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Inflammatory Joint Disease in Ayurvedic Medicine

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Abstract:

Amavata is a disease of chronic joint and body pain, accompanied by a swelling of some or all of the synovial joints. These symptoms are typically accompanied by immobility, a loss of taste, thirst. indigestion, a lack of enthusiasm, a feeling of heaviness, and fever. If the condition is allowed to progress the pains may begin to migrate from place to place, with an intense stinging and/or burning sensation. There may be scanty, frequent urination, and sleep may become disturbed. The digestion will continue to worsen, with bowel irritability and spasm, constipation, nausea and vomiting. There may be dizziness and/or angina, with profuse perspiration. extreme stiffness and episodic fainting (Srikantha Murthy 1995, 95).

Amavata displays many features in common with a collection of signs and symptoms that are typically diagnosed as rheumatoid arthritis (RA). The cause of RA is still the subject of some contention among pathologists, but most theories to date either advocate an autoimmune mechanism or an infectious agent. In conventional medicine RA is typically treated with non-steroidal antiinflammatory drugs (NSAIDs). corticosteroids, immunosuppressive drugs Despite the obvious similarities between it and RA, amavata is nonetheless very different. It is a distinct etiological and pathological model, based on the interaction and influence of the three *doshas*, vis. *Vata*, *Pitta* and *Kapha*. regimen Thus the treatment different for *amavata* has entirely objectives than those of RA, and very likely different results. Amavata may also include the features of other conditions. from the more pedestrian diagnoses such as arthritis, to more 'exotic' conditions such as ankylosing spondylitis.

Keywords: *Amavata*, RA, rheumatoid arthritis, *vatavyadhi*

Introduction:

History of Amavata

The basic clinical features of *amavata* have been recognized in Avurvedic medicine for thousands of years. The two oldest texts on Ayurvedic medicine, the Sushruta (11 cent. BCE?) and Charaka (8 cent. BCE) Samhitas, often refer to symptoms such as joint pain and swelling as diagnostic features of other syndromes, as *jvara* (fever), *rajayaksman* such (tuberculosis) and arsa (piles). Both of these texts however also describe features of *amavata* in syndrome а called *vatavyadhi*, a diverse group of symptoms that are organized according to the systemic and local manifestations dosha. According of Vata to the *Charaka's* description of vatavyadhi, when *Vata* affects the bones (*asthi*) there is painful swelling of the joints and immobility. The Sushruta Samhita adds that along with the inflammation. immobility and pain, the joints in vatavvadhi disorders eventually become deformed (Kumar 1997, 117-8). Much later another condition called *vatashonita* is described in the Ashtanga Hrdava (6 cent. CE) that has clinical features in common with amavata, but a different etiology. As a distinct clinical entity in and of itself amavata was only described in the 7 century CE in a diagnostic manual called the Madhava Nidanam (1).

The term *amavata* is a combination of two Sanskrit words: Vata and ama. Vata is one of the three doshas, or humors, in the Avurvedic humoral theory. The etymological origin of Vata is the Sanskrit word 'va.' which means 'to move.' the Thus Vata is prime mover and motivator in the human body, responsible for the initiation of every kind of activity. The other

two doshas (Pitta and Kapha) are said to be 'lame' without the motivating force 'seat' (sthana) of of *Vata*. The primary influence of Vata in the body is the area located from the umbilicus downwards, primarily in the large intestine. Here Vata allows for the proper elimination of wastes, which allows the body to function unobstructed.

Of the three doshas, Vata is the most subtle and easily affected. As the principle 'mover' of the body, Vata is usually the force behind any kind of physiological dysfunction. Vata is identified in the body by its cold (shita), light (laghu) and dry (ruksha) qualities. The very nature of *Vata* is therefore opposite to that of life, and the natural warmth (e.g. Pitta), heaviness and moistness (e.g. *Kapha*) of the human body.

Ama is a Sanskrit word that means 'undigested food.' As a concept, it arises out of an understanding that when the digestive fire (*jatharagni*) is weak, the result is the accumulation of 'uncooked food' in the stomach. This 'uncooked food' then enters into systemic circulation and into the various bodily tissues (dhatus), slowing, impairing or impeding a variety of metabolic processes. As *ama* accumulates further. the doshasbegin to become vitiated, and manifest as local or systemic symptoms.

Etiology of Amavata

The clinical description of a given disease and its cause (etiology) evolves from a within of study Ayurvedic branch medicine called nidana, a Sanskrit term 'causes.' In meaning practice however, nidana not only concerns itself with the etiology of disease but with its description (pathogenesis), and the clinical tools used to assess the patient. Unlike western medicine, Ayurveda will often advocate a variety of causes for a given condition, and in many cases these causes are a similar assortment of psychospiritual, dietary. and environmental lifestyle. factors.

The *Madhava Nidana* provides three basic causes of *amavata*:

•Weakness of digestion

The digestive fire is one of the prime motivators of all human function, ensuring the proper absorption and metabolism of nutrition. When digestion is impaired *ama*accumulates,

the *doshas* become vitiated, and the vitality (*ojas*) diminishes.

•Incompatible foods in the diet, including:

-foods consumed out of season or without respect to local bioclimactic factors

-unfamiliar foods (*asatmya*, i.e. opposite of the healthy norm (*satmya*), nontraditional foods -spoilt and contaminated foods, food additives, refined flour, feed-lot and farmed meat, etc.

•Lack of physical activity

Physical exercise (*vyama*) is considered an important aspect to *dinacharya*, the daily regimen recommended in Ayurveda. A lack of exercise predisposes one to *amavata*because the circulation of blood to and the removal of wastes (*ama*) from the periphery is impaired. *Pathogenesis of* Amavata

The Madhava Nidanam states that when the digestive fire is weak and ama is allowed to accumulate, it moves to the different locations of Kapha in the body (Srikantha Murthy 1995, 95). Kapha is Sanskrit derived from the root word 'shlesh, 'which means 'to embrace.' Thus, Kapha binds the joints together, nourishing and protecting the articular surfaces (Srikantha Murthy 1994, 169). *Kapha* is heavy (guru), Like *ama*. moist (snigdha) and cold (shita) in nature. For this reason, *amatypically* associates with Kapha before the other two doshas. According to the Madhava Nidanam, when *ama* is allowed to accumulate in the joints they become congested with a "...hard, waxy material" (i.e. Kapha). Soon the circulatory supply channels (dhamanis) that these regions become congested as well. Eventually this blockage affects the heart (hrdaya), which then becomes the "...seat of the disease" (2) (Srikantha 1995, 95). Once Kapha has Murthy vitiated become the other doshaseventually become involved. To restore homeostasis the body will initiate local inflammatory processes (i.e. Pitta) in the joint in order to 'cook' the accumulated *ama*. Despite the inflammatory component of this condition however, the hallmark of amavata is the progressive pathological influence of Vata in the synovial joints, and the resultant joint degeneration.

The pathogenesis of *amavata* bears some similarity to the recently described intestinal permeability syndrome (IPS). The impetus for IPS is a process by which some agent or combination of agents initiates an inflammatory response in the digestive tract. Persistent gastrointestinal inflammation eventually disrupts the integrity of the mucosal lining of the gut, and tiny perforations allow for molecules larger than usual to pass across this barrier, including molecules from dietary protein and fats, bacteria, parasites and fungi. In response to this infiltration, an immune response is initiated and the body begins to manufacture specific antibodies to these antigens. Unfortunately, many tissues have antigenic sites almost identical to those substances that pass across a permeable intestinal wall. Once activated, these antibodies then circulate and 'look' for more antigens. When an antigen is found, such as a tissue that has similar markers to an exogenous antigen, the antibody initiates an immune response and the tissue begins to be destroyed (Galland 1993). The differences between IPS obviously and *amavata* are significant, with each using an entirely different physiological model. Nonetheless if we can translate the antigens described in IPS the *ama* identified into in Ayurvedic medicine. the two models become strikingly congruent (Kumar 1997, 94). Although *amavata* is primarily a disease of Vata, it is differentiated into three basic subtypes, namely, Vata, Pitta, and Kapha. This differentiation allows the practitioner to identify a greater range of subtlety within the diagnosis and treatment of *amavata*. Where *Pitta* is involved the joints appear red and feel hot, and the patient complains of a burning, searing pain. With Vata the pain is severe, and migrates from place to place. With Kapha the pain is less, but there is more stiffness and immobility, often combined with sensations of itching. There may also be a combination of any two or

three

of

the *doshas*. If

one *dosha* is

involved the condition is said to be easy to cure. With two *doshas* the situation is more difficult, and with all three *doshas* in a state of vitiation the condition is said to be incurable. Similarly, when there is migrating pain and severe inflammation in the joints of the hands, feet, head, heels, waist, knees and thighs, *amavata* is said to be incurable (Srikantha Murthy 1995, 95-96).

Treatment of Amavata

The basic approach in the treatment of *amavata* is three-fold:

- 1. support and enhance digestion (*agni*)
- 2. facilitate the removal of *ama* from the body
- 3. repair damaged tissues and restore vitality (*ojas*)

To achieve these therapeutic objectives a variety of approaches are taken. The diet should be light and easy to digest, with plenty of pungent and bitter tastes to enkindle digestion and promote the removal of ama. Such measures include the use of herbs such as ginger and garlic in the preparation of food, as well as plenty of leafy green vegetables and other whole foods.Heavy, congesting and cold foods are to be avoided in most cases, including dairy, flour, pork, and other greasy or sticky foods that tend to promote congestion. In many cases there are specific items that need to be avoided by a particular individual, such as the solanaceous vegetables like tomatoes and peppers, certain cereals such as wheat or corn, or legumes such as peanuts or soy. down-regulate the To inflammatory response and support tissue restoration a number of nutritional supplements can be used alongside more traditional Ayurvedic therapies, such as halibut liver oil, vitamins C. chelated В and and multimineral supplements.

Ayurvedic Traditionally, medicine employs a method called pancha karma in of amavata. Pancha treatment the karma involves the usage of a variety of eliminative (shodhana) therapies, such as purgation (virechana), emesis (vamana), enemata (vasti) and errhines (nasva). These treatments are performed very carefully in an in-patient facility over a period of weeks. Prior to the application of these therapies, a variety of preparatory treatments (purva karmas) are utilized to prepare the body for pancha karma. These typically include a combination of oleation massage, snehana) and (oil sudation (diaphoretic, svedana) therapies.

While *pancha* karma is certainly considered the most effective treatment for *amavata*, a combination of eliminative (*shodhana*) and

palliative (*shamana*) treatments can be employed on an out-patient basis to good effect, and has several practical advantages over *pancha karma*.

Although karmas such the *purva* as *snehana* and *svedana* are typically used prior to pancha karma, they may also be used at the onset of treatment and on an ongoing basis in an out-patient regimen. Patients can be encouraged to apply medicated oils prescribed (e.g. kottamchukkadi taila, brhat saindhavadya taila) to the affected joints or the entire body. These areas are then treated with some kind of sudation therapy, locally, such as heated sand in a linen bag or a hot water bottle, or more generally, as in a sauna or steam bath. Similarly, medicated salves, fomentations and poultices may also be applied locally help facilitate the removal to of ama. Typical herbs used in such external preparations include pungent and bitter tasting herbs with rubefacient and decongestive properties, such as dhattura (Datura stramonium), shunthi (Zingiber haritaki (Terminalia officinalis), chebula), kushta (Saussurea lappa), rasna roxburghii) and vacha (Acorus (Vanda Important oils (taila) used calamus).

in *amavata* are castor and sesame oil. Other medicaments for external usage include *saindhava* (rock salt) and *dadhi* (milk curd). Care must be taken when applying some of these agents in periods of exacerbation, as they can make the symptoms much worse, even while assisting the removal of *ama*.

Another therapy used in *pancha* karma that has application on a clinical basis is enemata (vasti) therapy. The fluid used in enematavaries according to the specifics of each case, sometimes alternating between aqueous-based and oil-based medicated preparations. In most relatively common recipe cases a called madhutailika will suffice:

- 600 mL eranda mula (Ricinus communis root) decoction
- 230 grams each of honey and sesame oil
- 15 grams of *saindhava* (rock salt)
- 30 grams of *shadhakva churnam* (*Anethum graveolens* powdered herb).

The ingredients are mixed together well in a mortar, the heat from the *eranda* decoction allowed to cool to a suitable temperature before application.

Enema therapy is considered very important in *amavata*, as the colon is both the seat of *Vata*, and the primary organ that eliminates *ama* from the body.

Regular exercise is another important aspect in treatment of *amavata*, to enhance the circulation to and away from the affected areas and thereby facilitate the removal of ama. The Ashtanga Hrdaya recommends that one should exercise to one-half one's strength in the winter and spring, and to a lesser extent in summer and fall, until sweat appears on the forehead and in the axilla (Srikantha Murthy 1994, 25). gentle exercises such Typically, as simple *hatha* walking or *yoga* positions (asanas) are recommended at the outset of treatment. Depending on the severity of the condition however, the person would do well not to engage in any strenuous exercise that might damage the joints further.

Internally a variety of therapies may be recommended. from simples such as shunthi(Zingiber officinalis), to complex polyherbal formulations such as yogaraja guggulu, to specially purified mineral preparations (rasa) such as maha lakshmivilasa rasa. The commonality between these various remedies is their specific ability to assist in the removal of ama. Many of these herbs and formulas have additional properties, such as a special ability to reduce pain and inflammation, treat infection, or restore function to a damaged tissue or organ. The following are five botanicals that are commonly used in the treatment of amavata, chosen somewhat randomly from an enormous number of possibilities.

Botanicals

Guggulu - Commiphora mukul

Guggulu is the oleo-gum resin derived from the stem bark of Commiphora mukul, a shrubby tree with spiny branches found throughout the subcontinent of western Asia. Arabia and eastern Africa. On the subcontinent of India it is found in dry sub-tropical areas such as the Sindh, Baluchistan, and the Deccan plateau. The leaflets are arranged in clusters of one to three, the lateral leaflets typically half the size of the terminal ones. Similar species include C. molmol and C. myrrha. When fresh the oleo-gum resin is moist, fragrant and golden in colour. It burns in fire, melts in the sun and when mixed with hot water milky forms а emulsion. Guggulu is widely used in Ayurvedic medicine as an excipient, utilized in the manufacture of pills and pastes (Kirtikar and Basu 1935, 527; Nadkarni 1954, 167; Varier 1996, 164).

Guggulu gum has a bitter (*tikta*), pungent (*katu*) and

astringent (kashaya) taste (rasa), and а hot (ushna), dry (ruksha) and light (laghu) energy (virya). It is used in amavata to enkindle digestion (dipanapachana) and raise the body heat to the periphery (swedana). Guggulu acts to 'dry up' (soshana) and diminish (langhana)the accumulated ama in the channels (srotas). *Guggulu* is indicated in vitiations of Vata and Kapha, but is typically contraindicated in *Pitta* conditions. Fresh guggulu is considered to be anabolic and nourishing (brimhana), whereas old guggulu is considered depleting (lekhana). In the treatment of amavata, guggulu is typically combined with other ingredients in a formula called yogaraja guggulu (Dash 1991, 179; Nadkarni 1954, 167; Varier 1996, 164). Guggulu has been the subject of fairly extensive scientific research, and has been shown to exhibit antihelminthic, antiobesity. antioxidant and hypocholesterolemic effects in human clinical trials (Sheir et al 2001; Bhatt et al 1995; Singh et al 1994; Nityanand et al

1995, Shigh et al 1994, Nityalahd et al 1989). In experimental models, *guggulu* has displayed antitumor, antifungal, anti-inflammatory, musclerelaxing, cardioprotective, hypotensive, bradycardiac and thyrotropic effects (Abdul-Ghani and Amin 1997; al-Harbi et al 1997; Andersson et al 1997; Qureshi et al 1994; Lata et al 1991; Tariq et al 1986; Tripathi et al 1984; Sarbhoy 1978).

Haritaki - Terminalia chebula

Haritaki is the fruit of *Terminalia chebula*, a large deciduous tree found throughout India in both forested areas and dry slopes, from the Himalayas to southern India. The young leaves are covered in rust-coloured hairs that become glabrous when mature, and are rounded at the base, elliptic to oblong, tip acute. The small flowers are dull-white to yellowish in color, arising in simple spikes or panicles. The mature fruits are shiny, glabrous, yellow to orange-brown drupes. Both the immature and mature fruits are used medicinally (Kirtikar and Basu 1935, 1020-21; Varier 1996, 263).

Haritaki is perhaps the most important medicinal plant in the Ayurvedic materia medica, the first among plants listed in every *nighantu*. *Haritaki* possesses all tastes with the exception of salty (*lavana*). The post-digestive effect (*vipaka*) is sweet (*madhura*), and the energy (*virya*) is

hot (ushna), light (laghu) and

dry (*ruksha*).Similar to *guggulu, haritaki* is used to enkindle digestion (*dipanapachana*), and

remove ama by virtue of its dry and light properties. Haritaki is also used as a rejuventative (rasavana), taken on a daily basis to ward of old age and disease. It vitiation reduces the of all three doshas (humors), and is considered to strengthen the mind and nervous system (medhya rasayanam). The unripe fruit is more astringent and less purgative than the mature fruit, and is used in the treatment of dysentery and diarrhea (Dash 1991, 8; Nadkarni 1954, 1207; Varier 1996, 263).

Categorized by many as a simple purgative, *haritaki* has yet to stimulate much interest in the scientific community. Haritaki has been shown to have antimicrobial. antitumor and hypocholesterolemic effects in experimental models (Sohni et al. 1995; Phadke and Kulkarni 1989; Dutta et al 1998; el-Mekkawy et al 1995; Creencia et al 1996; Thakur et al 1988).

Guduchi – Tinospora cordifolia

Guduchi is a large deciduous perennial climber found throughout tropical regions of the subcontinent of India and south-east Asia. It has large succulent stems and papery bark; the leaves are glabrous and cordate; the male flowers yellow-white and clustered; the females flowers yellowwhite and solitary. As it climbs upwards *T*. *cordifolia* sends down long, pendulous fleshy roots. The stem is used medicinally (Kirtikar and Basu 1935, 77-78; Varier 1996, 283).

Guduchi is bitter (tikta), astringent (kashaya) and sweet (madhura) in taste (rasa). The post-digestive effect (vipaka) is sweet (madhura), and the energy (virya) is light (langhana) and mildly warming (ushna). Like many herbs that are used in treatment of amavata, the guduchi enkindles the digestive fire (dipana) and promotes the removal of ama through its light (langhana) properties. Although classified in many nighantus as warming in energy, the balance between its bitter and sweet tastes makes guduchi specific to disorders and deficiencies of the liver. blood, and skin, and to reduce the vitiations of Pitta. Guduchi is often used along with circulatory stimulants such as Zingiber officinalis in the treatment of conditions such as amavata, to reduce the symptoms of inflammation and pain. Taken with ghee guduchi is used to treat Vattic conditions, with jaggery for Pitta, and with castor oil for vitiated Kapha (Dash 1991, 14; Dash and Junius 1983, 139; Kirtikar and Basu 1935, 78; Varier 1996, 283).

Guduchi has demonstrated immunosupportive and hepatoprotective effects in human clinical trials (Rege et al 1993). Experimental models for Tinospora cordifolia and similar species have demonstrated adaptogenic, antioxidant, hypolipidemic, antimalarial. hepatoprotective and immunomodulatory effects (Prince et al 1999; Prince and Menon 1999, Rege et al 1999; Najib et al 1999; Thatte et al 1994; Bhattacharya et al 1991).

Ashwagandha - Withania somnifera

Ashwagandha is the root of Withania somnifera, an erect undershrub found in the drier parts and wasteland of the Indian subcontinent, the Middle East and Africa. It attains a height of between 0.3-1.5 meters, with simple, alternate ovate leaves up to 10 cm in length. The flowers are green to yellow, borne on short axillary clusters, fruits spherical, orange-red when mature, enclosed within the inflated calyx (Kirtikar and Basu 1935, 1774; Mills and Bone 2000, 596; Varier 1996, 409).

a

mildly

Ashwagandha has

bitter (tikta) and astringent (kashava) taste (rasa). The postdigestive effect is sweet (madhura) and the energy (virya) is warm (ushna) and heavy (guru) in nature. Although *Ashwagandha* has applications in many disorders it is considered a specific for mitigating Vata. Ashwagandha is an important herb in the treatment of *amavata*, although it has a different application and mode of action than some of the other herbs already discussed in this paper. Ashwagandha is reasonably contraindicated in *amavata* because of its anabolic (brimhana) effects in the body, a property that would tend to facilitate the production of ama. Ashwagandha should probably not be used at the outset of treatment. but only after much of the ama has already been removed. Its best application is as a rejuvenative (rasayana). used to restore damaged tissue, and as a palliative (shamana) in severe conditions. Ashwagandha has a special restore ability to the nervous system (medhya rasayana), relieve pain (vedanasthapana), and promote sleep. Ashwagandha has demonstrateda hypoglycemic, diuretic and hypocholesterolemic effect in human clinical trials (Andallu and Radhika 2000). combination with Boswellia In serrata stem, Curcuma longa rhizome and zinc complex (Articulinа F), Ashwagandha was shown to have a significant effect in the reduction of pain, in a randomized, double-blind, placebo controlled, cross-over study in patients with osteoarthritis (Kulkarni et al 1991). Experimentally Withania has demonstrated adaptogenic, anti-inflammatory,

antioxidant, antitumor, chemoprotective, gabanergic, analgesic, and immunomodulant effects in animal models (Archana and Namasivayam 1999; Davis and Kuttan 1998; Dhuley 1998; Bhattacharya et al 1997; Kulkarni and Ninan 1997; Devi 1996; Ziauddin et al 1996; al-Hindawi et al 1992, 1989; Mehta et al 1991).

Bhallataka - Semecarpus anacardium

Bhallataka is Semecarpus anacardium, the cashew tree. Although some sources indicate that this moderate sized semideciduous tree was brought to India from South America by the Portuguese, Bhallataka is clearly mentioned both in the Sushruta and Charaka samhitas, texts which antedate the Portuguese by more than a millennia (Varier 1996, 98). S. anacardium is now cultivated all over the world as a food, in moist tropical forests, and in the subcontinent ranging from the sub-Himalayas and Assam in the north, to the coast of Kerala in the south.. The grey bark exfoliates in small irregular flakes, and the leaves are simple, alternate, obovate-oblong, rounded at the apex, glabrous above and pubescent below. The greenish fruits are ovoid to oblong drupes that are attached to a swollen, fleshy receptacle that sits below it and turns vellow when ripe (Kirtikar and Basu 1935, 667; Varier 1996, 98). The toxic pericarp is a by-product of the cashew industry, where special care is taken to remove it from the cashew kernel. The juice of the pericarp is vesicant and eschariotic, used at one time in India as a marking ink (Nadkarni 1954, 1120-21). The sap of the tree has also been shown to be quite toxic, with one reported case in the literature of severe dermatitis, anuria, and renal cortical necrosis from skin exposure (Matthai and Date 1979). The whole fruit and pericarp are used medicinally.

Bhallataka fruitispungent (katu),bitter (tikta),astringent (kashaya) and

sweet (*madhura*) in taste. It has a sweet (*madhura*) post-digestive

effect (*vipaka*), and a light (*laghu*) and hot (*ushna*) energy (*virya*) (Dash 1991, 99; Nadkarni 1954, Varrier 1996, 98). The Ashtanga Hrdaya (7 cent CE) considers bhallataka fruit to be "...like fire in property, increasing intelligence and effectively

mitigating Vata and Kapha(Srikantha

Murthy 1994, 100). Bhallataka has long been considered an important remedy in the treatment of a variety of complaints including rheumatism, liver disorders and hemorrhoids, considered equal to mercury in action (Nadkarni 1954, 1120). The pericarp however contains a variety of toxic principles that can precipitate a skin rash and renal failure if the dose is too large or if the remedy is prepared incorrectly. Some individuals appear to be more sensitive to *bhallataka* than others. the many preparations Among that contain *bhallataka* is a rasavana mentioned by the Chakradatta(12 cent

CE)called amritabhallataka. In the preparation of this remedy bhallatakafruit is first boiled in water, the fruits removed, and the decoction mixed and cooked with milk, ghee and jaggery (Sharma 2002, 648). The Chakradatta states that this is "king preparation the of all rasayanas," and may be used on an ongoing basis to promote strength and longevity (Sharma 2002, 648). Toxicological studies for a milk extract of bhallataka called serankottai nei indicated that when processed correctly bhallataka is remarkably welltolerated (Vijayalakshmi et al 2000). Bhallataka has displayed anticarcinogenic,

neuroprotective, antioxidant, antiinflammatory, cytotoxic, and hypocholesterolemic effects in experimental models (Shukla et al 2000; Premalatha and Sachdanandam 2000: Premalatha and Sachdanandam 1999; Vijavalakshmi et al 1997; Sharma et al 1995; Smit et al 1995).

Yogaraja Guggulu

Although individual herbs and simple formulations are used in Ayurvedic medicine, more often herbal medicine takes the form of complex polyherbal formulations. Among the most important of these formulas in Ayurvedic medicine is *yogaraja guggulu* the recipe here taken from the *Chakradatta*:

Reduce to a fine powder and combine in equal weight): chitraka parts (by (Plumbago zeylanica root), pippalimula (Piper *longum* root), vavani (Trachyspermum *ammiseed*), Krishna jiraka (Carum *carvi* seed), vidanga (Embelia ribes fruit), ajamoda (Carum roburghianum fruit), devadaru (Cedrus *deodara* heart wood), chavya (Piper chaba stem), ela (Elettaria cardamomum fruit), saindhava (rock salt), kushta (Saussurea lappa root), rasna (Vanda roxburghii leaf and root), gokshura (Tribulus terrestris fruit). dhanyaka (Coriandrum sativum fruit), *triphala* (equal parts Terminalia chebula fruit, Terminalis bellerica fruit, Emblica officinalis fruit), musta (Cyperus rotundus rhizome). trikatu (equal parts *Piper longum* fruit, Piper nigrum fruit, Zingiber officinalis rhizome), tvak (Cinnamomum *zeylanicum* stem bark), ushira (Vetiveria zizanioides root), *vavakshara* (an alkaline preparation derived from Hordeum vulgare), patra (Cinnamomum

tamala leaf) and *talishapatra* (Abies webbiana leaf). Add a quantity of guggulu equal to the weight of the above combined ingredients, and then slowly add enough ghee while mixing in a mortar to form a soft mass.

The *Chakradatta* states that *yogaraja guggulu* is a medicine like "...nectar," useful in the treatment of *amavata*, and other conditions such as helminthiasis, chronic ulcers, splenomegaly, and

piles. *Yogaraja guggulu* "stimulates the digestive fire, promotes energy and strength, and overcomes *vatika* (*Vata*) disorders even if located in the joints and marrow" (Sharma 2002, 250). The Ayurvedic Formulary of India suggests a recommended dose of 3 grams (1978, 58-59).

Conclusion:

The signs and symptoms typically associated with rheumatoid arthritis are clearly mentioned throughout the Ayurvedic literature. If one conducts a survey of the most important texts, these symptoms are found grouped together in a variety of different ways, sometimes with other symptoms, and given different names. All of these conditions are clinically relevant in Ayurvedic medicine, but amavata specifically is most often correlated with rheumatoid arthritis. Unlike the typical approaches used in Western medicine, Ayurveda employs a variety of sophisticated holistic treatment methods to decrease joint pain and inflammation, and restore the joints to a degree of normalcy. While not all, many of the botanicals used by Avurvedic herbalists in the treatment of amavata have undergone a significant degree of scientific investigation. The vast majority of these studies, comprised of both human clinical trials and experimental models, have validated traditional Ayurvedic herbology in the treatment of inflammatory joint disorders.

References:

 Abdul-Ghani AS, Amin R. 1997. Effect of aqueous extract of Commiphora opobalsamum on blood pressure and heart rate in rats. J Ethnopharmacol 1997 Aug;57(3):219-22

- al-Harbi MM, Qureshi S, Raza M, Ahmed MM, Afzal M, Shah AH. 1997. Gastric antiulcer and cytoprotective effect of Commiphora molmol in rats. J EthnopharmacolJan;55(2):141-50
- al-Hindawi, M.K., S.H. Al-Khafaji, and M.H. Abdul-Nabi. 1992, Antigranuloma activity of Iraqi Withania somnifera. J Ethnopharmacol. Sep; 37(2):113-6
- 4. al-Hindawi, M.K., I.H. Al-Deen, M.H. Nabi, and M.H. Ismail. 1989. Anti-inflammatory activity of some Iraqi plants using intact rats. *J Ethnopharmacol*. Sep; 26(2):163-8
- Andersson M, Bergendorff O, Shan R, Zygmunt P, Sterner O. 1997. Minor components with smooth muscle relaxing properties from scented myrrh (Commiphora guidotti). *Planta Med* Jun;63(3):251-4
- 6. Andallu B, Radhika B. 2000. Hypoglycemic, diuretic and hypocholesterolemic effect of winter cherry (Withania somnifera, Dunal) root.*Indian J Exp Biol* Jun;38(6):607-9
- Archana, R. and A. Namasivayam. 1999. Antistressor effect of Withania somnifera. J Ethnopharmacol. Jan; 64(1):91-3
- 8. Berkow, Robert ed. et al. 1992. *The Merck Manual of Diagnosis and Therapy*. 17th ed. Rahway, NJ: Merck and Co.
- Bhatt AD, Dalal DG, Shah SJ, Joshi BA, Gajjar MN, Vaidya RA, Vaidya AB, Antarkar DS. 1995. Conceptual and methodologic challenges of assessing the shortterm efficacy of Guggulu in obesity: data emergent from a naturalistic clinical trial. *J Postgrad Med*Jan-Mar;41(1):5-7
- 10. Bhattacharya, S.K., K.S. Satyan and S. Ghosal. 1997. Antioxidant activity of glycowithanolides

from *Withania somnifera*. *Indian J Exp Biol*. Mar; 35(3):236-9

- 11. Bhattacharya SK, Maity PC, Mediratta P, Sen P. 1991. Modulation of humoral immune responses by Tinospora malabarica in experimental animals. *Indian J Exp Biol*Oct;29(10):971-2
- 12. Creencia, E., T. Eguchi, T. Nishimura, and K. Kakinuma. 1996. Isolation and Structure Elucidation of the Biologically Active Components of *Terminalia chebula* Retzius

(Combretaceae). KIMIKA 12:1-10

- 13. Dash, Bhagwan. 1991. Materia Medica of Ayurveda. New Delhi: B. Jain Publishers.
- 14. Dash, Bhagwan and Manfred Junius. 1983. *A Handbook of Ayurveda*. New Delhi: Concept Publishing.
- 15. Davis L. and G. Kuttan. 1998. Suppressive effect of cyclophosphamide-induced toxicity by *Withania somnifera* extract in mice. *J Ethnopharmacol*. Oct; 62(3):209-14
- 16. Devi, P.U. 1996. Withania *somnifera* Dunal (Ashwagandha): source potential plant of а promising drug for cancer chemotherapy and radiosensitization. Indian J Exp Biol.Oct; 34(10):927-32
- 17. Dhuley, J.N. 1998. Effect of Ashwagandha on lipid peroxidation in stress-induced animals. J Ethnopharmacol. Mar; 60(2):173-8
- Dutta, B.K., I. Rahman, and T.K. Das. 1998. Antifungal activity of Indian plant extracts. *Mycoses*. Dec; 41(11-12):535-6
- 19. el-Mekkawy, S *et al.* 1995. Inhibitory effects of Egyptian folk medicines on human immunodeficiency virus (HIV) reverse transcriptase. *Chem Pharm Bull* Apr; 43(4):641-8

- 20. Galland, Leo. 1995. Leaky Gut Syndromes: Breaking the Vicious Cycle. Towsend Letter for Doctors. Aug/Sep: 62-68
- 21. India. Ministry of Health and Family Planning. 1978. *The Ayurvedic Formulary of India*. Part 1. 1 ed. Delhi
- 22. Kirtikar KR and BD Basu. 1935. *Indian Medicinal Plants*. 2 ed. Vol. 1-4. Reprinted in 1993. Delhi: Periodical Experts.
- 23. Kulkarni RR, Patki PS, Jog VP, Gandage SG, Patwardhan B. 1991. Treatment of osteoarthritis with a herbomineral formulation: a double-blind, placebo-controlled, cross-over study. J *Ethnopharmacol.* May-Jun;33(1-2):91-5
- 24. Kulkarni, S.K. and I. Ninan. 1997. Inhibition of morphine tolerance and dependence by *Withania somnifera* in mice. *J Ethnopharmacol*. Aug; 57(3):213-7
- 25. Lata S, Saxena KK, Bhasin V, Saxena RS, Kumar A, Srivastava VK. 1991 Beneficial effects of Allium sativum, Allium cepa and Commiphora mukul on experimental hyperlipidemia and atherosclerosis - a comparative evaluation. J Postgrad MedJul;37(3):132-5
- 26. Matthai TP, Date A. 1979. Renal cortical necrosis following exposure to sap of the marking-nut tree (Semecarpus anacardium). *Am J Trop Med Hyg* Jul;28(4):773-4
- 27. Mehta, A.K., P. Binkley, S.S. Gandhi, and M.K. Ticku. 1991.
 Pharmacological effects of Withania somnifera root extract on GABAA receptor complex. Indian J Med Res. Aug; 94:312-5
- 28. Mills, Simon and Kerry Bone. 2000. Principles and Practice of Phytotherapy.London: Churchill Livingstone

- 29. Nadkarni, K.M. and A.K. Nadkarni. 1954. *Indian Materia Medica*. Vol 1. 3 rev. Mumbai (Bombay): Popular Prakashan Private Ltd.
- Najib Nik A Rahman N, Furuta T, Kojima S, Takane K, Ali Mohd M. 1999. Antimalarial activity of extracts of Malaysian medicinal plants. J

EthnopharmacolMar;64(3):249-54

- 31. Nityanand S, Srivastava JS, Asthana OP. 1989. Clinical trials with gugulipid. A new hypolipidaemic agent. J Assoc Physicians India. May;37(5):323-8
- 32. Phadke, S.A., and S.D. Kulkarni. 1989. Screening of in vitro antibacterial activity of *Terminalia chebula*, *Eclipta alba* and *Ocimum sanctum*. *Indian J Med Sci*. May; 43(5):113-7
- 33. Premalatha B, Sachdanandam P.
 2000. Potency of Semecarpus anacardium Linn. nut milk extract against aflatoxin B(1)-induced hepatocarcinogenesis: reflection on microsomal biotransformation enzymes. *Pharmacol Res* Aug;42(2):161-6
- 34. Premalatha B, Sachdanandam P. 1999. Semecarpus anacardium L. nut extract administration induces the in vivo antioxidant defence system in aflatoxin B1 mediated hepatocellular carcinoma. *J Ethnopharmacol* Aug;66(2):131-9
- 35. Prince PS, Menon VP. 1999. Antioxidant activity of Tinospora cordifolia roots in experimental diabetes *J* Ethnopharmacol 1999 Jun;65(3):277-81
- 36. Qureshi S, al-Harbi MM, Ahmed MM, Raza M, Giangreco AB, Shah AH. 1993. Evaluation of the genotoxic, cytotoxic, and antitumor properties of Commiphora molmol using normal and Ehrlich ascites carcinoma cell-bearing Swiss

albino mice. *Cancer Chemother Pharmacol* 33(2):130-8

- 37. Rege NN et al. 1999. Adaptogenic properties of six rasayana herbs used in Ayurvedic medicine. *Phytother Res* Jun;13(4):275-91
- 38. Rege N, Bapat RD, Koti R, Desai NK, Dahanukar S. 1993.
 Immunotherapy with Tinospora cordifolia: a new lead in the management of obstructive jaundice. *Indian J Gastroenterol* Jan;12(1):5-8
- 39. Sarbhoy AK, Varshney JL, Maheshwari ML, Saxena DB.
 1978. Efficacy of some essential oils and their constituents on few ubiquitous molds. *Zentralbl Bakteriol*[Naturwiss] 1978;133(7-8):723-5
- 40. Sharma A, Mathur R, Dixit VP. 1995. Hypocholesterolemic activity of nut shell extract of Semecarpus anacardium (Bhilawa) in cholesterol fed rabbits. *Indian J Exp Biol*Jun;33(6):444-8
- 41. Sharma, P.V. 2002. *Cakradatta* (*Sanskrit text with English translation*). 3 ed. Varanasi: Chaukhamba Orientalia.
- 42. Shukla SD, Jain S, Sharma K, Bhatnagar M. 2000. Stress induced neuron degeneration and protective effects of Semecarpus anacardium Linn. and Withania somnifera Dunn. in hippocampus of albino rats: an ultrastructural study. *Indian J Exp Biol* 2000 Oct;38(10):1007-13
- 43. Singh RB, Niaz MA, Ghosh S. 1994. Hypolipidemic and antioxidant effects of Commiphora mukul as an adjunct to dietary therapy in patients with hypercholesterolemia. *Cardiovasc Drugs Ther* 1994 Aug;8(4):659-64
- 44. Sohni, Y.R., P. Kaimal, and R.M. Bhatt. 1995. The antiamoebic effect of a crude drug formulation

ofherbalextractsagainst Entamoebahistolytica invitroandinvitrosinthnopharmacol.Jan; 45(1):43-52

- 45. Smit HF, Woerdenbag HJ, Singh RH, Meulenbeld GJ, Labadie RP, Zwaving JH. 1995. Ayurvedic herbal drugs with possible cytostatic activity. J *Ethnopharmacol* Jul 7:47(2):75-84
- 46. Srikantha Murthy K.R. 1996. Madhava Nidanam (Roga Viniscaya of Madhavakara). 2 ed. Varanasi: Chaukhamba Orientalia.1994.
- 47. Vagbhata's Ashtanga Hrdayam. Vol 1. Varanasi: Krishnadas Academy
- 48. Prince P, Menon VP, Gunasekaran G. Stanely Mainzen. 1999. Hypolipidaemic action of Tinospora cordifolia roots in alloxan diabetic rats. J Ethnopharmacol. Jan;64(1):53-7
- 49. Tariq M, Ageel AM, Al-Yahya MA, Mossa JS, Al-Said MS, Parmar NS. 1986. Antiinflammatory activity of *Commiphora molmol. Agents Actions* 17(3-4):381-2
- 50. Thakur, C.P., B. Thakur, S. Singh, P.K. Sinha, and S.K. Sinha. 1988. The Ayurvedic medicines Haritaki, Amala and Bahira reduce cholesterol-induced atherosclerosis in rabbits. *Int J Cardiol*. Nov; 21(2):167-75.
- 51. Thatte UM, Rao SG, Dahanukar SA. 1994. Tinospora cordifolia induces colony stimulating activity in serum. *J Postgrad Med*. Oct-Dec;40(4):202-3
- 52. Tripathi YB, Malhotra OP, Tripathi SN. 1984. Thyroid stimulating action of Z-guggulsterone obtained from Commiphora mukul. *Planta Med* Feb;(1):78-80
- 53. Varier, V.P.S. 1996. Indian Medicinal Plants: A Compendium of 500 species. Edited by PK

Warrier, VPK Nambiar and C Ramankutty. vol 5. Hyderabad: Orient Longman.

- 54. Indian Medicinal Plants: A Compendium of 500 species. Edited by PK Warrier, VPK Nambiar and C Ramankutty. vol 4. Hyderabad: Orient Longman, 1995b.
- 55. 1995a. Indian Medicinal Plants: A Compendium of 500 species. Edited by PK Warrier, VPK Nambiar and C Ramankutty. vol 3. Hyderabad: Orient Longman.
- 56. Indian Medicinal Plants: A Compendium of 500 species. Edited by PK Warrier, VPK Nambiar and C Ramankutty. vol 2. Hyderabad: Orient Longman, 1994b.
- 57. Indian Medicinal Plants: A Compendium of 500 species. Edited by PK Warrier, VPK Nambiar and C Ramankutty. vol 1. Hyderabad: Orient Longman, 1994a.
- 58. Vijayalakshmi T, Muthulakshmi V, Sachdanandam P. 2000. Toxic studies on biochemical parameters carried out in rats with Serankottai nei, a siddha drug-milk extract of Semecarpus anacardium nut. J Ethnopharmacol. Jan;69(1):9-15
- 59. Vijayalakshmi T, Muthulakshmi V, Sachdanandam P. 1997. Effect of

milk extract of Semecarpus anacardium nuts on glycohydrolases and lysosomal stability in adjuvant arthritis in rats. J

Ethnopharmacol Sep;58(1):1-8

60. Ziauddin, M., N. Phansalkar, P. Patki, S. Diwanay, B. Patwardhan. 1996. Studies on the immunomodulatory effects of Ashwagandha. *J Ethnopharmacol*. Feb; 50(2):69-76

Footnotes

1. About a century or so before the *Madhava* Nidanam the Ashtanga Hrdaya (6 century CE) described a condition called vatashonita (vatarakta) that is often

transliterated as gout, but bears many similarities to rheumatoid arthritis and inflammatory joint disease in general. Unlike the etiology described for *amavata*, *vatashonita* is caused by the simultaneous vitiation of *Vata* and *shonita* (blood).

2. According to the Ayurvedic understanding of physiology, the heart receives the immediate products of digestion (*rasa*) and transports it throughout the body as blood (*rakta*).

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