical analysis of objective parameters

S.No	Parameters	Ме	an	%of relief	S.	D	S.E		t-value	p-value	s
		BT	AT		ВТ	AT	ВТ	AT			
1	Tenderness	2.10	0.10	95.23	0.31	0.34	0.06	0.08	16.57	<0.0001	HS
2	Lumbar Flexion	2.40	0.83	65.41	0.50	0.53	0.09	0.10	17.02	<0.0001	HS
3	Lumbar Extension	2.53	1.43	43.47	0.51	0.50	0.09	0.11	12.53	<0.0001	HS
4	Left Lateral Movement	2.13	0.87	59.15	0.35	0.68	0.06	0.12	13.32	<0.0001	HS
5	Right Lateral Movement	2.03	0.50	75.36	0.18	0.51	0.03	0.09	14.69	<0.0001	HS
6	Lumbar Rotation	1.93	0.57	70.46	0.58	0.50	0.11	0.09	12.17	<0.0001	HS
7	Walking Time	2.33	0.70	69.95	0.48	0.53	0.07	0.09	18.25	<0.0001	HS
8	SLR	2.60	0.90	65.38	0.50	0.66	0.09	0.12	13.25	<0.0001	HS
9	VAS	2.43	1.23	49.38	0.50	0.63	0.09	0.11	11.93	<0.0001	HS
10	Sugar Baker &Barofsy	2.40	1.17	51.25	0.50	0.53	0.09	0.10	13.43	<0.0001	HS
11	Greenough& Fraser score	2.33	0.93	60.08	0.61	0.64	0.11	0.12	12.33	<0.0001	HS

# **Discussion:**

**Age:** The highest percentage of age group seen is 25-35 years and next is 45-50 years. This is because of irregular postures and habits of younger people and degenerative changes of disc in the 45-50 age group.

**Occupation:** It reveals that prolonged sitting and standing postures, increased mental activity, and stress and strain contribute in establishment of *Kati Graha*. House wives also are more prone to this disease because of irregular house hold work, sitting and standing postures.

**Diet:** Mixed diet contains *katu*, *tikta*, *kashaya rasa aahara*, *sushka mamsa* and *viruddha aahara* which can aggravate vata and contribute one of the *Vataroga* i.e *Kati Graha*.

**Prakruti:** From this data it can be inferred that *Vata Pitta and Vata Kaphaprakruti* people are more prone to *Kati Graha* as they are main *doshas* involved in *Kati Graha*.

**Nidana:** When it comes to *nidana*, 20 (66.66%) patients were having habit of *katu*, *tikta Kashaya rasa sevana*, 21 (70%) patients were having *dukha sayyasana*, 19(63.33%) were having *vega dharana*, 8 (26.66%) were having *ratri jagarana*, 16 (53.33%) were having atikrodha/ atichinta (stress), 4(13.33%) were having uchair bhashana and Ati adhva. This indicates that the present life style and dietary habits contribute more to *vatavyadhis* like *Kati Graha*.

**Disc level:** L4-L5 and L5-S1 discs are most commonly affected. This is because of weak reinforcement of posterior fibres by posterior longitudinal ligament. Especially L4-5 and L5-S1 where it is midline, narrow, unimportant structure attached to annulus.

**Area of lesion:** Posterior region is most common area of lesion. This is because of incomplete annular lamellae in this quadrant i.e. each lamellae end with fusion to an adjacent lamellae not completely circular. Fibres of annulus were deficient posteriorly.

# Probable Mode of Action of Vasti procedure and Drugs Maha Masha Tailam, Pravala Bhasma, Rasona Ksheerapaka in Kati Graha:

# On the basis of karma:

Asthi poshaka and asthi sandhana karma of pravala and lasuna and brimhana of mashamay doasthi dhatu poshana and modify degenerative changes of asthi.

#### On the basis of Dosha Karma:

All the ingredients were having *vata hara* property. Because of that *vyana* and *apanavata* were regulated. Symptoms like *ruja*, *graha* were relieved.

## Pravala Bhasma:

*Pravala* is a by-product of tiny sea anemone that takes in minerals from the ocean and deposit coral on the ocean floors. Because of this digestion process, coral calcium is unique in its ionic, easily absorbable form.

## Mode of action of vasti:

Vasti is known as Ardha chikitsa for Vatic disorders. It acts in the body in four dimensions i.e. by its action of unique Procedure, Drug, Sroto Sodhana and regulation of GUTBRAIN¹¹ system. If we administer Pravala orally, smaller amount will be absorbed than through rectal route. According to pharmaco kinetics, drug given in rectal route will have faster absorption and higher bio availability. Pravala is known as best asthirasayana. Based on Dalhana's quote pureeshadhara as asthidhara kala, we assume this as GIT contain more Na-Ca channels in its Auerbach plexus of muscle layers. When we give pravala in rectal route, large amounts of ionic calcium will be absorbed and action potential will be generated in short time. This will increase intestinal muscle contractions. When contractions increase, absorption of nutrients also will be more. So dhatu kshaya can be corrected. Through deepana, pachanaguna of pravala we can correct aama in shorter duration. Based on Asrayaasrayi Sambandha by increasing Asthi dhatu it alleviates Vatadosha. Thus, vasti acts as prospective treatment in management of IVDP.

# Conclusion

On completion of this study, conclusion drawn on the basis of deductive reasoning of data obtained from this clinical trial is as follows:

Kati Graha is considered as Inter Vertebral Disc Prolapse in the present study. It is a degenerative disorder involving dhatu kshaya and margavarodha. This is more prevalent in 25-30 years and 45-50 years aged Vata pitta and Vata Kapha prakrati persons and in people having mixed diet. Ratri Jagarana and Atiadhva were observed in most of the cases. Katu ruksha ahara sevana is one of the common aharaja nidana was observed in 66% of the cases. Vega dharana and Dukh as ayyasana are common viharaja nidana observed in 60% of cases. In the present trial, Matravasti with Maha Masha Tailam, Pravala Bhasma and oral Rasona Ksheerapaka were found effective. The present study by Matravasti with Maha Masha Tailam and Pravala bhasma and oral rasona ksheerapaka showed extremely statistically significant results (P<0.0001) in Ruja, Graha and Heena Gamana Shakti. No major hazardous side effects were noticed during the present study. So, it can be said to be a safer drug. As the sample size is too small (30) need of study with a bigger sample size to draw a valid conclusion.

# References:

- healthdata.org(home page on internet) available form <a href="http://www.healthdata.org/sites/default/files/files/policy\_report/2019/GBD\_2017\_Booklet.pdf">http://www.healthdata.org/sites/default/files/files/policy\_report/2019/GBD\_2017\_Booklet.pdf</a>
- 2. Vaidya Shodala; Gada Nigraha; By Indradeva Tripati; Edited by Ganga Sahay Pandey; Part
- 3. 3rd Edition 99; Varanasi; Chaukambha Sanskrit Sansthan; Pp;871; Page No 508. 3. Rasatarangini 23/129.
- 4. Susrutha Samhita sutrasthana 46/244 by Priyavat Sharma, Chowkambha orientals.
- 5. CharakaSamhitha Sutra sthana 27/217 by Vaidya Bhagavandas, Chowkambha Sanskrit orientals, 2007 reprint.
- 6. K. Shasthri. Charaka Samhita of Charaka. GangasahayaPandeya (ed). Reprint ed. Varanasi: Chaukhamba Sanskrit Sansthan; 2009 vol 2. p.175.
- 7. Tripathi Jagadishvaraprasad. Chakradatta of Chakrapanidatta, Mishra Brahmashankara (ed). 5th ed. Varanasi: Chowkhamba Sanskrit Series; 1983. p. 260.
- 8. ShastriVidyasagar Parasurama. Sharangdharasamhita of Sharangdhara. 7thed. Varanasi: Chaukhamba Orientalia; 2008. p.167.
- 9. Yadavjitrikamji Acharya. DravyagunaVignana, 2nd ed. Satyabhamabaipanduranganirnayasagaramudranayantralaya; 1947. Uttarardha paribhasha khanda. p. 33-34.
- 10. Jyotirmitra Kaviraja. Astanga Sangraha of Vagbhata Vridda, Dr. Shivaprasad Sharma 2nd ed. Varanasi: Chowkhamba Sanskrit Series; 2008. p. 617.
- 11. Wood, J.D: Physiology of enteric nervous system, In Johnson L.R, ed Physiology of gastrointestinal system, Vol.1, New York raves Press, 1981, pp-1-37.

# **Case Report**

# Effectiveness of Ayurveda modalities in the management of Gridhrasi (Prolapsed Inter Vertebral Disc): A Case Study

# <sup>1</sup>Ritu Kumari, <sup>2</sup>N.R.Singh, <sup>3</sup>Jitendra Varsakiya

<sup>1</sup>MD Scholar, PG Dept. of Kayachikitsa, Ch. Brahm Prakash Ayurveda Charak Sansthan, Khera Dabar, Najafgarh, New Delhi

<sup>2</sup>Professor and Head, PG Dept. of Kayachikitsa, Ch. Brahm Prakash Ayurveda Charak Sansthan, Khera Dabar, Najafgarh, New Delhi

<sup>3</sup>Assistant Professor, PG Dept. of Kayachikitsa, Ch. Brahm Prakash Ayurveda Charak Sansthan, Khera Dabar, Najafgarh, New Delhi .

Manuscript Received on 26/09/2019

Reviewed on 09/10/2019

Accepted on 04/11/2019

# **Abstract**

Due to globalization as well as urbanization people have sedentary lifestyle which is resulting in the problem of lower backache. The main cause of this problem is prolapsed intervertebral disc at the level of L4-L5 vertebrae. Protrusion of gelatinous nucleus pulposus through the rend of fibrous annulus fibrosus is called prolapsed intervertebral disc. This protruded gelatinous material compresses the roots of spinal nerves and this irritation produces the symptoms like lower backache radiating or non-radiating to unilateral or bilateral lower limbs. In Ayurveda, this condition resembles Gridhrasi. In the contemporary system of medicine NSAIDs and intradermal corticosteroid injection is the main treatment modality for pain management which causes a number of side effects. Aim: To evaluate the effectiveness of Ayurveda modalities in the management of *Gridhrasi*. Material and method: A case of PIVD (Prolapsed Inter Vertebral Disc) was treated with *Panchakarma* procedures and Ayurvedic oral drugs which have given satisfactory results with no side effects. The *Panchakarma* protocol designed for this particular patient was *Sarvanga Abhyanga* ( therapeutic massage) with *Laghu Vishagarbha Taila*, *Sarvanga Swedana* ( sudation therapy) with *Dashamoola Kwatha*, along with oral medication as *Trayodashanga Guggulu* 3 gm daily in three divided doses and *Eranda Taila* 10 ml twice a day with luke warm water for 14 days.

**Result**: Visual Analogue Scale, Oswestry Disability Index and SLRT were considered as parameters of assessment. There was sustained satisfactory improvement in patient's condition over 2 months of follow-up. **Conclusion**: This case study shows that PIVD can be successfully managed by Ayurvedic treatment modality. **Keywords-** Ayurveda, *Gridhrasi*, lower backache, prolapsed intervertebral disc

# Introduction

World wide up to 80% of people experience lower back pain at some point in time in their life, and according to Kelsey and White, an even larger proportion have found to have degenerative spine disease at autopsy. Chronic back pain which is defined as pain lasting more than 12 weeks has a prevalence of 5.9% to 18.1%. Recurrent meningeal or sinu-vertebral nerves are the branches of spinal nerve which supply pain fibers to the intraspinal ligaments, periosteum of bone, the outer layer of annulus fibrosus and the capsule of the articular facets. Although spinal cord itself is insensitive, but so many conditions produce pain by involving the adjacent structures. Among all the signs in detecting nerve root compression, straight leg raising and Laseague's sign is the most useful. The fully developed syndrome of the common prolapsed intervertebral lower lumbar disc is i.) pain in buttock radiating to bilateral or unilateral lower limb, ii.) a stiff or unnatural spinal posture, iii.) some combination of paraesthesia, weakness and reflex impairment. Pain management and resume of activity is the main aim in the management of lower lumbar disc disease. Pain management is done in many ways i.e. traction, oral analgesics as NSAIDs, uploads and corticosteroids. Which can result in lots of complications including rare disease i.e. fungal meningitis. These managements Provide short term improvement and the need for surgery is always there. Variants of hemilaminectomy with excision of the disc fragment are the most widely used surgical procedures for lumbar disc diseases . These uncertainties in the management of this disease made us think to explore some ayurvedic ways for the management of this disease

**Corresponding Author:** Dr. Ritu Kumari, MD Scholar, PG Dept. of Kayachikitsa, Ch. Brahm Prakash Ayurveda Charak Sansthan, Khera Dabar, Najafgarh, New Delhi, India, Email: <a href="mailto:ritu6285@gmail.com">ritu6285@gmail.com</a>

**How to cite this article:** Kumari R, Singh NR, Varsakiya J: *Effectiveness of Ayurveda modalities in the Management of Gridhrasi (Prolapsed Inter Vertebral Disc): A Case Study;* **AYUHOM**: Vol. 6, Issue 1 (Jan – June, 2019); 19 - 23

In Ayurveda this disease condition is related to *Gridhrasi* which is described under the umbrella of *Vatavyadhi*. Etiological factors for all the *Vatavyadhis* are common which includes inappropriate and excessive exercises, swimming, jumping, excessive walking, trauma, falling from vehicles. Symptoms of this disease are pain, stiffness, pricking or pulsatile sensation starting from the gluteal region then radiating to the waist, back, thigh, lower leg and foot. Snehana (Ayurvedic therapeutic massage) and *Swedana* (sudation therapy) is first general treatment modality of all the *Vatavyadhi*. Administration of *Mriduvirechak Dravya* (mild purgative drugs) is also advised in the form of *Eranda taila* (Castor oil) in the management of *Vatavyadhi*.

# **Material and Method:**

# **Case Report**

A 23-year-old Hindu, male patient visited CBPACS Kayachikitsa OPD on 13th March 2019 having central I.D. no. 28993 with the complaint of severe pain in waist radiating to bilateral lower limbs. The pain was severe agonizing type and he was unable to walk himself. The patient had pain for one year. Initially had mild pain but gradually pain increased and affected his daily activity. For one month he was unable to walk without support and had severe pain. For one year had consulted physicians who asked for intradermal injections and physiotherapy. But the condition gets worse. So, the patient visited here for better management for his ailment.

# Aims and objectives:

To evaluate the efficacy of Ayurveda remedies in the management of Gridhrasi (Prolapsed Inter-Vertebral Disc.)

# Case findings

The patient was admitted to the I.P.D. ward of *Kayachikitsa* with I.P.D. registration no. 18319. The general condition of the patient was not good. He was anxious, appetite was moderate, and tongue was coated. His blood pressure was 110/70 mm Hg. The pulse rate was 78 per minute and was full in volume and regular. The range of movement of bilateral lower limbs were restricted and painful. There was tenderness in the lumbar area. There was no difficulty in defecation and micturition. No numbness was present in the lower extremity. There was no history of any fall or trauma. During physical examination, SLR (Straight Leg Raising) Test was done which was 0 degree on the left and 50 degrees on the right. On muscle examination, there was coordination of upper and lower limbs. Muscle bulk, Power, Tone was normal. Deep tendon reflexes and superficial reflexes were normal. The patient was working as B.P.O. employee doing night shifts. His ESR was raised. He was vegetarian. *Dasha Vidha Pariksha* was done for patient assessment. The patient has *Kapha Pittaj Prakriti, Vikriti – Vata Kaphaj, Sama Pramana, Madhyama Sara*, with *Avara Vyayam Shakti*, and *Avara Bala*.

Table no. 1: Time line of events:

Duration	Particular and interventions
March 2018	First episode of lower backache
March 2018 - May 2018	Managed with allopathic treatment
December 2018	Lower backache increased gradually
January 2019 – February 2019	Severity of pain increases
11 <sup>th</sup> March 2019	Unable to get up and stand by his own
13 <sup>th</sup> March 2019	Gets admitted in hospital
14 <sup>th</sup> – 28 <sup>th</sup> March 2019	Treatment with Sarvanga Abhyanga with Laghu Vishagarbha Taila, Sarvanga Swedana with Dashmoola Kwatha, Trayodashanga Guggulu 3 gm in 3 divided doses and Eranda Taila 10 ml BD with luke warm water
5 <sup>th</sup> April 2019	First Follow up
13 <sup>th</sup> April 2019	Second follow up
20 <sup>th</sup> April 2019	Third follow up
27 <sup>th</sup> April 2019	Fourth follow up

# Diagnostic focus and assessment

The diagnosis was done on the basis of symptoms and MRI findings. In Ayurveda this condition resembles *Gridhrasi*. To reconfirm the diagnosis of prolapsed intervertebral disc, MRI was done. In MRI there was diffuse disc bulge at L3-L4 and L4-L5. And there were no signs of sacroillitis. Thus ankylosing spondylitis was excluded. In MRI there were no signs of any fracture, benign or malignant tumors, any degenerative diseases, any anatomical or congenital deformities. So, these things were excluded from the differential diagnosis.

## Treatment Plan

During the course of treatment *Panchakarma* procedures were done in form of *Sarvanga Abhyanga* (therapeutic massage) with *Laghu Vishagarbha Taila* and *Sarvanga Vashpa Swedana* (sudation therapy) with Dashamoola Kwatha, Trayodashanga Guggulu 2 gm per day in three divided doses with lukewarm water after meal, and *Eranda Taila* 10 ml BD with lukewarm water after meal for 14 days. (Table No.1)

Table no. 2: Details of procedures used during treatment:

No.	Name of procedure	Drug used	Method	No. of days
1.	Sarvanga Abhyanga (therapeutic massage)	Laghu Vishagarbha Tail	Gentle massage with Luke warm oil was given for 20 minutes per day.	14 days
2	Sarvanga Vashpa Swedana (sudation therapy)	Dasha moola Kwath	Sudation therapy was given with the steam of Dashmoola Kwath for 10 minutes or as long as the patient feels comfortable.	14 days

# Table no. 3 : Posology of Drugs used in the study:

No.	Name of drug	Dose	Anupan	Duration
1.	Trayodashanga Guggulu	2 gm daily in three divided doses ( 4 tablets each of 500mg)	Lukewarm water	14 days
2.	Eranda Taila (castor oil)	10 ml twice a day morning and night empty stomach	Lukewarm water	14 days

Table No. 4: Qualities of ingredients of Trayodashnag Guggulu

SI. no.	Name	Botanical name	Rasa	Guna	Virya	Vipaka	Karma
1	Abha	Acacia arabica Wild.	Kashaya	Guru, Ruksha	Sheeta	Katu	Kaphapitta Shamaka, Snehana and Kaphagna
2	Ashwagandha	Withania somnifera Linn.	Kashaya, Tikta	Laghu, Snigdha	Usna	Madhura	Kaphavatashamaka, Balya, Rasayana, Vednashamaka and Shotha hara
3	Hapusa	Juniperus communis Linn.	Katu,Tikta, Kashaya	Guru, Ruksha, Teekshna	Usna	Katu	Kaphavatashamaka, Shoolahara and Naditantrottejaka
4	Guduchi	Tinospora cordifolia Wild.	Katu,Tikta, Kashaya	Laghu	Ushna	Madhura	Tridoshaghna, Balya, Rasayan
5	Shatavari	Asparagus racemosus Wild.	Madhura, Tikta	Guru, Snigdha	Sheeta	Madhura	Vatapittashamaka, shotha hara, Nadi bala prada
6	Gokshru	Tribulus terrestris Linn.	Madhura	Guru, Snigdha	Sheeta	Madhura	Vatapittashamaka, Vedanasthapaka, Vatashamaka Anulomana, Shothahara
7	Vriddhadaru	Argyreia speciosa Sweet.	Katu, Tikta, Kashaya	Laghu Snigdha	Ushna	Madhura	Kaphavata Shamaka, Anulomana, Rasayana Nadi Balya
8	Rasna	Pluchea lanceolate Oliver & Hiem	Tikta	Guru	Usna	Katu	Kaphavatashamaka Vedanasthapana, Vataghna Vedanashamaka

9	Shatahva	Anethum sowa Kurz.	Katu, Tikta	Laghu, Teekshna	Ushna	Katu	Kaphavatashamaka, Vatanulomaka, Vedana sthapaka
10	Shati	Hedychium spicatum Buch- Ham	Katu, Tikta, Kashaya	Laghu, Teekshna	Anushna	Katu	Kaphavatashamaka
11	Yavani	Trachyspermum ammi Linn.	Katu, Tikta	Laghu, Teekshna	Usna	Katu	Kaphavatashamaka, Shoolprashmana Vatanulomana
12	Nagara	Zingiber officinale Roxb.	Katu	Laghu, Snigdha	Usna	Madhura	Kaphavatashamaka, Vatanulomana, Aamdosha hara
13	Kaushika	Commiphora mukul (Hook ex stocks)	Tikta, Katu, Kashaya	Ruksha,	Ushna	Katu	Tridoshaghna, Vedanasthapana Shothhara, nervine tonic
14	Goghrita		Madhura, Katu, Tikta	Snigdha , Sheeta	Sheeta	Madhura	Tridoshaghna, Agnivardhaka,

#### **Outcome Measure And Follow Up**

For pain assessment, VAS score and Oswestry Disability Score was done. Weekly assessment of pain and SLR was done. There was a remarkable improvement in his condition and after two weeks of hospital stay his SLR test was 90 degrees in both lower limbs and mild pain in the lower back only after walking or climbing stairs. His condition was satisfactorily stable during the follow up period of two months.

Table No. 5: Assessments obtained before and after treatment:

DATE	VAS Score	ODI Score	SLRT	
13 <sup>th</sup> March 2019(before treatment)	10	62%	Left – 0, Right -50	
20 <sup>th</sup> March 2019	7	54%	Left – 40, Right -70	
28 <sup>th</sup> March 2019	3	32%	Left – 90, Right -90	
5 <sup>th</sup> April 2019 (first follow up)	3	32%	Left – 90, Right -90	
VAS - Visual analog score, ODI Score - Oswestry Disability Index, SLRT - Straight Leg Raising Test				

# **Discussion**

Abhyanga (Therapeutic massage) is used to produce relaxation and strength to the muscles. In this process after applying medicated oil, gentle message of body is done. Laghuvishagarbha Taila is used in all the Vata Vyadhi. Swedana is very effective treatment for Vata Kapha Dosha. This procedure eliminates stiffness, coldness, heaviness, and promotes sweating. Dashmoola has Shothahara properties. In a study it has been concluded castor oil in dose of 0.9 ml TDS is efficacious in management of osteoarthritic pain with no adverse effects. Ricinoleic acid, the main component of castor oil, has been studied extensively in an experimental model for its anti-inflammatory action due to its inhibitory action on phospholipase A2, leukotriene B4. It alters mucous membrane permeability, stimulates adenyl cyclase-cyclic AMP system, .A study showed chemical capsaicinand acetic acid-induced and thermal pain were reduced by ricinoleic acid to an extent similar to capsaicin. Effect was may be due to the peripheral depletion of substance P in inflamed paws indicating a probable reduction of substance P in the periphery or that it is acting through VR1 (Vanilloid receptor 1) which is a critical mediator of thermal nociception following topical application. Ricinoleic Acid is chemically similar to prostaglandinE1. In Ayurveda it has been mentioned that Erand Tail is Vata Kapha Dosha Hara and it pacifies the perturbed Dosha in lower part of body and can used in management of Gridhrasi. Sopha (edema) is always associated with Vata and Kapha Dosha because there is mentioned in Ayurveda classis two type of Ghridhrasi one is Vata second one is Vata-Kaphaja.

**Trayodashang Guggulu** can be drug of choice in *Gridhrasi* which is named as because it contains thirteen ingredients. Most of the ingredients of *Trayodashang Guggulu* having the property of *Vatakaphahara* and *Vatanuloman*. Most of ingredient of *Trayodashang Guggulu* is *Rasayan* and *Balya* in its nature strengthen nerves. Thus, it is not only helpful in pain management but also helps in progress of the symptoms of this.

# **Conclusion:**

In modern era due to life style changes people are not focusing on health and resulting the ill condition. Due to wrong postures, improper exercises, disturbed sleep pattern, hours of commute, problems like Gridhrasi is very common. Conservative management by contemporary medicines have lots of side effects and those are not cost effective as well. Traditional system of medicine definitely effective in this type of condition without any side effect.

#### References:

- 1. Ropper AH, Samuels MA, Klein JP. chapter 11. In: Principles of neurology. 10th ed. china: Mc Graw Hill; p. 198.
- 2. Hooten WM, Cohen SP. Evaluation and Treatment of Low Back Pain. Mayo Clinic Proceedings. 2015;90(12):1699–718.
- 3. Ropper AH, Samuels MA, Klein JP. chapter 11. In: Principles of neurology. 10th ed. china: Mc Graw Hill; p. 209.
- 4. Shastri K, chaturvedi G editor. Charak Samhita of Agnivesh, chikitsa sthan chapter 2,. Varanasi, U.P.: Chaukhambha bharati academy; 2012. p. 779.
- 5. Shastri K, chaturvedi G editor. Charak Samhita of Agnivesh, chikitsa sthan chapter 28, Varanasi, U.P.: Chaukhambha bharati academy; 2012. p. 787.
- 6. Shastri K, chaturvedi G editor. Charak Samhita of Agnivesh, chikitsa sthan chapter 28. Varanasi, U.P.: Chaukhambha bharati academy; 2012. p. 787.]
- 7. The Ayurvedic Formulary of India. In: The Ayurvedic Formulary of India. 2nd ed. Delhi: The controller of publications; 2003. p. 68.
- 8. Final Report on the Safety Assessment of Ricinus Communis (Castor) Seed Oil, Hydrogenated Castor Oil, Glyceryl Ricinoleate, Glyceryl Ricinoleate SE, Ricinoleic Acid, Potassium Ricinoleate, Sodium Ricinoleate, Zinc Ricinoleate, Cetyl Ricinoleate, Ethyl Ricinoleate, Glycol Ricinoleate, Isopropyl Ricinoleate, Methyl Ricinoleate, and Octyldodecyl Ricinoleate. International Journal of Toxicology. 2007;26(3\_suppl):31–77.
- 9. Scott J, Huskisson EC. Graphic representation of pain. Pain. 1976;2:175–184. doi: 10.1016/0304-3959(76)90113-5.
- 10. Fairbank JCT, Couper J, Davies JB, O'Brian JP. The Oswestry low back pain disability questionnaire. Physiotherapy. 1980:66:271–3.
- 11. Ropper AH, Samuels MA, Klein JP. chapter 11. In: Principles of neurology. 10th ed. china: Mc Graw Hill; p. 201
- 12. Medhi B, Kishore K, Singh U, Seth SD. Comparative clinical trial of castor oil and diclofenac sodium in patients with osteoarthritis. Phytotherapy Research. 2009;23(10):1469–73.
- 13. Gaginella TS, Haddad AC, Go VL, Phillips SF. 1977. Cytotoxicity of ricinoleic acid (castor oil) and other intestinal secretagogues on isolated intestinal epithelial cells. J Pharmacol Exp Ther 201: 259–266.
- 14. Effect of ricinoleic acid and other laxatives on net water flux and prostaglandin E release by the rat colon. J Pharm Pharmacol 31: 681–685.
- 15. Caterina MJ, Leffler A, Malmberg AB et al. 2000. Impaired nociception and pain sensation in mice lacking the capsaicin. receptor. Science 288: 306–313.
- 16. Racusen LC, Binder HJ. 1979. Ricinoleic acid stimulation of active anion secretion in colonic mucosa of the rat. J Clin Invest 63: 743–749.
- 17. Shastri K, chaturvedi G editor. Charak Samhita of Agnivesh, chikitsa sthan chapter 28. Varanasi, U.P.: Chaukhambha bharati academy; 2012. p. 787.
- 18. Shastri A Editor. Bhaisjya Ratnawali., Vatavyadhi prakaran. In: Varanasi, U.P.: Chaukhambha prakashan; p.526.

# **Case Report**

# Ayurvedic management of Sub-Acute Pyelonephritis: A case report

## **Alok Kumar**

Lecturer, Dept. of Shalya Tantra, NEIAH, Shillong

Manuscript Received on 21/08/2019

Reviewed on 01/10/2019

Accepted on 08/10/2019

# **Abstract**

In *Ayurvedic* text *Sushruta samhita* most of the urological disorders are included in *Mutraghata/mutrakricha* and *Prameha*. Texts have wide description of these diseases in detail. Pyelonephritis is the disease of kidney where infection spreads from below to upwards. It can presents with fever, vomiting, frequent burning urination, abdominal pain which radiate to flanks and back of affected sites. In *Ayurveda* the condition is similar to the *Mutraghata* and *Prameha*. This is a case report of a 29 yrs old male bodybuilder patient, who came with complaint of pain in bilateral flanks, frequent burning urination and low grade fever. Ayurvedic medicines were used for above condition. The results were very encouraging.

Keywords: Mutraghata, Pyelonephritis, Prameha

# Introduction

Urinary tract infections (UTIs) include cystitis and Pyelonephritis<sup>1</sup>. Acute Pyelonephritis is an infection of the upper urinary tract, specifically the renal parenchyma and renal pelvis. Acute Pyelonephritis is considered uncomplicated if the infection is caused by a typical pathogen in a patient with compromised immune system who has normal urinary tract anatomy and renal function. A complicated Pyelonephritis is said to be present when it occurs in a patient with a structural or functional abnormality of the genitourinary tract and it leads to increase in the risk of unsuccessful treatment.

Acute Pyelonephritis<sup>2</sup> (APN) is a well-known disease, described extensively in literature. Earliest citing of the condition dates back to ancient Egypt, highlighting the destruction of the kidney parenchyma and its outcomes. In spite of this long history, the nomenclature of APN is still controversial, and the semantic ambiguities can still cause confusion.

Women are approximately five times more likely than men to be hospitalized with this condition however they have less mortality rates than men attributable to Pyelonephritis<sup>3</sup>.

APN is subdivided into uncomplicated and complicated. Severity of PN cannot be assessed by clinical or laboratorial parameters alone, radiological imaging such as Ultrasound (USG) abdomen, Computed Tomography (CT) is required to know the nature, extent and severity of disease and for planning interventions.<sup>4</sup>

Clinical Features: The clinical spectrum of acute Pyelonephritis is wide, ranging from a mild illness to sepsis syndrome. Mild Pyelonephritis can present as low-grade fever with or without lower-back or costo-vertebral angle pain, whereas severe Pyelonephritis can manifest as high fever, rigors, nausea, vomiting, and flank and/ or loin pain. Symptoms are generally acute in onset, and symptoms of cystitis may not be present. Fever is a distinguishing feature between cystitis and Pyelonephritis. Symptoms suggestive of cystitis (dysuria, urinary bladder frequency and urgency, and suprapubic pain) may also be present. In diabetics a rare but dreadful complication might occur which is called as emphysematous Pyelonephritis. This is an acute and severe infection of the renal parenchyma and peri-renal tissue, which results in gas within the renal parenchyma, collecting system or perinephrictissue. It is a life threatening condition if not treated promptly. Xanthogranulomatous Pyelonephritis occurs in association with chronic urinary obstruction (usually by staghorn calculi), together with chronic infection. It leads to granulomatous inflammation and destruction of renal tissue.

The clinical presentation of the Pyelonephritis may be correlated with the Usnavaat and the Mutroksaad

Corresponding Author: Dr. Alok Kumar, Lecturer, Dept. of Shalya Tantra, College of Ayurveda, North Eastern Institute of Ayurveda & Homoeopathy (NEIAH), Shillong, Meghalaya-793018, India, Email: dr.alokv@gmail.com

**How to cite this article:** Kumar A; Ayurvedic Management of Sub-Acute Pyelonephritis: A case report; AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 24 - 26

(pitta and kaphjanya) described by Acharya Sushruta<sup>5</sup> in Sushruta samhita. The causative factor for Usnavaat described by Sushruta are excessive exercise, longtime walking in bright sunlight. Due to above region the Vata and Pitta dosha aggravates and produce the Usnavaat. In Mutrosaad the causative doshas are Pitta and Kapha. Achary Sushruta advocated the general treatment<sup>6</sup> of Mutraghata by the different Kashay, kalka, ghrita, dugdha, kshara,aashava, aristha, upnaah,sweda, and bastichikitsa. These different kalpanas (formulations) are prepared with the various Ashmarinashak (Lithotripsic) drugs. The medicines used in this case study are described by Acharya Sushruta for the management of different mutra-ashmari.

# **Case report**

A 29 yrs old male professional bodybuilder patients came to OPD with complaint of pain in B/L flank, frequent burning urination and low grade fever since 1month. Patient has history of taken steroid upto 3 months for body building 6 month ago. Details of steroid were not available. No history of Diabetes Mellitus, Hypertension or any other chronic ailments. Patient was diagnosed as APN had taken some allopathic medicine for 10 days but condition was not cured completely so he visited our hospital. Blood test reveals WBC counts-11x103, Hb-8.9gm%, Blood urea 60mg/dl, Serum Creatinine 1.8 mg/dl, ESR-26mm/1hr. Urine examination reveals turbid, dark yellow color, 100/120 pus cells. Ultrasound of KUB region reveals sub-acute Pyelonephritis. The patient was advised to take plenty of water and following *Ayurvedic* medicine for four weeks.

Gokshuradi guggulu: 2 tab twice daily after food.

Syp. Neeri KFT: 3 tsf twice daily after food.

Chandraprbha vati 2: tab twice daily after food.

After four weeks of treatment patient was again assessed. He was symptom free and the WBC-6x103, Hb-12.50 gm%, ESR-10mm/1hr, Blood urea 32mg/dl, Serum Creatinine 1.0 mg/dl Urine became light yellow color, clear. Pus cells in urine 3-5. USG not repeated because patient was asymptomatic and patient refused to do.

# **Discussion**

When patient came to OPD he was not in acute phase of Pyelonephritis. He already has taken antibiotics for one week. He was in the sub acute phase of Pyelonephritis. The possible reason behind the infection may be the steroid which he consumed before six months for body building. Steroids compromise the immune system of the body that leads to the infection. The medicines used in the treatment of patient are combination of multiple drugs. Most of the drugs of combination are advised for the treatment of renal calculus and mentioned in *mutraghat* and prameha chikitsa in Sushruta samhita. The main drug of Gokshuradi guggulu is Gokshur (tribulus terrestris), and from Neeri-KFT the main drugs are Punarnava (Boerhaia diffusa), panchtrinmool (classical combination of five drugs), kansi (cichoriou mintybus), Makoya (solanumnigrum), Giloe (tinosporia cardifolia), kamalkandi (nelumo nucifera), Palaspushpa and Gokshur etc. the main composition of Chandraprabhavati is Shilajit (Asphaltum punjabianum) and Guggulu (commiphora mukul). Gokshuadi guggulu and Chandraprabha vati these are the two classical drugs mention in Ayurvedic text in treatment of Prameha/Mutrarog<sup>7</sup> (urinary disorders) and Neeri KFT is branded drugs by AIMIL pharmaceutical limited. The possible mode of action of the drugs is that these drugs stimulate the renal parenchyma cells for enhancing active and fast filtration that produce more urine. That produces a kind of flush action for kidneys as well as whole circulating fluid, which reduces the loads of pathogens. Gokshura<sup>8</sup> is well known for diuretic action. The drugs like Giloe<sup>9</sup> & Shilajit<sup>10</sup> are very good immune booster which augment the defense mechanism of the body to stop the pathogens. The drug like Guggulu<sup>11</sup> is a potent anti-inflammatory, analgesics, and antipyretics which reduce the pain, malaise and fever.

# **Conclusion**

Pyelonephritis is an infective disorder as per modern medical science. In the acute phase antibiotics drugs are required. In sub-acute condition *Ayurvedic* medicine have very good effect in clearing of the pathogens from kidneys because of enhancing the glomerular system of kidneys by diuretic action and boosting the renal function. These drugs enhance the immune system to improve the defense mechanism of body. So the results of this particular study, it was observed that Ayurvedic medicine work very efficiently to improve this particular condition. However to get more accurate and firm result study should be carried in multicenter with large sample size.

# References:

- Aggarwal R etall , Acute Pyelonephritis, Urology & Nephrology Open Access Journal, 2017, 5(5): 00187, doi:10.15406/ unoaj.201705.00187.
- 2. L UmeshaetallAcute pyelonephritis: A single-center experience, Indian journal of nephrology, 2018, 28 (6), 454-461. Doi:10.4103/ijn.IJN\_219\_16
- 3. Shields J, Maxwell AP, Acute pyelonephritis can have serious complications, Practitioner 2010 Apr; 254(1728):19, 21, 23-4, 2.
- LeelavathiVenkatesh, RamalingiahKaradakereHanumegowda, Acute Pyelonephritis Correlation of Clinical Parameter with Radiological Imaging Abnormalities, JCDR 2017-11 (6) doi: 10.7860/jcdr/2017/27247.10033
- 5. Kaviraj Dr. Ambikaduttashastri, SushrutaSamhitauttarardha, Ayurveda tattvasandipikahindi commentary, Chaukhamba Sanskrit samsthan, Varanasi, chapter 58/22-26 p-426.
- 6. Kaviraj Dr. Ambikaduttashastri, SushrutaSamhitauttarardha, Ayurveda tattvasandipikahindi commentary, Chaukhamba Sanskrit samsthan, Varanasi, chapter 58/27-28 p-427.
- 7. Dr. Brahmanand tripathi, sarngadhar-samhita of pandit sarngadharacarya, annoted with "DIPIKA" Hindi commentary, chaukhamba subharti prakashan Varanasi, chapter seven, page 137 & 133.
- 8. Saurabhchhatreetall, Phytopharmacological overview of Tribulusterrestris, Pharmacognosy review, 2014 Jan-Jun; 8(15): 45–51 doi: 10.4103/0973-7847.125530.
- S.S. SINGH etall, Chemistry and medicinal properties of tinosporacordifolia (guduchi), Indian Journal of Pharmacology 2003; 35: 83-91.
- 10. Carlos Carrasco-Gallardoetall, Shilajit: A Natural Phytocomplex with Potential Procognitive Activity, International Journal of Alzheimer's Disease, Volume 2012, Article ID 674142, doi:10.1155/2012/674142
- 11. Jain AnurekhaetallChemistry and pharmacological profile of guggul-A review, Indian Journal of Traditional Knowledge, Vol 5(4)-October 2006 -pp 478-483,

# **Case Report**

# An Ayurveda Approach to Anidra w.s.r. to Anxiety induced Insomnia: A Case Report

# <sup>1</sup>Gopesh Mangal, <sup>2</sup>Nidhi Gupta, <sup>3</sup>Pravesh Srivastava

<sup>1</sup>Associate Professor & Head, PG Department of Panchkarma, National Institute of Ayurveda, Jaipur-302002, Rajasthan

<sup>2</sup>PG Scholar, PG Department of Panchkarma, National Institute of Ayurveda, Jaipur-302002, Rajasthan <sup>3</sup>PG Scholar, PG Department of Panchkarma, National Institute of Ayurveda, Jaipur-302002, Rajasthan

Manuscript Received on 12/11/2019

Reviewed on 25/11/2019

Accepted on 27/11/2019

# **Abstract**

Ahara (food), Nidra (sleep) and Bramhacharya (celibacy) in Ayurveda are collectively described as the Trayoupastamba (three supportive pillars) of the life. Hence forth, sleep is one of the essential factors for a healthy life. Ayurveda also mentioned Nidra as one of the important dimensions for happiness and good health which finally leads to relax mental state. Anidra can be clinically correlated with insomnia. Insomnia is the most common sleep problem worldwide. Long term manifestation of insomnia can lead to many psychosomatic disorders like fatigue, high blood pressure, lack of concentration and ultimately reduce the productivity and hampers the quality of life. The national sleep foundation of America estimates that at least 1/3rd people of world suffer from sleep disorders. Ayurveda advocates some fruitful Panchkarma therapies for the treatment of Anidra. Hence an effort has been made to evaluate the efficacy of Shirodhara and Shamana Chikitsa in the management of the Anidra (insomnia).

This is single case study of 37 years old male suffering from sleeplessness for 10 years. He had associated complaints of excessive thoughts, indigestion and headache. Patient sleep was markedly disturbed leading to stress and anxiety hence he visited to the allopathic hospital and started anti-psychotic and sedatives drugs. Patient took medication regularly for 8 years but the symptoms didn't get subsided. Treatment given was *Shirodhara* with *Ashwagandha Taila* for 45 min for 14 days along with *Shamana Chikitsa*.

The therapy provided marked relief in stress and sleeplessness and patient quality of life was improved. On the basis of this case study it can be concluded that *Panchkarma* can play a vital role for the effective management of *Anidra* (insomnia). Since the single case is not enough more rooted study in this is required. **Keywords** - *Anidra*, *Shirodhara*, *Ashwagandha Taila*, Insomnia

# Introduction

Ahara (food), Nidra (sleep) and Bramhacharya (celibacy) in Ayurveda are collectively described as the Trayoupastamba (three supportive pillar) of the life<sup>1</sup>. Hence forth, sleep is one of the essential factors to lead a healthy life. Ayurveda has mentioned Nidra as one of the most important dimensions of health associated with happiness and good health which leads to relax mental state<sup>2</sup>. According to WHO (World Health Organization), health is a state of complete physical, mental, or social well being and not merely the absence of disease or infirmity<sup>3</sup>. In present era due to changing of lifestyle and environmental factors quality of sleep has been disturbed. Good quality of sleep is essential for good health and well being. The disturbance in Nidra might be related to the life style, mental tension, change food habits and day to day stress which ultimately disturb the biological rhythm of the sleep. Acharya Charaka has described Anidra as one of the 80 Vata Nanatmaja Vikara4. Some Acharya also mentioned Anidra as Lakshana of some disease5. Anidra can be correlated with insomnia on the basis of sign and symptoms as mentioned in ancient Ayurveda literatures. Insomnia also known as sleeplessness is a sleep disorder in which people have trouble sleeping<sup>6</sup>. Insomnia is the inability to obtain an adequate amount or quality of sleep. Trouble can be falling asleep, remaining asleep, or both. People with decrease sleep don't feel refreshed when they wake up. Now a day's insomnia is a common symptom affecting millions of people. Insomnia may be caused by many conditions like psychological causes- fear, anxiety, anger etc<sup>7</sup>. Insomnia often leads to fatigue, lack of energy, difficulty concentration, irritability etc. Women are more commonly affected than men, and its incidence is found to be increased with increase in age8.

Corresponding Author: Dr. Gopesh Mangal, Associate Professor & Head, PG Department of Panchkarma, National Institute of Ayurveda, Jaipur-302002, Rajasthan, India, Email: gmangal108@gmail.com

**How to cite this article:** Mangal G, Gupta N, Srivastava P: An Ayurveda approach to Anidra w.s.r. to anxiety induced Insomnia: A Case Report; AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 27 - 31

Approximately 30% of general population are affected from insomnia<sup>9</sup>. One out of every twenty Indian suffers from sleep disorder. 16% of the population suffered from insomnia in India. Different findings suggest that, sleeplessness is an epidemic that affects an estimated 150 million in developing world. In 20 years, over 260 million people will experience sleep disorders. 5% to 6% of people aged 50 years and above are affected by sleep disorders in India<sup>10</sup>.

Different types of tranquilizers are prescribed for the management of insomnia continuous and long term of which is having various adverse effects including drug dependency. In such a scenario there is a need for the efficient management of insomnia in a natural way without leading to further adverse effects. Ayurveda classics explains different modality for the management of Anidra (Insomnia). Ayurveda advocates some fruitful Panchkarma therapies like Murdhni Taila which include Shiroabhyanga, Shirosheka, Shiropichu and Shirobasti. Hence an effort has been made to evaluate the efficacy of Shirodhara and Shamana Chikitsa (Internal medicine) in the management of the Anidra (Insomnia).

# **Case Report**

This is single case study of 37 years old male (OPD Reg. No. 29321022019) suffering from sleeplessness for 10 years. He had associated complaints of excessive thoughts, Constipation, tiredness and headache. Patient sleep was markedly disturbed leading to excessive stress and anxiety hence he visited to the allopathic hospital and started anti-psychotic and sedatives drugs. Patient took medication regularly for 8 years but the symptoms didn't get subsided. Hence patient came for the Ayurveda treatment and visited the Panchkarma OPD, NIA, Jaipur. There was no any significant past history of HTN, DM, traumatic injury etc. Astavidha Pariksha [Table No. 1], Manas Bhav Pariksha [Table No. 2] and systemic examination [Table No. 3] was done.

Table No. 1: Astavidha Pariksha

Nadi	74bpm		
Mutra	Samyaka		
Mala	Nirama		
Jivha	Nirama		
Shabda	Spasta		
Sparsha	Samshitoshna		
Drik	Spasta		
Akriti	Madhyam		

Table No. 2: Manas Bhav Pariksha

Bhaya	Absent		
Krodha	Present		
Shoka	Absent		
Chinta	Present		

Table No. 3: Systemic examination

126/80 mmHg
98.6 0F
74/bpm
18/min.
72 kg
5'9"
Normal
Absent

**Investigation** - Haematological investigation was within normal limit. The case was diagnosed as *Anidra* (Insomnia) on the basis of signs and symptoms. Patient was admitted at male IPD ward, Panchkarma department of NIA. The treatment was given according to Ayurveda principles.

#### Intervention

Shirodhara along with the Shamana Chikitsa was given to the patient [Table no. 4]. Shirodhara is a type of Murdhani Taila that involves gentle pouring of warm liquids over the forehead. It is the process in which medicated oil is poured in a continuous stream of drip on the forehead in a specific manner and height. Shirodhara therapy is extensively used for alleviation of psychic and psycho-somatic ailments. In this study Ashwagandha Taila was used for the Shirodhara procedure.

Proper counselling about the procedure was done to patient. Before procedure vitals of the patient was measured. Procedure was done during the morning hours. *Ashwagandha Taila* was kept in *Dharapatra*. Eyes of the patient was closed with the help of cotton plug and bandage to prevent the flow of oil inside the eyes. *Dhara* was poured continuously on forehead neither very fast nor very slow. Luke warm temperature of oil was maintained throughout the whole procedure. *Dharapatra* was moved in the pendulum manner starting from one lateral side to the other lateral side during the pouring of oil. The Oil was collected in another vessel and was used to refill the *Dharapatra* before it becomes empty. Procedure was done for 45 min till 14 days. The oil was changed on every 7<sup>th</sup> day. Total 2 litre of *Ashwagandha Taila* was used during the procedure. *Shamana Chikitsa* was given continuously 14 days during the procedure and after the procedure to till follow up.

Table No. 4: Intervention

S. No.	Date of start	Drug	Dose	Frequency
1		Sarswatarista	4tsf	BD after meal
2		Bramhi Ghrita	10gm	OD after meal at night
3	12/3/2019	Shankha Bhasma Bhuvneshwar Churna	250mg 4gm	BD before meal
4		Ashwagandha Taila (For Shirodhara)	2 litre	At morning time

Follow up of the patient was done after 15th day of completion of procedure.

## Assessment

Assessment was done after the completion of procedure and after the follow up. Both subjective as well as clinical improvements were employed for the assessment of the impact of the procedure. All symptoms which were selected for assessment, their improvements were thoroughly examined and the severity of each of them was rated before and after the trial of the procedure. Grading was done for the assessment [Table no. 6]. Patient was assessed from subjective criteria [Table No. 5] and also from PSQI scale [Table No7]. Krodha and Chinta of the patient was markedly reduced.

Table No. 5: Assessment after treatment and follow up

S. No	Symptoms	ВТ	AT	After follow up
1.	Jrimbha (Yawning)	1	1	1
2.	Tandra (Drowsiness)	0	0	0
3.	Arati & Klama (Fatigue &Inertness)	2	1	0
4.	Shirashoola (Headache)	2	1	0
5.	Ajirna -Agnimandhya (Indigestion)	3	0	0
6.	Malabaddhata (Constipation)	2	0	0
7.	Quality of sleep	1	2	3

Table No. 6: Grading

Sleep	Gradation
No sleep	0
Mild sleep (1-3 hrs)	1
Moderate sleep (4-5 hrs)	2
Good sleep (More than 6 hrs)	3
Quality of sleep (Is sleep continous/disturbed)	Gradation
Continuous sleep	0
Severely disturbed sleep(Disturbed at 4-5hrs)	1
Moderate disturbed sleep(Disturbed at 3-4hrs)	2
Mild disturbed sleep(Disturbed at 1-2hrs)	3

**Table No 7:** Sleep quality index (PSQI Scale) Component of PSQI scale-

Component	Interpretation	B.T.	A.T.	After follow up
C1	Subjective sleep quality	3	2	2
C2	Sleep latency	3	1	1
C3	Sleep duration	2	1	1
C4	Sleep efficiency	3	2	2
C5	Sleep disturbances	2	1	1
C6	Use of sleep medication	3	2	1
C7	Day time dysfunction	4	3	2

Global PSQI scale is the sum of all the seven components of PSQI. The total score of "5" or greater is indicative of poor sleep quality.

# **Discussion**

Insomnia is a sleep disorder in which people have trouble to sleep. Physiologically it can be defined as the state of decrease functioning of thalamus and basal fore brain. Pharmacological studies suggest the role of different neurotransmitter for the disturbance of sleep. Catecholamine and serotonin level plays the vital role in the promotion of sleep.

Shirodhara is a procedure which comes under the external application of oil over the fore head mostly used for neuromuscular relaxation and nourishment. Shirodhara is a relaxation therapy which relieves mental exhaustion as well as pacifies the aggravated Vata Dosha. It helps to normalize the function of central nervous system by relaxing the nervous system and balancing the circulation of blood in the head. Shirodhara acts as anti-anxiety and stress relaxant and these effects are mediated by the brain wave coherence, Alfa waves, and a down regulation of the sympathetic outflow<sup>11</sup>. As it is proven that local ointments after application passes through stratum corneum and reaches the target organ same wise the continuous pouring of Ashwagandha Taila may also absorbed and stimulates the brain cortex, corrects the deficiency of neurotransmitter thus producing tranquilizing effect. Shirodhara also pacify the disturbed Manasika Bhava like Krodha, Chinta, Bhaya, etc promoting mental health<sup>12</sup>. Shirodhara produces calming effect as after procedure it makes patient relax and induces sleep may be due to raise in serotonin level. Shirodhara when continuously poured over fore head with particular pressure produces a kind of vibration. These vibrations reaches to the target organ through cerebrospinal fluids thus stimulating the thalamus and basal fore brain and helps in the correction of catecholamine and serotonin level<sup>13</sup>. Correction of neurotransmitters helps in promotion of sleep.

In *Anidra* (insomnia) the main vitiated *Dosha* is *Vata*, so balance of *Vata Dosha* is an important thing to treat in insomnia. *Ashwagandha Taila* having *Vatahara* properties helps in the suppression of symptoms like excessive thoughts, sleeplessness and headache. Main component of above oil is *Ashwagandha* which is having *Nadibalya*, *Mashtiska Shamak*, *Nidrajana* and *Vatahara* properties. Bhuvneshwar *Churna* and *Shankha Bhasma* were given to correct the *Agni* of the patient and to improve digestion. As per *Ayurveda* text medicines like *Bramhi Ghrita*, *Sarswatarista* are some of the widely indicated formulation for Psychiatric illness like *Unmada*, *Atatvaabhinivesha* etc. *Brahmi* is a proven brain tonic, act as neuro-activator and helps to improve mental & intellectual activity, further when administered in the form of ghee it is easily absorbed due to lipophilic nature. *Sarswatarista* was used as *Medhya Rasayana*. *Sarswatarista* having *Tridoshara*, *Rasayana*, *Ojaskara* and *Medhya* properties which provide proper nutrition to the brain and stimulate its function. All these together helps to improve brain functioning and provides nourishment to brain which is responsible for the suppression of aggravated *Vata Dosha* thereby overcomes stress & anxiety, calm down the mind and in turns improves quality and duration of sleep.

# **Conclusion**

In overall assessment it was found that *Shirodhara* with *Ashwagandha Taila* along with *Shamana Chikitsa* have beneficial role for the management of *Anidra* (Insomnia).

Conflicts of Interest: Not any

#### References

- Charaka Samhita, Vidyotini Hindi commentary vol.1,Sutrasthana, 11/3 Chaukhamba Bharati Akadmi; Varanasi By Dr. Gorakha Nath Chaturvedi & Pd. Kashi Nath Shashtri, page no.332
- 2. Charaka Samhita, Vidyotini Hindi commentary vol.1,Sutrasthana, 21/3 Chaukhamba Bharati Akadmi; Varanasi By Dr. Gorakha Nath Chaturvedi& Pd. Kashi Nath Shashtri, page no.332
- 3. Constitution of the World Health Organization in World Health Organization; Basic documents, 45th edition, Geneva; WHO 2005
- Charaka Samhita, Vidyotini Hindi commentary vol.1,Sutrasthana, 19/3 Chaukhamba Bharati Akadmi; Varanasi By Dr. Gorakha Nath Chaturvedi& Pd. Kashi Nath Shashtri, page no.332
- A case study insomnia, Kalpna V. Satpute et al; Internalnational Journal of Applied Ayurveda Research ISSN: 2347-6362.
- 6. Ayurvedic therapy (Shirodhara) for Insomnia: A Case Series, Sivarama Prasad Vinjamury et al. gahmj.2012.086%20(1).
- Nidranasha (insomnia) cause, consequences &management an Ayurvedic perspective, Nirgude Rajendra et al; International journal of Herbal Medicine, ISSN 2321-2187.
- 8. Ohayon MM. Epidemiology of Insomnia: What we know and what we still need to learn. Sleep. Med Rev. 2002; 6(2): 97
- 9. Roth T. Insomnia: definition, prevalence, aetiology and consequences. J Clin Sleep Med. 2007.
- 10. Insomnia rampant in third world countries: Study. http://www.Health.india.com (accessed 4 March 2013).
- 11. Dhuri, K. D., Bodhe, P. V., & Vaidya, A. B. (2013). Shirodhara: A psycho-physiological profile in healthy volunteers. Journal of Ayurveda and integrative medicine, 4(1), 40–44.
- 12. Kundu, C., Shukla, V. D., Santwani, M. A., & Bhatt, N. N. (2010). The role of psychic factors in pathogenesis of essential hypertension and its management by Shirodhara and Sarpagandha Vati. Ayu, 31(4), 436–441.
- 13. Pokharel, S., & Sharma, A. K. (2010). Evaluation of Insomrid Tablet and Shirodhara in the management of Anidra (Insomnia). Ayu, 31(1), 40–47.

# **Case Report**

# Verruca Vulgaris of scalp annihilated by Homoeopathic Medicine - A case report

Partha Pratim Pal

Research Officer (H)/ Scientist -I, Dr. Anjali Chatterji Regional Research Institute (H), Kolkatta

Manuscript Received on 19/05/2019

Reviewed on 11/09/2019

Accepted on 14/10/2019

# **Abstract**

Verrucas (warts) are cutaneous lesions caused by human papilloma viruses which can occur on any part of the body by skin contact directly or indirectly. Treatments available are mostly invasive i.e. either surgical excision or external application of some corrosive material. Only judicious use of homoeopathic medicines can be effective in absolute eradication of these types of pathological growths. A male patient, 51 years of age, who was coming to one of the *Swasthya Rakshan* camp, complained of fleshy, sessile, indurated mass on the upper occipital part of the scalp which was growing conspicuously for the last 6 months. After detailed case taking, miasmatic evaluation and repertorization, Carcinosin was given in centesimal potency. In a span of four months, the verruca was nowhere to be found on the scalp.

Keywords - Carcinosin, Case report, Individualization, Scalp, Swasthya Rakshan camp, Verruca.

# Introduction

Verruca commonly known as warts is caused by infection of skin by human papilloma virus (HPV: ds- DNA viruses growing only in the epidermis). It can occur in adults and children and infection is transmitted by direct contact although it has a low infectivity and therefore a casual contact may not result in the local malady1. HPV infections may be clinical, subclinical or latent 2. Clinical types are visible to the naked eye and can be diagnosed correctly by history and physical examination alone3. They can manifest in different forms and on basis of location they can be categorised as - verruca vulgaris (sessile, dome-shaped and usually a centimetre in diameter) 4, palmoplantar warts (multiple, confluent/discrete, painless) verruca plana (flat, smooth papular), filiform warts (thread like/finger like projections), epidermo-dysplasia verruci formis (genetically predisposed, extensive lesions), and anogenital warts (sexually transmitted) 5. Their morphology and texture may be soft, hard, flat, smooth, rough, scaly etc. Warts also have diagnostic value in distinguishing different stages in a disease. The verruca vulgaris appear at the time of second dentition in cases of children suffering from hereditary sycosis. The verruca filiform is comes as a tertiary lesion in an acquired form of sycosis. The verruca plana juvenilis (pigmented, disseminated and in irregular unilateral groups) is another form found on the back of hands and faces of children and young people<sup>6</sup>. Management of warts is based on their clinical appearance, location and the immune status of the patient2. The various modalities available for the treatment of warts in various systems of modern medicine include electrocoagulation, cryosurgery, curettage, applying salicylic acid, liquid nitrogen and podophyllin, CO2 laser surgery and interferon injection7.

# **Case Report**

A male patient (PT) of 51 years of age came to the OPD of Swasthya Rakshan camp(note),complained of fleshy, sessile, indurated mass on the scalp(at the upper occipital area) which has developed to a size of 3cm x 2 cm within a period of 6 months. Surface was rough and margins diffuse. There was no pain, itching or burning sensations. Bleeding in tinge was found occasionally when got struck by comb or hand accidently.

### Past history

Patient has history of chicken pox at the age of 14 years which healed spontaneously with proper rest and diet. He has tendency to develop vesicular eruptions after mosquito bite.

# Family history

Grandfather died of throat carcinoma (cannot exactly specify whether it was pharyngeal or laryngeal or anything else). Mother suffers from uric acid diathesis and obesity. Father is also suffering from diabetes mellitus for last 10 years.

**Corresponding Author:** Dr. Partha Pratim Pal, Research Officer (H)/ Scientist –I, RRI (H), Agartala, Under CCRH, India, Email: justdoit.partha007@gmail.com

**How to cite this article:** Pal PP: *Verruca Vulgaris* of scalp annihilated by Homoeopathic Medicine - A case report; AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 32 - 37

### **Personal history**

Patient was government employee at Group-B level in an administrative section. He was a singer also and occasionally he conducts stage performances on contract basis.

### Generalities

# Physical generals

His appetite was good and regular. There was a strong desire for meat and aversion towards fruits and vegetables. He also had the predisposition of developing flatulence in abdomen towards evening. Bowel movements were irregular with so much hard stool that at times he was reluctant to go for defecation. Patient also had tendency of sweating all over the body but especially during sleep. Sleeplessness was prominent due to activity of thoughts which keeps on coming to his minds on lying down and hence on waking feels un-refreshed. He always felt better in winter season, cannot tolerate the heat in any form and hence to be considered as hot patient.

### Mental generals

Mentally he was very joyful and becomes more cheerful during stormy and windy weathers. He felt more energised and singing spontaneously comes to him during such conditions. He described himself as an 'Explorer' who always wants to travel to new places to see original facts and to learn innovative things. At the same time he was a perfectionist, very particular about his physical presentations and performances. Inspite of all this positivity in the mental sphere he had a strong fear of having cancer. Whenever he had problems regarding the voice like hoarseness, cracking of voice etc he got anxious. He kept on thinking that it might turn fatal and he may die because of throat cancer just like his grandpa.

# Local and systemic examination

On inspection fleshy mass (on the upper occipital area on the left side) was visible on the removal of the hair with hands. It was reddish on appearance. No ulcerative spot on the mass was detected. On palpation, it was sessile, fixed to skin of the scalp, no fluctuation found and was having firm consistency. Tongue was clean and moist. Skin was having numerous black moles all over.

### Analysis of the case

After analysing the symptoms of the case, the characteristic mental and physical generals and particular symptoms were considered for framing the totality. Fear of cancer, cheerful mood during thunderstorm, desire to travel, and fastidiousness were the mental generals. Desire for meat, aversion to fruits and flatulent abdomen during evening, hard stool, profuse perspiration mainly during sleep, unrefreshing sleep, sleeplessness from activity of thoughts and thermal reactions of being a hot patient were the physical generals in the totality. Regarding the particular symptoms, warts on scalp and numerous black moles all over body were considered.

### Miasmatic analysis

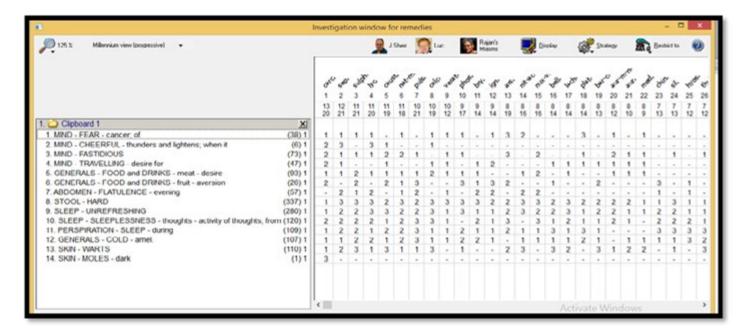
Miasmatic evaluation of all the presenting symptoms was done with the help of "Chronic miasms in homoeopathy and their cure with classification of their rubrics/symptoms in Dr. Kent's repertory by Dr. RP Patel" showed the predominance of psoricmiasm with syphilitic adulteration8 hence it is quite rightly stated that the miasm of psora unfolds itself in many countless disease forms in all human race.

Table no 1 -

Symptoms/Rubrics	Miasm
Mind: Cheerful – thunders and lightens; when it (+++)	Psora
Mind: Travelling – desire to (+++)	Psora
Generals: Desires meat (++)	Psora
Generals: Aversion fruits (+)	Psora
Abdomen: flatulence – evening (++)	Psora
Stool: hard (++)	Psora
Stool: odour, offensive (++)	Psora
Sleep: unrefreshing (++)	Latent psora
Sleep: sleeplessness – from activity of thoughts (++)	Psora, Syphilis
Generalities: warm aggravation (+++)	Syphilis
Skin: Warts - bleeding (+++)	Psora

# Reportorial analysis

Table no 2 -



Considering the above symptomatology, The Synthesis repertory 9.1 version was selected and RADAR software used for repertorization<sup>9</sup>. Therepertorisation chart is provided above. After repertorisation, many medicines were competing with each other, namely Carcinosin, Sepia, Sulphur, Lycopodium, Causticum, Natrummuriaticum etc. Carcinosin covered maximum symptoms (13) but Sepia and Sulphur scored the maximum (21). But Carcinosin was preferred as evidences stated that it acts favourably and modifies all cases in which history of cancer is present in the family<sup>10</sup>. H/O diabetes mellitus in the family also goes in favour of the prescription<sup>11</sup>. Moreover there are evidences of successful prescription of carcinosin on warts. A pedunculated painless wart on anterior side of neck for one year was cured by Carcinosin 1M in two months<sup>12</sup>. Two doses of Carcinosin 1M given in sac lac pulvis was prescribed, to be taken in empty stomach, OD x 2 days, followed by placebo for one month on 20.08.15, after considering the reportorial totality and miasmatic background. Unfortunately no picture was documented on day one. The following pictures are from day two onwards.

Follow-Up Date	-Up Date Indications for Prescription Medicine with Doses	
1 <sup>st</sup>	Picture 1 – bleeding spots still	Carcinosin 1M/ 2 doses in sac lac
follow-up	found. Size almost the same. No	pulvis
01/10/15	such remarkable improvement.	OD x 2 days, followed by one month
		placebo.



Follow-Up Date	Indications for Prescription	Medicine with Doses
1 <sup>st</sup>		Placebo continued for one more
follow-up	more occurring. Margins seem to	month for once daily.
01/10/15	condense and the mass seemed to	
	shrink.	





Indications for Prescription	Medicine with Doses
Picture 4 and 5 - no bleeding episodes. The growth decreased to a great extent and almost	month for once daily.
	Picture 4 and 5 – no bleeding episodes. The growth decreased





Follow-Up Date	Indications for Prescription	Medicine with Doses
4 <sup>th</sup> follow-up 07/01/16	Picture 6 – growth vanished 2 weeks earlier and patient came just to inform.	·



# Discussion

Homoeopathic treatment of warts is mainly through constitutional medicine i.e. medicine selected is based on characteristic history and totality of the symptoms obtained from complete study of the patient as an indivisible whole <sup>13</sup>. It can be classified as external local malady which refers to one-sided diseases having alterations and ailments on the outer parts of the body. Their cause is an internal suffering requiring homoeopathic medicines taken internally only (no external means should be directly applied) 14. Several published articles have shown the effectiveness of homoeopathic treatment on warts. Out of 62 cases in from an outpatient study, in 47 cases warts disappeared completely and medicines prescribed was mainly Thuja, Ruta, Antim crud, Calc carb, Nitric acidum and Opium in 30th and 200th potency 15. Five successful case reports on warts located in left and right arms, nose, upper lip, and cheek was reported in a journal. Medicine prescribed was Thuja (2 cases) and Causticum (3 cases) in centesimal and LM potency<sup>16</sup>. Another such publication of two case reports on warts located over feet and eyelid was also found in another journal. In these cases, the medicines were Nitricumacidum and Causticum in 30th potency. In both the case report publication, evidence-based documentation was done by photography before and after the treatment 17. A study of randomized double-blind placebo-controlled trial to evaluate the efficacy of a homoeopathic treatment of plantar warts which surprisingly concluded that homoeopathic treatment was no more effective than the placebo. The trial lasted for 6 weeks and the conclusion was surprising for the homoeopaths. The reason behind such a negative result may be due the mode of prescription done as it was a polypharmacy (Thuja 30, Antim crud 7 and Nitricum acidum 7) consisting of 3 remedies. Secondly it is not necessary that these medicines must be effective in all cases of warts. A recent study of homoeopathic treatment for various types of warts found that 18 out of 19 people with plantar warts were cured in on average of 2.2 months. The most common remedy was Ruta (12 out of 19 cases), along with Thuja (3 cases) and Antim crud (2 cases) 18. From above studies the importance of individualisation is evident and constitutional medicine becomes

the most selected similimum in all occasion. Hence, treatment with a single well indicated remedy as per the characteristic totality is desirable for a favourable outcome.

# **Conclusion**

Verrucas are expressions of deranged vitality reflected on skin and should not be treated as regional maladies. They have been considered as local maladies and treated accordingly with suppressive applications. They may disappear within months or may take several years and can also recur at times, but a well selected suitable remedy is actually required to cure the lesion in its whole extent.

Note: Swasthya Rakshan Program has been initiated through Central Council for Research in Ayurvedic Sciences (CCRAS), Central Council for Research in Unani Medicine (CCRUM), Central Council for Research in Homoeopathy (CCRH) and Central Council for Research in Siddha (CCRS) in selected districts/ villages with the objective of promoting health and health education in villages as per the directions of Ministry of AYUSH.

#### References

- Gupta R, Manchanda RK. Textbook of Dermatology for Homoeopaths. 5thedition. New Delhi. B. Jain Publishers Pvt. Ltd.; 2004. Pa no 90
- 2. James WD, Berger TG, Elston DM. Andrew's diseases of the skin clinical dermatology. 10th edition. Canada. Saunders Elsevier; 2006. Pg no 403
- 3. Reichmann RC. Human papilloma virus. In: Kasper DL, Fauci AS, Longo DL, Braunwald E, Hauser SL, Jameson JL, et al., editors. Harrison's Principles of Internal Medicine. Vol. II. 17th ed., New Delhi: McGraw Hill; 2008. Pg no 1117
- Calvin MO, Lawley TJ. Eczema, Psoriasis, Cutaneous infections, Acne and other common skin disorders. In: Kasper DL, Fauci AS, Longo DL, Braunwald E, Hauser SL, Jameson JL, et al., editors. Harrison's Principles of Internal Medicine. Vol. II. 17th ed., New Delhi: McGraw Hill; 2008. Pg no 319.
- 5. Khanna N. Illustrated synopsis of Dermatology and sexually transmitted diseases. 1st printed edition in India. New Delhi. RELX India private limited; 2016. Pg no 287
- 6. Oza PM. Homoeopathic management of warts. Indian journal of drugs in dermatology. January-June 2016. Volume 2. Issue 1. Pg no 45-47
- 7. Sterling JC, Handfield Jones S, Hudson PM; British Association of Dermatologists. Guidelines for the management of cutaneous warts. The British Journal of Dermatology 2001; 144:4 11
- 8. Patel RP. Chronic miasms in homoeopathy and their cure with classification of their rubrics/symptoms in Dr. Kent's repertory. Indian edition. Indian Books and Periodicals Publishers. New Delhi. 1996. Pg no 10, 82, 436, 440, 493, 576, 1167, 1168, 1203, 1225, 1258
- 9. RADAR 10. Archibel Homoeopathic Software. Belgium. 2009
- 10. Murphy R. Lotus Materiamedica. 2nd revised edition. Reprint edition 2007. New Delhi: B. Jain Publishers (P) Ltd.; 2010. Pg no 389
- 11. Dey SP. Bronchial asthma An integrated approach. pg no 27.Available from http:// books.google.co.in [ last cited on 19.09.19]
- 12. Rasal AP. A case of Carcinosin. June 2008. Available from http:// homoeopathyworldcommunity[ last cited on 19.09.19]
- 13. Principles of Prescribing. National health portal. Available from www.nhp.gov.in (last cited on 10.10.19)
- 14. Hahnemann S. Organon of Medicine. 26thimpression, Translated from the fifth edition with an appendix by R. E. Dudgeon, with additions and alterations as per sixth edition, translated by William Boericke and introduction by James Krauss. New Delhi: B. Jain Publishers (P) Ltd.; 2010: 105-106
- 15. Gupta R, Bhardwaj OP, Manchanda RK. Homoeopathy in the treatment of warts. British Homoeopathic Journal. April 1991. Volume 80. Pg no 108
- Shaikh MI. Case studies for treatment of warts with homoeopathy. Indian Journal of Research in Homoeopathy. October-December 2016. Volume 10. Issue 4. Pg no – 272- 277
- 17. Mondal A. Homoeopathic treatment of warts. Homoeopathy 360. January 2017. Available from http://www.bjain.com [ last cited on 19.09.19]
- 18. Labrecque M, Audet D, Latulippe LG, Drouin J. Homeopathic treatment of plantar warts. Canadian medical association journal.1992 May 15; 146(10):1749-53.

### **Case Report**

# Homoeopathic approach to common viral skin diseases- Case studies

<sup>1</sup>Sangita Saha, <sup>2</sup>Koushik Bhar, <sup>3</sup>Mahakas Mandal

<sup>1</sup>Reader, Department of Organon of Medicine, The Calcutta Homoeopathic Medical College and Hospital <sup>2</sup>Postgraduate trainee, Dept. of Homoeopathic Materia Medica, National Institute of Homoeopathy, Govt. of India

<sup>3</sup>Postgraduate trainee, Dept of Practice of Medicine, The Calcutta Homoeopathic Medical College and Hospital

Manuscript Received on 19/08/2019

Reviewed on 19/10/2019

Accepted on 14/11/2019

# **Abstract**

Skin diseases are among the most common of all human health afflictions and affect almost 900 million people in the world at any time and are associated with disfigurement, disability, stigma and psychological distress. Viral skin diseases range from simple superficial exanthemas to complex systemic diseases affecting people of all ages. The diagnosis is based on typical patterns of presentation, occasionally requires laboratory testing and treatment is based on the patient's comorbidities, the extent, location and progress of the disease. Homoeopathy can play a major role in the treatment of viral skin diseases and it considers the man as a whole while forming the totality of symptoms of a case of disease. Five cases of common viral skin diseases namely Molluscum contagiosum, Herpers zoster, Filiform warts, Digitate wart and Flat warts were treated with internal homoeopathic medicines selected on the basis of totality of symptoms of each case and without any external application. The treatment not only removed the local skin lesions but also showed considerable improvement in the comorbidities of the patient.

Key words: Homoeopathic treatment, Individualistic approach, Viral skin disease.

# Introduction

Skin diseases are numerous and a frequently occurring health problem affecting all ages.¹ The Global Burden of Disease project has shown that the skin diseases continue to be the 4<sup>th</sup> leading cause of non-fatal disease burden worldwide.² Skin and subcutaneous diseases were the 18<sup>th</sup> leading cause (1.79%) of global Disability-Adjusted Life Year (DALY), of which viral skin diseases accounted for 0.16%.³ Viruses are obligatory intracellular parasites and viral skin infections occur when a virus penetrates the stratum corneum and infects the inner layers of the skin,resulting in herpes simplex, shingles (herpes zoster), molluscum, warts, etc.⁴-6 Some systemic viral infections, like chicken pox and measles, may also affect the skin.⁵

**Wart:** Skin warts are benign tumours caused by infection of keratinocytes with human papilloma virus (HPV), visible as well-defined hyperkeratotic protrusions. They are common throughout the world and are more common in immuno-compromised patient. They spread by direct or indirect contact and are a significant cause of concern and frustrations for patients. Viral warts show acanthosis and hyperkeratosis, usually with the characteristic feature of koilocytosis of upper keratinocytes. Warts on the skin may present in a number of different morphological forms dependent of virus type, body site, environmental influences and immunological status of the patient. They may present as common warts, periungual warts, plantar warts, plane (flat) warts, filiform or digitate warts, etc.<sup>4,5,7</sup> Social activities can be affected or lesions can be uncomfortable and often the treatment available can become painful and frustratingly ineffective.<sup>4</sup>

**Herpes zoster:** Herpes zoster results from reactivation of varicella-zoster virus (that also causes chickenpox). In the beginning one may notice a tingling sensation or burning pain on one side of the body or face. Within days, tiny clusters of red bumps quickly develops into a group of painful blisters and get crusty with pus. Lesions follow a dermatomal distribution, with thoracic and lumbar roots being the most common. Unless the patient is immuno-compromised, the rash almost never crosses the midline of the body (it's unilateral), as it is localized to one nerve root. The area can become very painful, itchy, and tender. Pain severity correlates with the extent of lesions and elderly persons tend to have severe pain. <sup>4</sup> After one to two weeks, the blisters heal and form scabs, the duration depends on patient's age, severity of eruption and presence of underlying immune-suppression<sup>4,5,7</sup>. Up to 15% of people with herpes zoster develop deep pain called post herpetic neuralgia that follows after the

Corresponding Author: Dr. Sangita Saha, Reader, Department of Organon of Medicine, The Calcutta Homoeopathic Medical College and Hospital, India, Email: dr.sangita@rediffmail.com

How to cite this article: Saha S, Bhar K, Mandal M: Homoeopathic approach to common viral skin diseases- Case studies); AYUHOM: Vol. 6, Issue 1 (Jan – June, 2019); 38 - 46

infection has run its course. It can continue for months or even years, especially in older people. The incidence of shingles and of post herpetic neuralgia rises with increasing age. Shingles usually occurs only once, although it has been known to recur, usually in people with weakened immune systems.

**Molluscum contagiosum** – A viral skin infection that causes either single or multiple raised umbilicated, pearlike bumps (papules), averaging 3-5 mm on the skin. It is caused by closely related types of Pox virus (MCV-1 to MCV-4) and is common in children <sup>4</sup>. It occurs worldwide, but seems much more frequent in geographic areas with warm climates<sup>6</sup>. Usually it is transmitted by contagion through direct contact and can spread on a single individual through scratching and rubbing. <sup>4,5,8</sup> Lesions tend to be more on the face,trunk and extremities. Irritated lesions may become crusted and even pustular, simulating secondary bacterial infection. It has a characteristic histopathology affecting follicular epithelium with presence of Molluscum bodies or Henderson- Paterson bodies in the cytoplasm. The diagnosis of Molluscum contagiosum can be done with distinctive central umbilication of the dome shaped lesion and can be further confirmed with the pasty core when the lesion is expressed, squashed between two microscopic slides, stained with Wright, Giemsa or Gram stain. <sup>4</sup>

Homoeopathy treats the patient, not the disease. So skin manifestations are also to be treated by means of internal medication following the principles of Homoeopathy. In the management of such cases along with the exact character of the local affection, all the changes, sufferings and symptoms observed in the patient are to be taken into consideration for the formation of totality of symptoms. The medicine thus administered internally not only removes the general morbid state of the body but also the local affection at the same time.<sup>9</sup>

# **Case Reports**

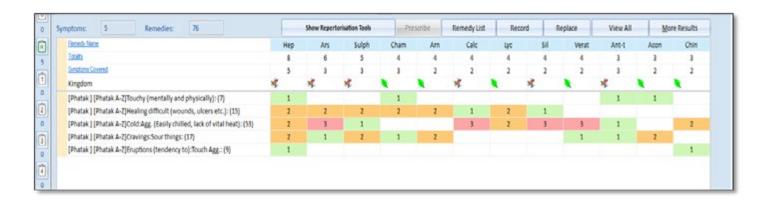
#### Case no. - 1

Mrs. R.D. Female, aged 27 years, presented on 04/02/2019 with papular eruptions at the right side of nose since last 7 days. Her symptoms included pain and itching at that part along with sense of dryness. Pain and itching were worse at night. Local examination revealed white papular umbilicated eruptions at the right side of nose with dryness of the part and painful to touch. No excoriation.

History revealed that the patient suffered from jaundice at the age of 14 years, treated by home remedies and recovered.

Patient's appetite was poor with desire for sour things. She was chilly and had tendency to slow healing. She was very sensitive both physically and mentally.

# Diagnosis: Molluscum contagiosum



After repertorization through Phatak repertory <sup>10</sup>, consulting Materia Medica and considering the whole case Hepar sulphuris was chosen.

Date of visit	Symptoms with intensity in Numeric Rating Scale – NRS (0to10)	Treatment	Photograph
04/2/2019	<ul> <li>Papular eruption at the right side of nose. (NRS- 7)</li> <li>Part very sensitive to touch and painful. (NRS-9)</li> <li>Itching at that part &lt; night. (NRS-5)</li> <li>Sense of dryness, &lt; night. (NRS-5)</li> </ul>	Hepar sulphuris 200, 1dose followed by Placebo	
08/2/2019	<ul> <li>The papules started drying up.         (NRS- 3)</li> <li>Pain and sensitiveness both reduced.         (NRS-5)</li> <li>Itching and Sense of dryness also reduced significantly (NRS-1)</li> </ul>	Placebo	
15/2/2019	There were no sign of papules, all dried up. Skin became normal.	Placebo	

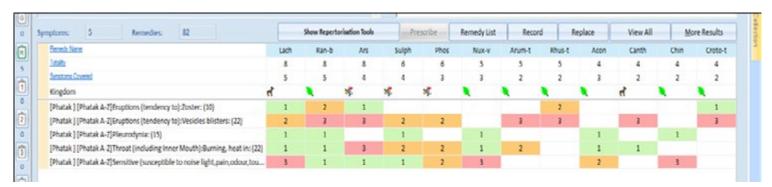
### Case no. - 2

Mrs. P. S., Female, 48 years, Housewife, presented on 11/03/2019 with vesicular eruptions at the left side of back along with extensive burning pain and itching since 2 days. Patient also complained of fever, myalgia and burning sensation in throat for last 2 days. She was unable to lie, turn or move owing to pain in left intercostal regions. On examination, 20-25 numbers of vesicular eruptions containing fluid were found along the left intercostal regions of back and were extensively sensitive to touch, with burning and itching.

In the past history, she suffered from chicken pox at the age of 16 years. Father had suffered from COPD and expired.

Patient was not feeling thirsty, tongue flabby, appetite moderate, ambi-thermal with normal bowel movement.

Diagnosis: Herpers zoster.



After Repertorization through Phatak repertory <sup>10</sup>, consulting Materia Medica and considering the whole case Ranunculus bulbosus was chosen.

Date of visit	Symptoms with intensity in Numeric Rating Scale – NRS ( 0 to10)	Treatment	Photograph
11/03/19	<ul> <li>20-25 numbers of vesicular eruptions at the left side of back. (NRS 7)</li> <li>Severe burning, itching and sensitivity at the site of eruption. (NRS 8)</li> <li>Myalgia, unable to lie or turn on sides. (NRS 8)</li> <li>Severe burning pain in throat. (NRS 8)</li> <li>Raised body temperature. (100° F)</li> </ul>	Ranunculus bulbosus 200, 1 Dose followed by placebo	
18/03/19	<ul> <li>Vesicular eruptions reduced in numbers and size. (NRS 4)</li> <li>Itching, burning and sensitivity reduced at the site of eruption. (NRS 4)</li> <li>Myalgia reduced, movement improved. (NRS 4)</li> <li>No pain in throat. (NRS 0)</li> <li>No fever (Body temp - 98° F)</li> </ul>	Placebo	
22/3/19	<ul> <li>Eruptions dried up maximally.(NRS 2)</li> <li>Itching, burning and pain disappeared. (NRS 0)</li> <li>General improvement of health.</li> </ul>	Placebo	
29/03/19	She presented with only altered skin colour, no other complaints.	Placebo	

#### Case no. - 3

Miss. T. H., 8 years, presented on 21/12/2018 with multiple granular, protruded warts just below the nose since 6 months. Numbers of warts were gradually increasing; even some appeared on fingers but were flat. No itching or scaling was present.

**Local examination:** 20-21 numbers of small granular warts beneath the nose, few coalescing together. 4 small granular warts on margin of upper lip. No local tenderness or bleeding found on touching the warts. Patient had taken other homoeopathic medicines earlier, but without effect.

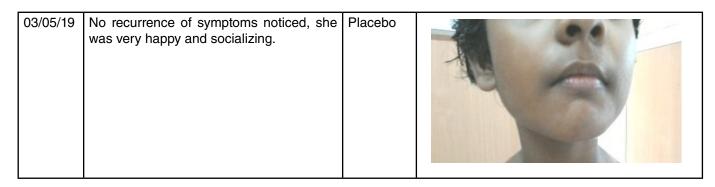
After the development of this type of wart just below the nose she felt very embarrassed and become melancholic, was avoiding socialization. She was chilly, had craving for fatty food and salt. Tongue fissured and cracked. Along with these she was losing weight gradually. In the past child suffered from measles at the age of 3 years. Family history showed her grandmother died from Carcinoma breast.

Diagnosis -Filiform warts and Flat warts



After repertorization through Phatak repertory <sup>10</sup>, consulting Materia Medica and considering the whole case Nitric acid was chosen.

Date of visit	Symptoms with intensity in Numeric Rating Scale – NRS ( 0 to 10)	Treatment	Photograph
21/12/18	<ul> <li>20-21 numbers of small granular warts beneath the nose few coalescing together.</li> <li>4 small granular warts on margin of upper lip.</li> <li>Felt very embarrassed melancholic (NRS-7)</li> </ul>	Nitric Acid 200 / 1dose	
17/01/19	<ul> <li>18 numbers of small granular warts. Clearing of 2, 3 warts at the sides.</li> <li>4 small granular warts on margin of upper lip- same.</li> <li>Appearance of flat wart at the base of fore finger dorsum of lefthand.</li> <li>Embarrassed and melancholic (NRS-7)</li> </ul>	Placebo	
21/02/19	Patient presented with slight improvement.	Placebo	
18/04/19	<ul> <li>Small granular warts 10 in number.         Scattered clearing of 8,9 warts</li> <li>Complete disappearance of warts on hand</li> <li>General improvements but was in standstill condition, since last 1 month.</li> <li>Embarrassed and melancholic reduced (NRS-3)</li> </ul>	Nitric acid 200/ 1 dose	



### Case no. - 4

Mr. P.S., Male, 65 years, Hindu, businessman presented on 16.09.16 with wart at lateral side of his right forearm, near the elbow joint, which was jagged, blackish colour in nature. The wart appeared 10 years ago as size of a pea, but was gradually increasing in size. Earlier the patient tried various allopathic medications but got no such improvement and stopped any kind of treatment. Owing to the increasing size and irregular, jagged edges he was facing difficulty in wearing clothes.

In his past history, there was Haemorrhoids, anal fistula, GB stone; all of them operated. There was also history of jaundice and recurrent malarial attacks.

Family history showed his mother had pleural effusion and his father had diabetes and arthritis.

Patient was chilly. He had cravings for oily, spicy foods, mutton; was a chronic smoker. He had history of bleeding per rectum with sticking pain, even after his operation for Haemorrhoids and fistula, with occasional bleeding per rectum. His stool, urine, sweat was also very offensive. Mentally he was very irritable and becomes angry easily as revealed by his wife. She also said that he always remains under business stress; becomes very much irritable, even curses others aggressively and even desires revenge. His irritability increased after death of his best friend from cancer of lung, 3 years back.

# Diagnosis- Digitate wart.



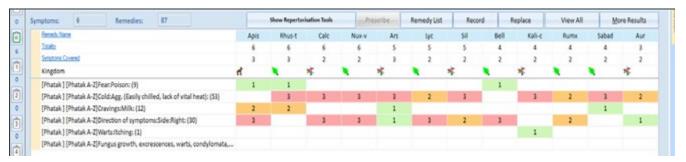
After repertorization through Phatak repertory <sup>10</sup>, consulting Materia Medica and considering the whole case Nitric acid was chosen.

Date of visit	Symptoms with intensity in Numeric Rating Scale – NRS (0 to10)	Treatment	Photograph
16.9.16	<ul> <li>Wart at lateral side of right forearm, near elbow joint jagged blackish and was gradually increasing in size. (NRS-9)</li> <li>Anger, cursing in slight contradiction (NRS-7)</li> <li>Offensive discharge of body secretions. (NRS-7)</li> </ul>	Nitric acid 200, one dose followed by Placebo	
21.10.16	<ul> <li>Wart at lateral side of right forearm near elbow joint, persisting- jagged blackish with slight reduction of size of projections. (NRS-7)</li> <li>Anger, cursing in slight contradiction slight reduced (NRS-5)</li> <li>Offensive discharge of body secretions reduced. (NRS-4)</li> </ul>	Placebo	
16.12.16	<ul> <li>Wart at lateral side of right forearm near elbow joint, persisting- blackish colour and projections reduced with reduction of size. (NRS-4)</li> <li>Anger, cursing same as that of last visit (NRS-5)</li> <li>Offensive discharge same as that of last visit NRS-4)</li> <li>But recurrence of bleeding per rectum with pain (NRS-6)</li> </ul>	Nitric acid 200, one dose	
January 2017 – August 2017	He visited the OPD once in two months, wart was gradually decreasing.	Placebo	
15.09.17	<ul> <li>The wart was almost levelled with the skin and the jagged appearance gone completely.</li> <li>Anger, cursing reduced (NRS-4)</li> <li>Offensive discharge reduced (NRS-3)</li> <li>No bleeding per rectum with pain (NRS-0)</li> </ul>	Placebo	

### Case no.- 5

Mr. M,D., 60 years old male patient visited the OPD on 25/01/2019 with multiple flat warts on the sides of right hand since 2 years. They were multiple in numbers of varying sizes (approx 8 big and 6 small)which were gradually increasing in size. Surface of warts were rough and with thickened skin over it. Itching and burning was present particularly at night. He applied allopathic and homoeopathic ointments but without any positive effect. In past he suffered from malaria 2 times at age of 28 and 45 years. He was a chain smoker, diabetic, non hypertensive, chilly person having desire for milk. He was always restless, anxious, did work very hurriedly and was having fear of being poisoned.

# **Diagnosis - Flat warts**



After Repertorization through Phatak repertory <sup>10</sup>, consulting Materia Medica and considering the whole case Rhus toxicodendron was chosen.

Date of visit	Symptoms with intensity in Numeric Rating Scale – NRS (0 to10)	Treatment	Photograph
25/01/19	<ul> <li>Flat warts on the sides of right hand(approx 8 big and 6 small), Surface of warts rough with thickened skin over it. (NRS-7)</li> <li>Itching and burning over warts, particularly at night. (NRS-5)</li> <li>Anxiety and restlessness (NRS-6)</li> </ul>	Rhus toxicodendron 10 M / 1 dose	
21/02/19	<ul> <li>Warts on sides hand flattened (approx 8 big and 6 small), smoothening of skin over the wart. No new warts (NRS-3)</li> <li>No Itching and burning. (NRS-0)</li> <li>Anxiety and restlessness reduced (NRS-2)</li> </ul>	Placebo for 1 month.	
15/03/19	All warts disappeared with return of normal skin colour.	Placebo	

# **Discussion and Conclusion**

Cases of five common viral skin disease conditions treated with Homoeopathic medicines were included in the case study. The indicated medicines were prescribed in different potencies as per the susceptibility of each individual patient and guidelines of Organon of Medicine. Placebo was prescribed as long as improvement continued. The medicines were not only efficient in removal or dissolution of the skin lesion of the viral disease, but also provided relief to the associated symptoms of the patient. All the patients were followed up for more than 3 months without any medication. No patient complained of any adverse report during the course of treatment or follow up period. Consent was obtained from the patient or their parents (in case of minor) before taking their photographs. The medicines namely Nitric acid, Rhus toxicodendron, Ranunculus bulbosus and Hepar sulphuris have been reported in the literature as useful for treatment of skin lesions as per their indications. These cases again substantiated the effectiveness of individualised Homoeopathic treatment in treatment of common viral skin diseases. Randomized controlled trials with larger sample size are warranted for validation of the result.

#### References:

- Hay RJ, Johns NE, Williams HC, Bolliger IW, Dellavalle RP, Margolis DJ, Marks R et al. The global burden of skin disease in 2010: an analysis of the prevalence and impact of skin conditions. J Invest Dermatol. 2014;134:1527–34. doi:10.1038/jid.2013.446.
- 2. Seth Divya, Khatiya C., Brown D., Esther E.F. Global burden of skin disease: inequalities and innovations; Current Dermatology Reports; 2017; 6 (3): 2014-210.
- 3. Karimkhani C., Dellavalle R. P., Coffeng Luc E., Flohr C., Hay R. J., Langan S. M., et.al. Global Skin Disease Morbidity and Mortality- An Update From the Global Burden of Disease Study 2013 JAMA Dermatol. 2017;153(5):406-412.
- 4. James William D., Berger T.G., Elstom Dirik M., Neuhaus M. Isaac; Andrew's Diseases of the Skin: Clinical Dermatology; Viral Diseases:12th ed; Elsevier: 2016: 359-417.
- 5. Rook's Textbook of Dermatology, Edited by Griffiths C, Barker J., Bleiker T., Chalmers R., Creamer D., Vol-1; 9th ed; Wilkey Blackwell 2016, Chapter 25, Viral infections; 25.46-25.50.
- 6. Wouden JC, Sande R, Kruithof EJ, Sollie A, Suijlekom Smit LWA, Koning S. Interventions for cutaneous molluscum contagiosum; Cochrane Database Systematic Reviews 2017, Issue 5.; Art No. CD004767.
- 7. Kasper, D. L., Fauci, A. S., Hauser, S. L., Longo, D. L. 1., Jameson, J. L., &Loscalzo, J. Harrison's Principles of Internal Medicine; 19th ed; New York: McGraw Hill Education; 2015.
- 8. Walker, Brian R; Davidson's Principles and Practice of Medicine, 22nd ed; Churchill Livingstone Elsevier; 2014.;1278.
- 9. Hahnemann.S., Organon Of Medicine,5th& 6th ed; B.Jain Publishers (Pvt.Ltd). New Delhi;2004.
- 10. Phatak S.R; Concise Repertory of Homoeopathic Medicines; Hompath Firefly
- 11. Numerical Rating Scale- An overview; Available from: https://www.sciencedirect.com.
- 12. Gagnier JJ, Kienle G, Altman DG, Moher D, Sox H, Riley D; the CARE Group. The CARE Guidelines: Consensus-based Clinical Case Reporting Guideline Development. Available from:https://data.care-statement.org/wp-content/uploads/2019/03/CARE-checklist-English-2013.pdf.

# **Instruction to Author for submitting Manuscript**

**AYUHOM** (ISSN 2349-2422) is a Peer Reviewed Bi-annual Research Journal of Ayurveda & Homoeopathy which is published by North Eastern Institute of Ayurveda & Homoeopathy (NEIAH), Shillong, Meghalaya-793018, an autonomous institute under Ministry of AYUSH, Government of India (Website: <a href="http://neiah.nic.in/ayuhom.html">http://neiah.nic.in/ayuhom.html</a>) with an objective of updating/ highlighting the latest developments in Medical sciences in general and Ayurveda / Homoeopathy in particular to the professionals in the field of Health Care Systems.

# Researchers may submit (1) Clinical Research Articles (2) Review articles (3) Case Reports (4) Pharmaceutical / Drugs Research

All submissions should contribute to advancement or should illuminate a particular aspect in any of the above mentioned fields. Every submission should adhere to the journal format & style, legibly written in good English, comprehensive, concise and complete. Contributors are strongly encouraged to read these instructions carefully before preparing a manuscript for submission. The manuscripts should be checked carefully for grammatical errors. Failure to follow them may result in papers being delayed or rejected. All papers are subjected to peerreview.

### **Types of Manuscripts**

**Research articles** should present new clinical/ experimental studies in elaborate form that constitute a significant contribution to knowledge. Research Papers should not exceed 08 pages.

Review articles should bring up the most important current topics or present interpretative and critical accounts, but not simple compilation, on subjects of general interest and should not exceed 08 pages.

# **Case Reports**

New, interesting and rare cases can be reported. They should be unique, describing a great diagnostic or therapeutic challenge and providing a learning point for the readers. Cases with clinical significance or implications will be given priority. These communications should have the following headings: Abstract, Keywords, Introduction, Case report, Discussion, Reference, Tables and Legends in that order. And pages should not exceed 06 pages.

### Total number of authors should not be more than 5.

Submission of an article to AYUHOM is understood to imply that it has not been either published or not being considered for publication elsewhere. The author's permission to publish his/her article in this journal implies exclusive authorization to the publisher to deal with all issues concerning copyright therein. Manuscripts with multi authors imply the consent of each of the authors.

Prepare the manuscript in A4 size page with margins 1 inch on all sides, Times New Roman font using a font size of 12. Title shall be in a font size 14, bold must be typed in single column, 1.5 spaced single-spaced throughout, including tables, graphs and figures. All section titles in the manuscript shall be in font size 12, bold. Subtitles in each section shall be in font size 12, bold face sentence case. **Cite LATIN Names and Ayurvedic terminology only in italic style font.** Justify all text by using (Ctrl+J).

The responsibility for all aspects of manuscript preparation rests with the authors. Extensive changes or rewriting of the manuscript will not be undertaken by the Editorial Board.

Standard International Units could be used throughout the text. Please do not put any hyperlinks and footnotes throughout manuscript.

### **Requirements for Article**

Manuscript should be starting with the title page and the text should be arranged in the following order:

- [1] Title
- [2] Author's name(s) & address.
- [3] Abstract
- [4] Keywords
- [5] Introduction
- [6] Materials and Methods
- [7] Results and Discussion
- [8] Conclusion
- [9] Reference/ Bibliography

### **Title Page**

Title must be brief and comprehensively represent the findings and description as written in the abstracts. Title page must contain all the desired information. Running title provided (not more than 50 characters). Do not use abbreviations in the title or abstract and limit their use in text.

Complete name Author (s) and Corresponding Author.

Page numbers included at bottom. Number all pages sequentially beginning with the title page.

# **Abstracts:**

As a summary of not more than 250 words abstracts, should be clear and factual in content. **Abstract must present the reason of the study (aims & ideas), the main findings and principal conclusions.** Emphasis may be made on new and important aspects of the study or may highlight some important observations. No abbreviations or references should be cited in the abstract.

# **Key Words:**

To identify the most important subjects covered by the article. (5-6 keyword maximum in alphabetic order)

# **Introduction:**

A concise account or a preview is required from the background of the subject, its significance and its relationships to earlier works clarified with pertinent references. Clearly state the purpose of the article. Do not review the subject extensively in the introduction.

# **Material and Methods:**

The manuscript should be presented with sufficient clarity and detail. The section of Clinical/ Experimental in Full Length Papers should include concise details on the methodology adopted; sufficiently elaborate to repeat the experiment. Data must be adequate and experimental design should be proper and accurate. Methods for which adequate references from published work can be cited are not to be described. All Physical and Spectral data should be reported. Method of Analysis should be validated.

### All possible effort must be made to give mechanism of actions.

In case of work related to plant materials, a sample of the authentic materials is to be deposited at any one of the designated institutions and their accession number or a reference of the same be quoted in the manuscript. Rationale for selection of certain solvent extracts of herbs/plants along with characterization (by way of spot tests, TLC pattern etc.) of such extracts evaluated for any activity should form part of manuscript. Use of positive and negative controls in experiments should be highlighted.

# **Results:**

The original and important findings should be stated in a logical sequence. Illustrate the results with figures or tables where necessary, but both must be kept to the minimum. Result must be precise and comprehensive and should not suffer from vagueness.

# **Discussion:**

It should contain a critical review of the results of the study with the support of relevant literature.

The principal conclusions drawn from the results and their important implications should be discussed. Do not repeat in detail data already stated in results. But if repetition is required then recommends to where it is appropriate may be included. Use generic names of drugs only unless the specific trade name of a drug used is directly relevant to the discussion.

# **Conclusion**

A brief Conclusion is desirable; this fragment should obviously state the foremost conclusions of the exploration and give a coherent explanation of their significance and consequence. It must be specific to the study.

**Illustrations:** All illustrations must be numbered using Roman numerals in their order of citation in the text. All Tables and figures must have a title and a legend to make them self-explanatory and they should be given numbers.

### **Tables**

Only MS word table format should be used for preparing tables. Tables should be numbered consecutively and bear a brief title. Tables should not be very large that they run more than one A4 sized page presented. The Journal reserves the right to crop, rotate, reduce, or enlarge the photographs to an acceptable size.

### **Figures**

Graphs and bar graphs should preferably be prepared using Microsoft Excel and submitted as Excel graph pasted in Word. As far as possible, please avoid diagrams made on white drawing paper, cellophane sheet or tracing paper with hand written captions or titles. Symbols, arrows or letters used in photomicrographs should contrast with the background.

Photographic illustrations should be in JPG format with sufficient resolution with at least 300 dpi.

#### References:

References should be numbered consecutively in the order in which they are first mentioned in the text (not in alphabetic order). Please cite the reference in introduction/ main text in Vancouver format in superscript without bracket (Ayurveda 1).

# Few Examples of writing References or Bibliography are given below:

- 1. Articles in Journals: Devi KV, Pai RS. Antiretrovirals: Need for an Effective Drug Delivery. Indian J PharmSci 2006: 68:1-6.
- 2. Volume with supplement: Shen HM, Zhang QF. Risk assessment of nickel carcinogenicity and occupational lung cancer. Environ Health Perspect 1994; 102 Suppl 1:275-82.
- 3. Issue with supplement: Payne DK, Sullivan MD, Massie MJ. Women's psychological reactions to breast cancer. Semin Oncol 1996; 23(1, Suppl 2):89-97.
- 4. Books and Other Monographs
  - a) Personal author(s): Ringsven MK, Bond D. Gerontology and leadership skills for nurses. 2nd ed. Albany (NY): Delmar Publishers; 1996.
  - b) Editor(s), compiler(s) as author: Norman IJ, Redfern SJ, editors. Mental health care for elderly people. New York: Churchill Livingstone; 1996.
  - c) Chapter in a book: Phillips SJ, Whisnant JP. Hypertension and stroke. In: Laragh JH, Brenner BM, editors. Hypertension: pathophysiology, diagnosis, and management. 2nd ed. New York: Raven Press; 1995. p. 465-78.

### **Electronic Sources as reference:**

1. Journal article on the Internet

Abood S. Quality improvement initiative in nursing homes: the ANA acts in an advisory role. Am J Nurs [serial on the Internet]. 2002 Jun [cited 2002 Aug 12];102(6): [about 3 p.]. Available from:http://www.nursingworld.org/AJN/2002/june/Wawatch.htm

### 2. Monograph on the Internet

Foley KM, Gelband H, editors. Improving palliative care for cancer [monograph on the Internet]. Washington: National Academy Press; 2001 [cited 2002 Jul 9]. Available from: http://www.nap.edu/books/0309074029/html/.

# 3. Homepage/Web site

Cancer-Pain.org [homepage on the Internet]. New York: Association of Cancer Online Resources, Inc.; c2000-01 [updated 2002 May 16; cited 2002 Jul 9]. Available from: <a href="http://www.cancerpain.org/">http://www.cancerpain.org/</a>

### **CONDUCT OF HUMAN/ ANIMAL STUDY:**

**Ethics:** The ethical standards of experiments must follow the guidelines provided by the CPCSEA (animal) and ICMR (human). Animal be as human as possible and the details of anesthetics and analgesics used should be clearly stated. The journal will not consider any paper which is ethically unacceptable. A statement permission and ethical practices must be included in all research articles under the 'Materials and Methods' section.

Authors publishing results from in vivo experiments involving animals or humans should state whether due Permission for conduction of the experiment was obtained, from the relevant ethics committees. The experimental protocol was approved by Institutional Animal Ethical Committee (IAEC) of Name of College and Place (Letter No....) with CPCSEA Registration No....

In addition, authors wishing to publish research work involving human studies should also send a letter of approval from the Institutional Ethics Committee and details of registration in CTRI (Clinical Trials Registry-India)

### **SUBMISSION OF MANUSCRIPTS**

To facilitate speedy and cost-effective submission of the full manuscript, an online submission via the Email is being offered. Authors are strongly encouraged to submit their manuscripts (the SOFT COPY IN MS WORD FORMAT) electronically, Email to: <a href="mailto:ayuhom.neiah@gmail.com">ayuhom.neiah@gmail.com</a>. All the communication will be done through Email only.

### Manuscript submission, processing and publication charges:

Journal does not charge the authors or authors' institutions for the submission, processing and/or publications of manuscripts in AYUHOM.

# Copyrights

The entire contents of the AYUHOM Journal are protected under Indian and international copyrights. The Journal, however, grants to all users a free, irrevocable, worldwide, perpetual right of access to, and a license to copy, use, distribute, perform and display the work publicly and to make and distribute derivative works in any digital medium for any reasonable non-commercial purpose, subject to proper attribution of authorship and ownership of the rights. The journal also grants the right to make small numbers of printed copies for their personal non-commercial use.

### **Peer-review policies**

All submitted manuscripts are evaluated by the Editorial Board and appropriate manuscripts will send for Review. The Editor calls upon at least two reviewers for their comments. We make every effort to reach an initial decision within **three month** of submission. Based on the reviewers comment, the Editorial Board accepts or request revisions of the manuscript. Editorial board reserves the right to reject any manuscript at any time without assigning any reasons thereof.

Copyright Transfer Agreement- Author (s) will be asked to sign a copyright form (<a href="http://neiah.nic.in/ayuhom/Copyright%20Agreement%20Form.pdf">http://neiah.nic.in/ayuhom/Copyright%20Agreement%20Form.pdf</a>) when the manuscript is accepted for review/ publication. All authors must read and agree to the conditions of copyright form, return the signed scanned Copyright form within 05 days via email: <a href="mailto:ayuhom.neiah@gmail.com">ayuhom.neiah@gmail.com</a>. Any article accepted for publication/published in the AYUHOM will be the copyright of the journal. The journal has the right to publish the accepted articles in any media (print, electronic or any other) any number of times.























# Published by

North Eastern Institute of Ayurveda & Homoeopathy (NEIAH)
Mawdiangdiang, Shillong, Meghalaya -793018 (INDIA)
(An autonomous Institute under the Ministry of AYUSH, Government of India)
E-mail: ayuhom.neiah@gmail.com / neiahshillong@gmail.com / dir-neiah@nic.in
Telephone: +91-364-2538134; Website: www.neiah.nic.in