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INTRODUCTION

Medicine has become as indispensable part of human being since ancient period of all systems of medicine, siddha system is unique since it is the one which deals with both Internal ie, Man and External body ie, Nature.

Nature has provided many plaints which are indispensable to man for his life. Also, in our siddha system of medicine, as many plants contain small amounts of minerals in them, they are first used as medicine & then used metals as medicine.

It is known from the following lines,

"வேர் பாரு தழைபாரு மிஞ்சினக் கால் மெல்ல மெல்லப் பற்பச்செந் தூரம்பாரே"

In this aspect, the author has selected a plant drug, Jatamansi, for the treatment of Eraippu Erumal as per our texts ie. Gunapadam – Mooligai (P.No - 416) (Mooligai No 266).

Jatamansi is used in siddha medicine for a long time. Spikenard or Indian nard is the common name of the plant. It's botanical name is Nardostachys jatamansi. It belongs to family valerianaceae. It is a native of himalayan region and is found to far south as the Deccan plateau India.

Jatamansi has two varieties

1. Himalayan i.e. Indian variety - N. Jatamansi

2. North American variety - F. Valeianaceae

Aralia racemosa

Family: Araliaceae

Jatamansi is used in India as drug and also in perfumery.

In this dissertation study, **Sadamanjil ver Chooramam** is taken for the study for the treatment of Eraippu Erumal.

AIM AND OBJECTIVE

The aim of this dissertation is to establish that the drug Sadamanjil Verchoornam is an effective remedy for the disease Eraippu Erumal.

Eraippu Erumal (Bronchial Asthma) is as common in India as in any other country. It is proably one fo the commenest of the major affections in our country. This forms the single biggest clinical group among the chest diseases that is treated in our college hospital. Millions of people suffer from it.

Bronchial Asthma is just an allergic manifestation in an over simplification of this difficult problem. In every case of Asthma there are a few known factors and many unknown factors, and so an aetiological classification of Asthma in any given case is difficult and other faculty.

The treatment of the disease is to be of successful has necessarily got to be individualized as no two cases of Asthma are alike either in their response to treatment or in manintaining the improvement. So the entire course of the disease is unpredictable and each case has got to be tackled by utilising all the resources both modern and Ancient.

Keeping this in mind the author selected "Sadamanjil Ver Choornam" which is not much used by the physicians.

The drug is easily available so the main aim of the study is to do a pharmacological and clinical study of **Sadamanjil Ver Choornam** on **Eraippu Erumal**.

The study is done in the following Aspects,

- 1. Botanical aspects.
- 2. Pharmacognostical aspects.
- 3. Phytochemistry.
- Pharmacological Aspects (Review of update work on Jatamansi all over the world)
- 5. Gunapadam Aspects.
- 6. Bio chemical Analysis.
- 7. Pharmacological Analysis.
- 8. Clinical Assessment.

BOTANICAL ASPECT

Nardostachys jatamansi de candolle

According to the Bentham and Hooker (1876) classification, N. Jatamansi DC is classified as follows.

Kingdom - Plant kingdom

Division - Angiosperms

Class - Dicotyledons

Sub -class - Gamopetalae

Series - Inferae

Order - Rubiales

Family - Valerianaceae

Genus - Nardostachys

Species - Jatamansi, decandolle.

Vernacular Names

Common names : Indian nard, balchar, spikenard

English names : Muskroot, Indian Spikenard

Sanskrit : Jatamansi, Mamsi

Hindi : Jatamansi, bal-chir, jatalasi

Kannada : Jatavasi, Jatamamsi

Malayalam : Jatamanci, Jatamamsi, manci

Telugu : Jatamamsi, Jatamsi

Bengali : Jatamansi

Marathi : Jatamavshi

Gujarathy : Jatamansi, Kalichhad

Kashmiri : Bhutijatt, Kukilipot

Garhwal : Masi

Nepal : Haswa, Naswa, Jatamangsi

Bhutan : Pampe, Jatamansi

Tamil : Jatamashi, (ஜடாமாஷி) Sadamanjil

(சடாமாஞ்சில்)

Geographic origin of the plant

Central Nepal.

Method of Growing

Wild.

Distribution and Habitat:

It is commonly distributed in an elevation range of 3500m to 4500m in the northern aspect of the sub-alpine and alpine pastureland of the Himalayas in Nepal. The plant is mostly found growing in steap areas with a 25° - 45° slope. It grows well on open, stony and grassy slopes and on the turf of glacial flats. Also found as far south as the Deccan plateau.

Varieties

1. Himalayan variety - N.jatamansi

Family: Valerianaceae.

The Indian variety is a Himalayan plant whose underground stem produce a perfume used in Eastern aromatic oils.

2. North American Variey - Aralia racemosa

Family: Araliaceae.

It has fragrant roots.

Habit and features

An erect perennial herb, 10 - 60cm in height, dwarf, and hairy, rhizomatous herb forming a group of thick roots covered with reddish brown fibres. Leaves opposite, spathulate. Stipules absent. Flowers capitate. Calyx epigynous. Corolla gamopetallous and tubular, stamens on the coroalla tube. Anthers 2 – celled opening lengthwise. Ovary inferior, 3 celled style simple and slender Endoperm absent.

External Morphology

Root stock

Woody, long stout, covered with fibres from petioles of withered leaves.

Stem

10 - 60cm more or less pubscent upwards, often glabrate below, subscapose .

Leaves

Entire radical leaves elongate, spathulate cauline few, Sessile, few oblong or subovate.15 – 20cm, longitudinally nerved, Glabrous or slightly pubescent narrowed into the petiole.

Flowers

Rosy, pale pink or blue. 1-30 in no. fine capitate, heads in cymes, bracts, 6mm oblong free or nearly so pubescent.

Calyx

5 lobed in fruit enlarged membranous veined.

Corolla

Tubular companulate, base sub equal, 5 lobes, spreading rosy, somewhat hariy.

Androecium

4 stamens, stamen on the corolla tube alternate with the corolla lobes.

Gynoecium:

Infereior ovary 3 celled, 1 ovuled, style linear, Slender, Stigma capitate.

Fruit:

4mm long covered with ascending white hairs, crowned by the ovate, acute dentate calyx, obovate, compressed, 3 – celled, 1 seeded, the 2 barren cells smaller than the fertile.

Seed: obovate, compressed.

Parts used: Rhizome.

Rhizome: Long stout and woody. It has an agreeable odour

with bitter aromatic taste.

GROWTH OF THE PLANT

The flowering takes place during June – July and fruiting in August -October. In the begining of October, all leaves, turn yellow and become ready for pereniation. During the winter, the herb sheds all leaves, gets buried under the snow and remains dormant with the melting of the snow in the begining of the summer, Jatamansi starts growing.

Regeneration

Natural regeneration takes place by rhizome and seeds. Jatamansi is a wild plant but is occasionally cultivated in India and china. The plant can be cultivated from the cuttings of underground parts or rhizomes as well as from seeds. The plants coming from the cuttings of rhizome grow faster than that from the seeds.

Harvesting

The appropriate time for harvesting jatamansi is october through December. Rhizomes harvested from the 2-3 year old plants give higher yield than young plants.

RHIZOME – JATAMANSI

Synonyms

Nard, Indian spikenard.

Biological name

Nardostachys jatamansi.

Biological source

Jatamansi consists of dried rhizomes of Nardostachys jatamansi DC family valerianaceae.

Geographical source

These plants are found in the alpine himalayas at an altitude of 3000 - 5000m. It is grown from punjab to sikkim and in Bhutan.

Cultivation and collection

Jatamansi is a perennial herb propagated by cuttings of the underground parts. The favourable altitude for the Luxurious growth of the plant is 3000 – 5000m.

The rhizomes are collected from the wild grown plants only. The plant is about $10 - 60 \, \text{cm}$ in height and with stout and long woody root stocks.

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Macroscopic features

The drug is usually covered with red to brown fibres which are the

accumulated remains of leaf bases. In the fibres remains of aerial shoots

are also seen.

Rhizome is cylindrical and with brown to deep greyish fibres, is 1-5

cm long and 0.5 – 3cm in diameter. The internal colour of rhizome is red to

brown.

If all the leaf bases, aerial shoots, and adventitious roots are

removed the rhizome shows rough surface with transvers rings. These

rings represent the scars of nodes, leaf bases and the adventitious roots.

Adventitious roots are thin, branched and red to brown in colour

odour is slight and aromatic and Taste Aromatic and pungent.

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Commercialization:

The rhizome of jatamansi is used in the preparation of medicinal oils

and in perfumery. The dried rhizomes are steam – distilled to yield between

1-2% of essential oil, commercially known as spikenard oil. It can be used

in perfumes with an oriental basis, heavy florals, animal amber types etc., It

blends well with cedar wood and Lavender.

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PHARMACOGNOSTICAL ASPECT

Commercial sample of jatamansi comprises the rhizomes of Nardostachys jatamansi detailed pharmacogonostic characters of the rhizome have been reported by Mehra & Garg (1962). The rhizomes occur with or without a bit of the tap root attached and covered all over with fibres which are the reminants of the leaf hases. A striking feature is the presence of Interxylary cork in the mature rhizome, in the form of continuous branching angular or fluted tubes occuring within each other. The fission of the rhizome into separate cards in the older regions is due to joining of the outermost.

Interxylary cork ring with innermost cortical cork ring in the primary medullary ray region. Schizogenous cavities are present in the inner cortex. Phloem and xylem parenchyma of the young rhizome. Minute droplets of oil are present in the cork cells. The aerial shoot is generally hollow with a ring of vascular bundles which are embedded in a thick continuous zone of fibrous tissue. The tap root resembles the rhizome in the formation of successive rings of cortical as well as interxylary cork, but does not split into cork.

Macroscopic characters

Colour - Dark grey rhizomes are crowned with reddish

brown tufted fibres

Odour - Highly aggreable, aromatic

Taste - Acrid, slightly bitter and aromatic

Size - Rhizomes are 2.5 to 7.5cm in length

Shape - Elongated and cylindrical

The fibres present on the rhizomes are the remaining of leaf hases.

Rhizomes break easily and internally they are reddish brown in colour.

Microscopic characters

Transverse section of the rhizome is characterised by the presence of outermost 2 – 5 layers of cork cells containing oil globules.

The cortex is marked by the presence of schizogeneous canals. Phloem is in the form of patches and xylem is characterized by vessels vessels have typical scalariform thickening. Cambium is continuous and well marked.

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Epidermis is absent in mature rhizome shows ring or rings of cortical cork broken at some places Between the successive cork rings cortical remains of cortical parenchyma with leaf traces are seen Inter Xylary and Medullary cork is the characteristic feature of this drug.

Interxylary cork occurs in the form of continuous, branched, angular or fluted tubes arranged in one another, outermost interxylary cork ring fuses with innermost cortical ring in the area of Medullary rays and thus in that part rhizome is broken in small cords in the order regions.

 Pharmacognosy by J.S.Qadry shah and Qadry's.

Adulterant

Mukerjee wrongly described rhizome selinum vaginetum (umbelliferae) found also at higher attitude as jatamansi Morphologically it resembles jatamansi, but differs microscopically in that it contains vittae in young and old rhizomes and roots and fan – shaped crystals of calcium oxalate; not found in jatamansi.

Further it doesnot contain interxylary and medullary cork and stele doesnot break in cords it contains volatile oil. Volatile oil is used as sedative and analgesic.

 Pharmacognosy by J.S.Qadry shah and qadry. Pharmacognostic studies of the commercial samples of N.jatamansi revealed the presence of adulterant very similar to genuine N.jatamansi in external appearance. The adulterant was found to be the rhizome of selinium vaginetum. The rhizome is covered with a dense tuft of bristly fibres which represent the skeletal remains of leaf bases as in N.jatamansi.

Microscopically, however the two can be easily distinguished (Mehra & Jolly, 1963) by the features described below.

The distinguishing characters of N.jatamansi and S.Vaginetum.

N. Jatamansi	S.Vaginetum
Presence of Inerxylary & Medullary	Absence of Interxylary of Medullary
cork which results in the splitting of	cork. No splitting of the stele in old
the stele into 4-6 seperate cords in	rhizome.
the old rhizome.	
Schizogenous cavities in the young	Secretary candles lined with
rhizome in the inner cortex and	epithelial cells present in the cortex,
parenchyma of phloem and xylem	pith and secondary phloem in both
but without any epithelial cells.	the old and young rhizomes and in
	the root.
Fan shape crystals of calcium	Fan shaped crystals of calcium
oxalate absent in the parenchyma	oxalate present in parenchyma cells.
cells oil globules cork cells present	
Oil globules in cork cells present.	Oil globules in cork cells absent.

N.Jatamansi & S.Vaginetum can also be differentiated by the T.L.C

pattern of the petroleum ether extracts of the two plants S.Vaginetum

contain a number of coumarins viz.angelicin, oroselon, lomatin, selindin

and vaginidin, and a flavinoid selenone which are absent in N.jatamansi

(Seshadri, 1969).

Chemical Test:

On microchemical studies, N.jatamansi cortex cells appear blue and

brown in colour, when the section is treated with strong lodine. The xylem

tissue turns into reddish violet colour and the cork tissue pink by the action

of phloroglucinol and Hydrochloric acid (Sharma, 1972).

The 80% alcoholic extract of drug shows bluish - white fluorescence

under ultravoilet light.

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PHYTOCHEMICAL ASPECT

In a preliminary chemical study, Nardostachys jatamansi was reported to yield 2% of volatile oil, containing an unidentified ester and alcohol, and two alkaloids. (Bose et al., 157 a).

The rhizomes of N.Jatamansi yielded 'Jatamanshic acid' (Chaudry et al., 1951) which was assigned a bicycle azulenic sesquiterpene structure.

The petroleum ether extract of the rhizomes gave a ketonic principle called 'Jatamansone' (Govindhachari et al., 1958) which was present also in the roots of Valeriana officinalis.

The hydrocarbon fraction of the petroleum ether extract N.Jatamansi rhizome yielded a known sesquiterpene Viz., Seychellene and a new hydrocarbon, seychelene. The hexane extract of the rhizomes of N.jatamansi yielded β.Sitosterol (Anjaneyulu et al., 1965).

The neutral fraction of the concrete of roots gave a number of compounds viz., Valeranone, Valeranal, Nardal, Calarenol, Nardostachone, N- hexacosanyl arachidate, n-hexacosanol, calarene, n-hexacosene, isovalerate and β -sitosterol.

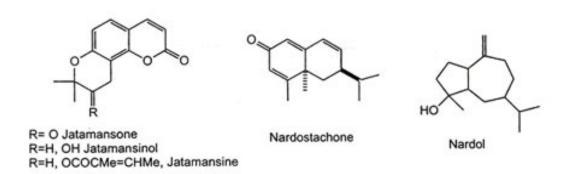
The roots of extraction with light petrol yielded norseychelanone, seychellen, parchouli alcohol, and also ∞ - & β pachoulenes (Rucker et al., 1976). The isolation of a sesquiterpene ketone 1 (10) - aristolen – 2- one

along with β - sistosterol and three unidentified compounds $C_6H_{24}O_3$; C14H24O; C15H22O) from the petroleum ether extract of the roots has also been reported (Maheswari & saxena, 1980).

The oil from the N.Jatamansi roots yielded terpenic coumarins, oroselol and a new one named as 'Jatamansin' (Shanbhag et al., 1964).

Nardostachnol, 9 – dehydroaristolene, 1(10) – dehydroaristolene, 2 β - Maalenine, and 1,2,9,10 – Tetrahydroaristolene identified in essential oil. Aristolen - 2 – one, β – sitosterol and three unidentified compounds isolated from roots.

Compendn Indian medicinal plants
Vol.2, Rastogi & Mahrota
PID, New Delhi 1991, P . 480.



New sesquiterpene ketone – jatamansone – isoloated from rhizomes, β - Maaliene and calarene from oil, a new terpenine coumarin – Jatamansin, oroselol, from roots; ∞ - pinene, β - Pinene, carene , β - eudesmol, elemol, a C_{30} hydrocarbon, β - sitosterol, Jatamansin, angelicin,

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Jatamansinol from roots, Nardol from roots. A new diethenoid bicyclic ketone - nardostachrone from roots.

Glossary of Ind. Med Plants
 Chopra, Nayar PID,
 New Delhi, 1956 P.173

$$\beta$$
 – Maaliene Angelicin

Jatamansi contains volatile oil and volatile oil contains a ketone '
Jatamansone' which is same as valeranone mentioned in valerian possibly it contains ' Jatamansic Acid'.

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Jatamansi contains 1 -2% of pure yellow volatile oil, resin, sugar, starch and bitter principle, an alcohol and its isovaleric ester.

It also contains jatamansic acid and ketones jatamansone and nardostachone.

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PHARMACOLOGICAL ASPECT

(Review of update works on Jatamansi)

Various extracts (Light petroleum,benzene and ethanol) of Nardostachys jatamansi, root showed both sedative action in rats as revealed by physical inactivation and potentiation of phenobarbitol sodium sleeping time in rats, the hypotensive activity in cats. The ethanolic extract was found most active (Hamied et al., 1962).

The alcoholic extract potentiated hexobarbital narcosis in rat, reduced the rat brain serotonin content, markedly increased the reaction time of trained rat in escaping through the tunnel (in a columbia obstruction box), decrease the conditional avoidance performance in cats and also abolished the tonic extensor response in rats subjected to maximal electroshock seizures. The effect of extract was less than that of diphenyl hydantoin sodium and was associated with signs of neurological deficit (Gupt , 1966).

The ethanolic extract (50%) of N.jatamansi rhizomes had no effect on the CNS of Mice. (Bhakuni et al., 1969).

The essential oil from the rhizomes had a depressant action on the CNS of guinea pigs and rats (Chopra et al., 1954).

Jatamansone, the sesquiterpene from N.jatamansi was shown to exert tranquilising activity in Mice and Monkey, hypothermic activity in Mice and antiemetic effect in dog (Arora et al., 1962 a).

Further studies on the effect of jatamansone on the biosynthesis and metabolism of serotonin in rabbit brain revealed an impairment of biosynthesis of serotonin in the brain tissue, thus leading to a reduction in the brain levels of 5 hydroxytryptamine. The degradation of serotonin was unaffected (Arora et al., 1962).

Jatamansone was found to be more effective than diphenyl hydantoin sodium and essential oil of Jatamansi in maximal electro shock seizures. However these compounds were ineffective against metrazol seizures. The LD₅₀ of Jatamansone & essential oil of jatamansi by the I.P route in Mice were 350mg / 1kg and 900mg/ kg respectively. (Arora et al., 1958a).

A compound herbal preparation with N.jatamansi, Acorus calamus and valeriana wallichi as ingredients showed CNS – depressant activity in rabbits and also inhibited the post – isolation syndrome in Mice (Moghe et al., 1981).

In another study, aqueous, alcoholic, volatile oil and alkaloidal fraction of N.jatamansi rhizomes and roots were studied for sedative and CVS effects. The alkaloidal fraction showed a significant and sustained hypotensive action in dogs. The fraction also produced a marked relaxation of plain muscles and depression of CNS and mild degree of relaxation of the skeletal Muscle. (Bose et al., 1957b). The action of the aqueous and alcoholic extracts and the total alkaloidal fraction of the tissue respiration in rat brain, liver and heart was studied. The total alkaloids

produced maximum inhibitory effect on tissue respiration leading in higher doses to complete cessation of respiratory activity in these tissues. A sedative action was also observed in rats after parenteral injection of the total alkaloids (Bose et al., 1975c).

The essential oil obtained from the rhizomes of N.jatamansi exerted prolonged and pronounced hypotensive effect in dogs. It did not depress the vasomotor centre but blocked the proprioceptive blood pressure regulating reflexes. The oil had negative ionotropic and positive chronotropic effect on the heart of the frog and dog. In moderate hypotensive doses it didn't lead to any ECG changes in dogs. It didn't bliock the ganglionic transmission but showed some adrenolytic action. The oil didnot depress the respiration but on the other hand caused some initial stimulation. (Arora et al., 1958 a).

The oil-free aqueous extract of N.jatamansi showed a transient hypotensive effect and electro cardiographic changes in dog's heart, apart from contracting frog's rectus Muscle. The CVS effect of the extract was similar to that of potassium. It could not be ascertained, however, whether all the CVS effects of N.jatamansi extract could be explained by the presence of potassium alone. The potassium content of the aqueous extract was found to be 5-67mg/m1 of the extract (sheth & kekre, 1956)

Jatamansone showed a potent and prolonged hypotensive activity in normotensive and hypertensive rats, normotensive dogs and cats. The hypotensive action was evident in both anaesthetized and unanaesthetized animals (Arora, 1965a).

Anti arrhythmic activity of the volatile oil of N.jatamansi was first reported against acetyl — choline induced fibrillations in monged dogs (Arora and Madan, 1955). The volatile oil was compared with Quinidine in another study (Arora and Madan, 1956) and was found to be less active than quinidine as an antiarrhythmic agent against isolated rabbit auricular fibrillation, experimental auricular flutter in anaesthetized dogs, auricular fibrillation induced by aconitine and acetylcholine in dogs. The volatile oil had no effect on Digitalis induced ventricular arrhythmia in dogs. It showed on advantage over quinidine in causing a lesser degree of slowing as tested by the ECG changes in the cat. The acute I.V toxicity of the oil was also lower than that of Quinidine (Arora & Madan, 1956).

Jatamansone was found to be more effective than quinidine as well as the essential oil of N.jatamansi in suppressing ectopic ventricular activity in unanaesthetized dogs induced by the two stage coronary ligation. It was however less effective than quinidine against acetyl – choline induced auricular fibrillation in anaesthetized dogs (arora et al., 1958 b).

A compound herbal preparation (khamira Abresham Arshadwala) having N.jatamansi as one of the ingredients lowered the blood pressure in hypertensive rats & also exerted and anti – arrhythmic action (Siddique, 1964).

The alkaloidal fraction of N. jatamansi showed a bronchodilatory effect on isolated tracheal chain of guinea pig. It also partially relieved histamine - induced bronchoconstriction (Bose et al., 1957a). Bronchodilator effect of powdered N. jatamansi fumes and aerosols against histamine induced bronchial Asthma in guinea pigs has been reported. In two different sets of experiments N. jatamansi fumes could not only relieved guinea pigs with acute dyspnoea induced by histamine aerosol, but also protect the animals (When pre treated with N. jatamansi alcoholic extract) against developing dyspnoea on exposure to histamine aerosol (Gupta et al 1961).

The alcoholic extract of N. jatamonsi inhibited the constrictor response induced by histamine, serotonin and acetylcholine in isolated smooth muscles (ie Trachea, Colon, Intestine & uterus).

The extract also showed a direct papaverine - like antispasmodic effect on the intestine (Gupta et al., 1962). There was also reduction in perfusion pressure through the isolated lungs of rats or guinea pig. The alkaloidal fraction of N. jatamansi also antagonized the perfusion outflow caused by histamine and serotonin (Gupta et al., 1963).

Anti bacterial activity in the alcoholic and aqueous extracts of N. jatamansi roots was reported against staph aureus and E.coli perhaps for the first time by George and pandalai (1949). In later studies, the alcoholic extract (20%) showed antibacterial activity against strep pyogenes, sal. posteurella multocida and ps. aeruginosa, although the anti bacterial activity of the aqueous extract was not confirmed (Naung et al., 1962) Ethanolic extract (50%) of the rhizomes was reported by the CDRI, Lucknow, to be devoid of anibacterial activity against B.subtils, staph aureus, sal typhi, E.coli, A. tumefaciens and Mycobacterium tuberculosis H37 Rv (Bhakuni et al., 1969).

The essential oil of N.jatamansi didnot show any antituberculous activity against M.tuberculosis H37 RV (Ramaswamy & Susi, 1967). The oil showed weak antibacterial action against staph aureus, E.coli, Sal typhosum, Vib cholera, shigella flexneri (chopra et al., 1954) and against vibrio cholerae, salmonella faecalis, salmonella typhi, clostridum diptheriae and streptococcus pyogenes (Girgune et al., 1978 a).

The ethanolic extract of (50%) the rhizomes was devoid of antiprotozoal activity against Entamoeba histolitica (Bhakuni et al., 1969) whereas the oil had a weak action against p ccaudatum (Chopra et al., 1954).

The ethanolic extract (50%) of the rhizome showed antifungal actiity agianst candida albicans, cryptococcus neoformans, Trichophyton mentagrophytes, Microsporum canis and Aspergillus niger (Bhakuni et al.,

1969), whereas the essential oil have been reported to reveal potent antifungal activity against Helminthosporium turcicum and Alternaria helianthi (Girgune et al., 1978 b).

The essential oil of N. jatamansi showed anthelmintic activity against Taenia solium (Girgune et al., 1978C).

The aqueous extracts of N.jatamansi root exhibited diuretic action in rats which compared favourably with that of potassium acetate and was more potent than urea (Gujral et al., 1955).

The ethanolic extract (50%) of the rhizomes showed hypotensive effects in cat / dog and antispasmodic action on isolated guinea pig ileum but was devoid of any hypoglycemic, antiviral or anticancer activities . The LD $_{50}$ of the extract in Mice was found to be > 1000mg ./ Kg IP (Bhakuni et al., 1969).

The semicarbazone derivative of Jatamansone revealed antioestrogenic activity in mice by inhibiting the increase in uterine weight caused by estrogen alone. When administered alone, the compound had no estrogenic activity but decreased the uterine weight, revealing an antiestrogenic activity (Agarwal et al, 1973).

- Glossary of Indian Medicinal plants

R.N.Chopra, S.L.Nayar

I.C.Chopra.

EFFECTS OF N.JATAMANSI ON BIOGENIC AMINES AND INHIBITORY

AMINO ACIDS IN THE RAT BRAIN

The effect of acute and sub chronic administration of an alcoholic

extract of the roots of N.jatamansi on nor epinephrine (NE), dopamine

(DA), serotonin (5-HT), 5 - hydroxy indoleacetic acid (5 - HIAA),

Gamma - amino butyric acid (GABA) and taurine were studied in male

albino wistar rats. The acute oral administration of the extract did not

change the level of NE and DA but resulted in a significant increase in the

level of 5 – HT and 5 – HIAA. A significant increase in the level of GABA

and Taurine were observed in the drug treated groups when compared to

the controls. A 15 - day treatment resulted in a significant increase in the

levels of NE, DA, 5 HT, 5 – HIAA, Jatamansi causes an overall increase in

the levels of central monoamines and inhibitory amino acids.

Authors :

Prabhu.v, Karanth K.S, Rao .A,

Dept. of Bio chemistry,

Kasthuribai Medical college,

Karnataka

Journal name:

Plant med 1994 Apr; 60 (2) 114 – 7.

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HYPOLIPIDAEMIC EFFECTS OF CURCUMA LONGA AND

NARDOSTACHYS JATAMANSI, DC IN TRITON - INDUCED

HYPERLIPIDAEMIC RATS

50% ethanolic extract of curcuma longa (tuber) and N.jatamansi (

whole plant) feeding elevates HDL - Cholesteorl / Total cholesterol ratio.

The extracts also caused a significant reduction in the ratio of total

cholesterol / phospholipids curcuma longa exhibited better cholesterol and

triglycoride lowering activity (Ch - 85%, Tg - 88%) as compared to

N.jatamansi in triton - induced hyperlipidaemic rats . In view of the

protective action of HDL against heart disease and atherogenicity, curcuma

longa consumption is recommended.

PMID : 3215683, UI : 89108563.

Authors : Dixit VP, Jain P, Joshi SC.

Journal name : Indian J physiol pharmacol 1998.

oct - Dec 32 (4); 299 - 304.

30

NARDOSTACHYS JATAMANSI PROTECTS AGAINST LIVER DAMAGE INDUCED BY THIOACETAMIDE IN RATS

Ali S, Ansari KA, Jafry MA, Kabeer H, Diwakar G. (Department of Biochemistry, Faculty of Science, Hamdard University, New Delhi).

Nardostachys jatamansi is a medically important herb of Indian origin used for centuries in Ayurvedic and Unani systems of medicine for the treatment of various ailments. In the present paper, a 50% ethanolic extract of the rhizomes of N. Jatamansi is shown to possess hepatoprotective activity. Pretreatment of rats with the extract (800mg/kg body wt, orally) for three consecutive days significantly ameliorated the liver damage in rats exposed to the hepatotoxic compound thioacetamide. Elevated levels of serum transaminases (aminotransferases) and alkaline phosphatase, observed in thioacetaminde alone treated group of animals, were significantly lowered in N. Jatamansi pretreated rats. Pretreatment of the animals with the extract also resulted in an increase in survival in rats intoxicated with LD90 dose of the hepatotoxic drug.

ISOLATION AND PHARMACODYNAMIC ACTIVITY OF THE SESQUITERPENE VALERANONE FROM NARDOSTACHYS JATAMANSI DC

[Article in German] (Rucker G, Tautges J, Sieck A, Wenzl H, Graf E)

The known sesquiterpene valeranone (= yatamanson) was isolated from the tubterranean parts of Nardostachys yatamansi (DC). It was pharmacologically investigated in animal experiments of sedative, tranquilizing and antihypertensive properties. In some experiments, typical for tranquilizers, certain activities could be demonstrated such as the prolongation of barbiturate hyponosis, the impairment of rotarod performance, an anticonvulsive activity on electric shock and potentiation of the body – temperature lowering activity of reserpine. In three other pharmacological models an anti ulcer action was detected. In general the activity of valeranone was lower than those of the standard substances used . As regards the hypotensive property only a weak activity was demonstrated . In toxicological studies on rats and mice an oral LD $_{50}$ of greater than 3160 mg/kg was fond, which suggestes the possibility of a therapeutically useful dose ratio.

THERAPEUTIC USES

Jatamansi

Nardostachys jatamansi.

English Name

Musk Root, Indian Spikenard.

Family

Valerianaceae.

Part used

Rhizome, Rhizome oil.

Action

Aromatic Expectorant

Antispasmodic Anodyne

Carminative Antiseptic

Deobstruent Deodorant

Digestive Somniferous

Diuretic Antipyretic

Emmenagogue Tonic

Nervine Tonic Aphrodisiac

Uses

Convulsions

Digestive disease

Epilepsy

Flatulence

Gastric disorders

Tleart palpitations

Jaundice

kidney stones

Respiratory diseases

skin conditions

Typhoid

Seminal debility.

Therapeutic uses

Jatamansi has the power to promote awareness and calm the mind. It is very useful for palpitation, Tension, headaches, restlessness and is used for promoting awareness and strengthening the mind. It aids in balancing the body of all three ayurvedic doshas. The herb's sedative properties increases awareness as opposed to valerian that dulls the mind.

Jatamansi is a useful hair tonic and is commonly used in hair oil, promoting hair growth and lustre. It is also used in oils and pastes that improve complexion and general health of the skin.

No side effects have been noted so far.

It is very interesting to note that in condition of insomnia, and restlessness this drug was used by sushruta to produce tranquility and sedation Infusion prepared from the fresh roots is employed in the treatment of spasmodic hysterical states, palpitations and tension headache. It is also said to be useful in menopausal disturbances.

Clinical trials were carried out with jatamansone in essental hypertension. Jatamansone has been used in febrile delirium and also in delrum tremens In dysmenorrhoea, it is used for pain relief and a smooth menstrual flow. Nardostachys was recommended in the Ayurvedic tradition for nervous and spasmodic symptoms, such as heart palpitations, headache, shaking and convulsims. The active constituents of Nardostachys are similar to those found in valerian. In India, modern research with the herb has been aimed at examining new uses rather than the traditional ones; It is being examined for its liver protective effects, ability to increase nerve growth factor and Lipid lowering effects.

The oil possess antiarhythmic and hypotensive activity Jatamansone an active principle of N.Jatamansi, brings forth a significant reduction in hyperactivity, restlessness and aggressiveness in hyperactive children.

Cosmetic application

Jatamansi is a useful hair tonic and is commonly used in hair oils, promoting hair growth and lustre. It promotes hair growth and imparts black colour to the hair. It is also used in oils and pastes that improve complexion and general healthy of skin.

In Germany and Japan, some interest in this herb as an alternative to valerian has been shown, in that preliminary experiments (in Laboratory animals) show that it has an even lower toxicity than valerian (which already has low toxicity).

- www.wikipedia.org

Jatamansi is useful in vitiated conditions of pitha and vadha, burning sensations, cough, Asthma, Bronchitis, Pectoralgia, Cephalgia, Inflammations, Somatalgia, Dyspepsia, colic, flatulence, Hepatopathy, Nephropathy, Strangury, Amenorrhoea, dysmenorrhoea, Lumbago, pharyngopathy, dermatopathy, Leprosy, erysipelas, epilepsy, hysteria, convulsions, neurosis, hypertension, Grey hair, falling of hair, Intermittent fever, General debility.

Indian Medicinal plants
 (A compendium of 500 species)
 orient longman

Jatamansi roots should also be used fresh as an aromatic adjunct in the preparation of Medicinal oils & in perfumery. Jatamansi is a good substitute for the official valerian. Infusion prepared from fresh roots is employed in the treatment of spasmodic hysterical affections, especially palpitation of heart, Nervous headache, chorea, flatulence etc., in doses of 1-2 ounces 3 times daily.

It is said to be useful also in menopause disturbances, hystero - epilepsy & similar nervous and convulsive ailments.

Dose: 10 -20 grains in powder. It may be usually combined with a few grains of camphor & cinnamon.

Voltile oil from rhizome can be used in many diseases of digestive, respiratory organs and in jaundice, also in leprosy. It is also employed mixed with sesame oil, it is rubbed on the head as a nerve sedative. It also promotes growth & imparts blackness to hair.

- The Indian materia medica

A.K. Nadkarni.

Spikenard oil possess antiarrhythmic activity in cases of auricular flutter, it is also effective than quindine, but has the advantage of being less toxic.

Jatamansone is more potent than the oil & is also more active than quinidine ventricular Tachycardia resulting from acute MI in experimently induced arrhythmias, it is as effective as quinidine except in the Acetyl choline induced Auricular fibrillation in which it is considerably weaker.

Jatamansone possess Anticonvulsant action as well. The oil exerts a hypotensive effect & in moderate doses it has a distinct depressant action on the CNS. Lethal doses causes deep nacrosis & Ultimately death within few hours. The root extracts show sedative properties.

A tincture of rhizome is given in intestinal colic & flatulence. The rhizome is used as an aromatic adjunct in the preparation of Medicinal oils.

- Wealth of India Vol VII.

Therapeutic Uses

In other system of Medicine

Ayurveda

Roots are acid, bitter with a flavour tonic, cooling, antipyretic, cure 'Tridosha', 'kapha', Biliousness, diseases of blood, burning sensation. Erysipelas, leprosy, skin diseases, throat troubles, ulcers, improve the complexion.

They also have antispasmodic effect and are often employed in treatment of epilepsy, hysteria, convulsive affection, palpitation of heart, intestinal colic. It enters into the composition of a compound powder which is burnt and used for inhalation in bronchial affection.

The rhizome, in combination with other drug is prescibed in snake bite and scorpion sting.

- Charaka, Sushruta.

Unani:-

The roots have a bitter sharp taste, tonic stimulant, diuretic, emmenagogue, carminative, and stomachic, laxative.

If increases the lusture of the eyes.

If promotes growth, blackness of hair.

Also useful in gleets, cough, chest pain intestinal inflammation.

Kidney and lumbar, troubles, dry wounds and increase the appetite.

- Indian Materna Medica

A.K. Nadkarni Vol – 2.

Therapeutics:

- 1. It is prescribed as nervine tonic and aromatic adjunct in the preparation of Medicinal oils and gritas.
- 2. In doses of 45gms, it is used as an expectorant in coughs and cold.
- 3. It is used in treatment of epileply, hysteria, and convulsve affections.
- 4. It is used in palpitation of the heart.
- It is administered suspended in mucilage with cinnamon water and is given as a carminative in cases of flatulence and as reflex stimulant in vomiting palpitation.
 - Pharmacopoeia India, K.C.Bose P.No. 122.

SPIKENARD OIL

Synonyms: Nardostachys jatamansi root oil

Nard root oil

Indian valerian root oil

Method of extraction

Jatamansi oil is obtained by steam distillation of dried rhizomes of N.Jatamansi DC.

Organoleptic properties

Appearance: Fluid to slightly viscous liquid.

colour : Varies from amber to deep blue or greenish blue.

Aroma: Heavy, Sweet - woody and spicy - animal odor.

Physio – chemical properties

Specific gravity: 0.9300- 0.9587 at 25°C.

Refractive index : 1.5055 – 1.5458 at 25° C.

Acid number : 1.5 - 8.

Ester number : 6-45.

Ester number after

Acetylation : 40-65.

Solubity : Soluble in 0.4 – 1.5 vol of 90% alcohol.

Chemical constituents

Spikenard oil contains an alcohol (C₁₅H₂₄O) and its isovaleric ester,a saturated bicyclic sequiterpene ketone, jatamansone (C₁₅H₂₆O b- p 108°) has been isolated from the rhizomes.

Actions

Stimulant

Antiseptic

Insect repellant

Spikenard oil - medicinal uses:

- The oil possess antiarrhythmic activity with possible therapeutical usefulness in cases of auricular flutter. It is less effective than Quinidine but has the advantage of being less toxic.
- The oil exerts a hypotensive effect and in moderate doses it has a distinct depressant action on the CNS.
- 3. It is also believed to be useful for leprosy.
- 4. It promote hair growth & helps in maintaining its color as well.
- It can be used in many diseases of digestive, respiratory organs and in jaundice.
- It is also employed mixed with sesame oil for rubbing on the head as a nerve sedative.
- 7. It is used in the treatment of stomachache, constipation and cholera.

GUNAPADAM ASPECT

°¼¡Á¡ï°¢ - Jatamansi

Nardostachys jatamansi

§ÅÚ | ÀÂ÷ jû - Synonyms

- ❖ °¼¡Á;ï°¢
- ❖ f¼;Á;ï°¢
- ❖ "À°;°¢
- ♠ Á¡Á¢°¢
- ♣ â¾S¸°¢É¢
- ۰ê"Ä
 - ̽À;¼õ ãÄ¢", ÅÌôÒ 6õ À¾¢ôÒ,2002 À.±ñ:

311

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- ❖ þãò¾ à¢ò¾ °ÁÉ¢

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- «ìÉ¢§Å°Ã¢ý °Ã,°õ†¢"¾ 3õ À̾¢
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À.±ñ : 332 Ó¾Ø À¾¢ôÒ ¬¸Šð 1987

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- ̽À;¼õ ãÄ¢", ÅÌôÒ 6õ

À¾¢ôÒ,2002

À.±ñ : 311

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6% Å%¢ôÒ ¬¸Šð 1987Å.±ñ :

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- ̽À;¼õ ãÄ¢", ÅÌôÒ À.±ñ: 311

6õ À¾¢ôÒ,2002

°¡¾¡Ã½ Á¡õ...¢ : ¸;÷ôÒ, ¸°ôÒ Í "Å |¸;ñ½Đ

Í ó¾ °¼¡Á¡õ...¢ : °ôÒîÍ"Å, ÌÙ"Áò¾ý"ÁÔõ,

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Ó¾ø À¾¢ôÒ ¬¸Šð 1987À.±ñ: 322

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" | Á;Æ¢°¼; Á;ï°¢ §Áý"Á Ó¾¢÷Í"Å ,";¾¢ì¾õ ÅÆ¢¾Õ ÁĐÃï °£¾Å£Ã¢Âí ,ÎÅ¢À; ,|ÁýÀ |Á;Æ¢,Àõ Ãò¾À¢ò¾ |Á;Õ ÅÕÅ;ö× Ì‰¼õ

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°¼¡Á¡ï°¢. ¸"¡Â, ¾¢ì¾, ÁĐà Í"Ÿ "ÇÔõ, °£¾Å£÷Âò"¾Ôõ, ¸ÎÅ¢À¡¸ò"¾Ôõ $^{-}$ "¼ÂĐ. ¸Àõ, þÃò¾À¢ò¾õ Å;ö×, ̉¼õ, Å¢"õ, Å¢…÷ôÀõ §À¡ììõ.

",Õ¾¢Â ÅÄ¢"Á ÒÒ¾¢ ,;Ó¾¢Ô Á;ÌÌ ÁÝÈ¢

ÅÕĐÂ÷ â¾Á;¾¢ §¾;"Óõ ÅÈðÊ §Â;ðÎõ

Á¢ÕĐÅ; Á¢"à ÅÇ÷ΰ¢ Á¢ÌÒ¾¢Îí ,Ó¾Á;õ...¢

|ÀâĐÂ÷ ̽ÓÁ£§¾ |ÂÝÈÉ÷ |Àâ§Â;÷¾;§Á".

- À¾;÷ò¾ Àï°Ì½ Áï°Ã¢

À.±ñ: 181-182.

°;¾;ý °¼;Á;õ...¢ - â¾ò¾;ø §¾;ýÚõ ĐýÀí, "ÇÔõ, ,;öî° "ÄÔõ, ¿ï "°Ôõ §À;ìÌõ.

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Ó¾ø À¾¢ôÒ ¬¸Šð 1987À.±ñ: 322.

வழக்கு:

சீமை இனத்தைப் பொடி செய்து 10 - 20 குன்றி எடை நீரில் கலந்து நாள் ஒன்றுக்கு இரண்டு, மூன்று முறை கொடுக்க பழைய சுரம், வலிப்பு நீங்கும்.

முக்கால் வராகனெடை கொடுக்க கோழையை வெளிப்படுத்தும்.

நாட்டு இனத்து சடாமாஞ்சியை நசுக்கி, இரண்டு வராகணெடை எடுத்து ஓர் ஆழாக்கு வெந்நீரில் 1 மணி நேரம் ஊற வைத்து வடிகட்டி, அரைக்கால் முதல் காலாழாக்கு வீதம் தினம் மும்முறை கொடுத்துவர சூதக சன்னி, வலிப்பு இவை நீங்கும்.

இதிலிருந்த ஒரு வகை மஞ்சள் நிறம் பொருந்திய எண்ணெய் எடுக்கப்படும் இதை மேற்கூறிய நோய்களுக்கு கொடுப்பதுடன் நரை மயிர் கறுப்பதற்கும் இதைத் தடவுவதுண்டு.

> - குணபாடம் மூலிகை வகுப்பு ப.எ.416. 6õ À¾¢ôÒ, 2002

சடாமாஞ்சில் வேரை கொத்துமல்லிக் கீரைச் சாற்றில் உரைத்து வடிகட்டிக் கண்களில் 1-2 துளிவிட கண் சிவப்பு மாறும் . பார்வை தெளிவுபடும்.

தலையில் தடவ மயிர் வளரும்.

இந்தச் சரக்கு விசேஷமாக , காக்கை வலி, சூதக சந்தி, சுவாசகாசம், மந்தாக்கினி, குடைச்சல், இருதய துடிப்பு , மேல் சுவாசம், மூர்ச்சா ரோகம் முதலியவைகளுக்கு பயன்படுகிறது.

- சித்த வைத்திய பதார்த்த குண விளக்கம்.

À¾¢ôÒ, 1998 ப.எ.304.

சடாமாஞ்சில் வேர் உபயோகம் : ½ பலம் வேர் எடுத்து நொறுக்கி , ½ படி நீரிலிட்டு ¼ படியாய் வற்றக் காய்ச்சி வடித்துக் கொடுக்க இரைப்பு , இருமல் சுவாசம் சுரம் இவற்றிற்கு கொடுக்க நன்று.

- அனு- வை பிரம்ம ரகசியம் ப.எ. **284** Ó¾üÀ¾¢ôÒ Á¡÷î 1999

சடாமாஞ்சில் வேரை இடித்து சூரணம் செய்து வேளைக்கு ½ வராகன் தேனில் இருவேளை கொடுக்கலாம்.

(அல்லது)

2 வராகன் சடாமாஞ்சியை நசுக்கி ஒரு மட்கலயத்தில் போட்டு ¼ படி சலம் விட்டு 1/8 படியாய் சுண்டக் காய்ச்சி வடிகட்டி வேளைக்கு 1-2 அவன்ஸ் வீதம் தினம் இருவேளை கொடுக்கலாம்.

> இவை இரைப்பைக்கும் , ஈரலுக்கும் பலம் கொடுக்கும். உதிரசிக்கலை நீக்கும். நீரை வறட்டும்.

தீபனத்தை உண்டாக்கும்.

இதனை அதிக அளவாயும் , நீடித்தும் தரக்கூடாது அங்ஙனம் கொடுக்க தலைவலி , மயக்கம் , பார்வை மந்தம் உண்டாகும் இன்னும் இதை இதர சரக்குகளுடன் கூட்டி உபயோகப்படுத்துவதுமுண்டு. எங்ஙனமெனில் ,

சடாமாஞ்சில் துரணம்

சடாமாஞ்சில் - 4 வராகன்

கருவாப்பட்டை - 1 வராகன்

வால்மிளகு - 1 வராகன்

சோம்பு - 1 வராகன்

சுக்கு - 1 வராகன்

இடித்து சூரணித:து 2 வராகன் சர்க்கரை கூட்டி வேளைக்கு ½ - 1½ வராகன் தினம் இரு வேளை கொடுத்து வர, வயிற்றுப்புசம் , வயிற்றுவலி , மூர்ச்சாரோகம் குணமாகும்.

- Indian Materia Medica A.K.Nadkarni, Vol-1. Print 1993 சடாமாஞ்சில் வேர் சேரும் காசத்திற்கான மருந்துகள்:

1. தாளிசபத்திரி சூரணம்

இரு வேளை திரிகடி வீதம் கொடுக்க காசம் - 5, ஷயம் - 4 தீரும்.

-அனுபோக வைத்திய பிரம்ம இரகசியம் முதற் பதிப்பு மார்ச் 1999

- கோஷாயி ப.எ.102.

2. தூதுவளை கிருதம்

2 வராகன் வீதம் இரு வேளை கொடுக்க, ஈளை, இருமல், காசம் - 5, சுவாசம் - 5, இரைப்பு, பொருமல் தீரும். - அனு . வை. பிர. ரகசியம் - கோஷாயி . ப.எ: 117.

3. புனர்னவாதி லேகியம்

காசம் , சுவாசம் குணமாகும்.

- அனு வை தேவ ரகசியம் (ப.எ.: 383).

4. பிரஹல்லவங்காதி சூரணம்

காசங்கள் குணமாகும்.

- அனு . வை. தேவ ரகசியம் (ப.எ.391).

5. தாளிசாதி சூரணம்

இருமல், ஷயம் தீரும்.

- அனு. வை. தேவ ரகசியம் (ப.எ. 401).

6. இலவங்காதி தூரணம்

இருமல் , ஷயம் தீரும்.

- அனு. வை. தேவ ரகசியம் (ப.எ.401).

7. அசுவகந்தி சூரணம்

- அனு . வை. தேவ ரகசியம் (ப.எ. 365).

8. கற்பூராதி சூரணம்

- அனு . வை. தேவ. ரகசியம் . (ப.எ.365).

9. சருவாங்க எண்ணெய்

தலை முழுகி வர கபம், காசம், பீனிசம் தீரும்.

- மருந்து செய்முறைகள் பதிப்பு அக்டோபர் 1985 (ப.எ.38).

10. கண்டங்காரி நெய்

3 கழஞ்சி உண்டுவர சுவாசகாசம் தீரம்.

- மருந்து செய்முறைகள் (ப.எ.38).

11. வில்வாதி குழம்பு

புன்னைக் காயளவு இருவேளை உண்டுவர இளைப்பு, இருமல், ஈளை குணமாகும்.

- மருந்து செய்முறைகள் (ப.எ. 62,63).

12. ஈளை, இருமலுக்கு லேகியம்

- உயிர்காக்கும் சித்த மருத்துவம் முதற்பதிப்பு அக்டோபர் 2000 (ப.எ.20).

13. முதண்ட லேகியம்

கழற்சிக் கொட்டை பிரமாணம் அந்திசந்தி உண்டு வர இருமல், சுவாசம், தொண்டைக் கட்டு தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் (ப.எ.195).

14. குழந்தைகளுக் குமாந்தக் கியாழம்

1-2 தேக்கரண்டி வீதம் தினம் 3-4 வேளை கொடுத்துவர, கபத்தினுபரி, சுரம் தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் (ப.எ.86).

15. ஜீவரஷாமிர்த தூரணம்

ஈரல்குலை நோய்க்கு நன்மை தரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் (ப.எ.121).

16. மஹாவில்வாதி லேகியம்

வேளைக்கு கொட்டைப் பாக்களவு சாப்பிட, 42 சயகாசம் பித்தம், அன்னத்துவேஷம் தீரும்.

- சிகிட்சராத்ந தீபம் 2ம் பாகமாகிய வைத்திய சிந்தாமணி (ப.எ.178).

17. அரக்குத் தைலம்

வாரம் 1 முறை ஸ்நானம் செய்துவரின், சுவாசகாசம், புராதன சுரங்கள், நளிர்சுரம் தீரும்.

> - சிகிச்சாரத்ந தீபம் இரண்டாம் பாகமாகிய வைத்திய சிந்தாமணி (ப.எ.193).

18. டிக்காமல்லித் தைலம்

வாரம் ஒரு முறை தலைமுழுகி வர, அளவு கடந்த தும்மல், தலைபாரம் குணமாகும்.

- சிகிச்சாரத்நதீபம் 2ம் பாகமாகிய வைத்திய சிந்தாமணி (ப.எ.198).

19. இலவங்காதி சூரணம்

வேளைக்கு ½ தோலா தினம் இருவேளை 20 நாள் உட்கொள் இருமல் , இரைப்பு தீரும்.

- சிகிச்சாரந்த தீபம் எண்ணும் வைத்திய நூல், பதிப்பு 1991 (ப.எ.123). 20. கஸ்தூரி மாத்திரை:

குழந்தைகட்கு மார்பில் கோழை மிகுந்திருப்பின் முசுமுசுக்கைச் சாற்றில் இரும்பை சிவக்கக் காய்ச்சி தோய்த்து வடித்த ரசத்தில் உரைத்துக் கொடுக்க நிவர்த்தி ஆகும்.

- சிகிச்சாரத்ந தீபம் வைத்திய நூல் பதிப்பு 1991 (ப.எ.127).

21. திரிபலாதி சூரணம்

திரிகடி பிரமாணம் அந்திசந்தி தேனில் மத்தித்து சாப்பிட்டு வர சீதள சம்பந்தமான ஈளை, காசம், இருமல் தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் (ப.எ.116).

MATERIALS AND METHODS

(Sadamanjil Ver Choornam)

Collection of the Drug

Rhizomes of the plant were collected from Gopalan Asan raw drug store, Nagercoil after identification.

Purification of the raw Drug

After collection it was cleaned thoroughly with fresh water and cut into small pieces (app. 1-2 inches), allowed to dry completely under sun shade for about 6-7days till the moisture was completely lost.

Preparation of the Test Drug

The dried purified sadamanjil ver pieces were made into a fine powder (chooranam) and filtered by a white cloth (Vashthirakayam)

Purification of the Test Drug (Chooranathooimai)

Sadamanjil ver chooranam was moistured with cowmilk. An earthern pot was taken and half filled with a mixture of cow milk and water. The mouth of the pot was covered with a cotton cloth and tied around its neck. The chooranam was placed on the cloth and another earthern pot was placed over the mouth of the pot completely covering the chooranam and the edges of the pots were covered with a moistured cloth. Then the

contents were boiled till the chooranam was fully cooked (pittaviyal). Then it was taken and dried in sunlight.

Route of Administration

Enteral.

Dose

One gram thrice a day with hot water after food. The prepared Sadamanjil ver chooranam used for the treatment of Eraippu Erumal was analysed by the following methods.

- 1. Bio chemical analysis
- 2. Pharmacological analysis and
- 3. Clinical assessment.

BIO-CHEMICAL ANALYSIS

BIO-CHEMICAL ANALYSIS OF SADAMANJIL VER CHOORANAM

PREPARATION OF THE EXTRACT

5gms of chooranam was weighed accurately and placed in a 250ml clean beaker. Then 50ml distilled water was added and dissolved well. Then it is boiled well for about 10 minutes. It is cooled and filtered in a 100ml volumetric flask and then it is made up to 100ml with distilled water. This fluid was taken for analysis.

QUALITATIVE ANALYSIS

S.NO	EXPERIMENT	OBSERVATION	INFERENCE
1.	TEST FOR CALCIUM		
	2ml of the above prepared		
	extract is taken in a clean	No White precipitate	Absence of
	test tube. 2 ml of 4%	is formed.	calcium.
	Ammonium oxalate solution		
	is added to it.		
2.	TEST FOR SULPHATE:		
	2ml of the extract is added	No white precipitate	Absence of
	to 5% barium chloride	is formed.	sulphate.
	solution.		
3.	TEST FOR CHLORIDE		
	The extract is treated with	No white precipitate	Absence of
	silver nitrate solution.	is formed.	chloride.

4.	TEST FOR CARBONATE	No brisk	Absence
	The substance is treated	effervessence is	Of
	with concentrated HCL.	formed.	Carbonate.
5.	TEST FOR STARCH The extract is added with weak iodine solution.	Blue colour is formed.	Indicate the presence of starch.
6.	TEST FOR IRON-FERRIC The extract is treated with concentrated Glacial acetic acid and potassium ferro cyanide.	No blue colour is formed.	Absence of ferric iron.
7.	TEST OF IRON FERROUS: The extract is treated with concentrated Nitric acid and ammonium thio cynate.	No Blood red colour is formed.	Absence of ferrous iron.
8.	TEST FOR PHOSPHATE The extract is treated with ammonium Molybdate and concentrated nitric acid.	No yellow precipitate is formed.	Absence of phosphate.
9.	TEST FOR ALBUMIN The extract is treated with Esbach's reagent.	No yellow precipitate is formed .	Absence of Albumin.
10.	TEST FOR TANNIC ACID The extract is treated with ferric chloride.	No blue black precipitate is formed.	Absence of Tannic acid.

11.	TEST FOR UNSATURATION Potassium permanganate solution is added to the extract.	It gets decolourised.	Indicates the presence of unsaturated compound.
12.	TEST FOR THE REDUCING SUGAR 5ml of Benedict's qualitative solution is taken in a test tube and allowed to boil for 2 mts and added 8-10 drops of the extract and again boil it for 2 mts.	No colour change occurs.	Absence of Reducing Sugar.
13.	TEST FOR AMINO ACID: One or two drops of the extract is placed on a filter paper and dried it well. After drying, 1% Ninhydrin is sprayed over the same and dried it well.	No Violet colour is formed.	Indicates the presence of Amino acid.

INFERENCE

The given sample of Sadamanjil Ver Chooranam contains Starch, Amino acid and unsaturated compounds.

PHARMACOLOGICAL ANALYSIS

ANTI-SPASMODIC EFFECTS OF SADAMANJIL VER CHOORNAM ON ISOLATED RABBIT ILEUM

Aim

To find out the anti-spasmodic effects of Sadamanjil Ver Choornam on isolated rabbit ileum.

Preparation of the test drug

1gram of the Sadamanjil Ver Choornam was dissolved in 10ml of water and boiled for 15minutes. The filtrate was used for the experiment.

Solution required

Aceteyl - choline - 1mg/ml. Atropine 0.5mg/ml.

Test drug (Sadamanjil ver choornam) 100mg/ml.

Nutrient solution

Tyrode – 1 to 2 litres.

Tissue used

Rabbit ileum.

Apparatus Required

Student's organ bath, sherrington rotating drum.

Procedure

A rabbit was starved for 48 hours and was allowed water ad-libturn. It was sacrified by a blow on the head and by carotid bleeding. The abdomen was quickly opened and the ileo-caecal junction was found out. A small piece of ileal portion was cut, removed and placed in a dish containing warm aerated **Tyrode Solution**. The luman of the ileum was gently rinsed out by pushing **Tyrode Solution** into it, 3cms length segment was cut from this part of ileum and was tied with thread on both ends without closing the luman and the tissue was mounted in the organ both containing **Tyrode Solution** maintained at 37°C bubbled with air by an oxygen tube.

First the drum was allowed to run for 1 minute from the baseline. Drugs are given to study the inhibiting effect of Acety-choline 0.2ml (1mg/ml) of Acetyl-choline was added and allowed to run the drum for 30 seconds. Thus the tissue was standardised and then the drum was stopped and the Acetyl – choline was washed out.

Again the **Tyrode Solution** was added to the organ bath till the lever comes to the baseline. The drum was allowed to run for 1 minute.

To the organ bath 1ml of test drug and 0.2ml (1mg/ml) Acetyl-choline was simultaneously added and the drum was allowed to run for 30 seconds. The response was recorded. Then the drum was stopped and the Acetyl-choline solution and test drug solutions were washed out. Then the

above experiment was done for 0.2ml dose of Acetyl-choline. The drum was allowed to run for 30 seconds. The response was recorded.

Then 0.2ml of Atroprine and 0.2ml of Acetyl-choline was added and the drum was allowed to run for 30 seconds. There is no elevation in the graph and it seems to be baseline. Then 0.2ml of Acetyl-choline was added to standardise the tissue. Then the tracing was labelled and fixed.

Inference

From the graph it is inferred that the test drug antagonise the effect of Acetyl-choline when added together. So the drug has got **significant** anti-spasmodic activity.

ANTI – HISTAMINE STUDY OF SADAMANJII VER CHOORANAM ON ISOLATED GUINEA PIG ILEUM

Aim

To study the anti-histamine effect of sadamanjii ver chooranam on isolated Guinea pig ileum.

Preparation of the Test drug

1 gm of Sadamanjil ver choornam was dissolved in 10ml of water and boiled for 15 minutes. The filtrate was used for the experiments.

Solutions required

Histamine – 1 in 1,00,000 strength.

Anti histamine (pheniramine maleate 22.75mg/ml).

Test drug (Sadamanjil Ver Choornam) 100mg/ml.

Nutrient Solution

Tyrode – 1 to 2 litres.

Tissue Used

Guinea Pig ileum.

Apparatus Required

Student's organ bath.

Sherrington rotating drum.

Procedure

An overnight fasted Guinea pig weighing about 400 gms was sacrificed by a blow on the head and by carotid bleeding. The abdomen

was suddenly opened and ileo caecal junction was found out. A Small piece of ileal portion was cut and placed in a dish containing warm aerated **Tyrode Solution**. The lumen of the ileum was gently rinsed out by pushing **Tyrode Solution** into it, 3cm length segment was cut from this part of ileum, and was tied with thread on both ends without closing the Lumen and the tissue was mounted in the organ bath containing **Tyrode Solution** maintained at 37°C and bubbled with air by an oxygen tube.

First the drum was allowed to run for 1 minute from the baseline. Drugs were given to study the inhibiting effect of Histamine. 0.2ml (10mg/ml) of Histamine was added and allowed to run the drum for 30 seconds. Thus the tissue was standardised and then the drum was stopped and the Histamine was washed out.

Again the **Tyrode Solution** was added to the organ bath till the lever comes to the baseline. The drum was allowed to run for 1 minute.

To the organ bath 1ml of test drug and 0.2ml (10mg / ml) Histamine was simultaneously added and the drum was allowed to run for 30 seconds. The response was recorded. Then the drum was stopped and the histamine solution and test drug solutions were washed out. Then the above experiment was done for 0.2ml dose of histamine. The drum was allowed to run for 30 seconds. The response was recorded.

Then 0.2ml of Anti-Histamine and 0.2ml of Histamine was added and the drum was allowed to run for 30 seconds. There is no elevation in the

graph and it seems to be baseline. Then 0.2ml of Histamine was added to standardise the tissue. Then the tracing was labelled and fixed.

Inference

From the graph it is inferred that the test drug antagonise the effect of Histamine when added together. So the drug has got **significant Anti-Histamine activity**.

CLINICAL ASSESSMENT

A clinical trial was done on 30 cases of different age and of both sexes. They were clinically diagnosed as Eraippu Erumal, according to the Siddha literatures. Among them 20 patients were treated in the out – patient department and 10 patients were treated in the In-patient department.

Patients were thoroughly examined, enquired and all the clinical features, complete history, hygienic conditions, surroundings, occupation were noted. personal habits, previous illness, dietary details and allergy to specific things, if any were recorded.

They were of different severity of signs and symptoms like difficulty in breathing, cough with expectoration, wheezing, sneezing, tightness of chest and sometimes having other upper respiratory tract diseases. The duration of illness was also variable.

The routine blood and urine investigations were done in each case.

Mantoux, sputum for AFB, and radiological investigations were carried out to rule out other causes and diseases.

The cases were screened as per the following criterias and selected from the outpatient and In- patient departments of the Government siddha medical college hospital, Palayamkottai.

During the course of clinical study, other ailments, if any occured were treated with conventinal siddha medicines.

Including criteria in the case of Eraippu Erumal

- 1. Cough Nocturnal cough, paroxysm of cough.
- 2. Difficulty in breathing
- 3. Expectoration
- 4. Sputum colour and quantity without gross abnormalities such as blood stained sputum, abnormally large quantities of sputum etc.,
- 5. History of Allergy
- 6. Sneezing
- 7. Allergic rhinitis
- 8. Differential count, especially Eosinophilia
- 9. Respiratory system examination added sounds Rhonchi
- Radiological investigation Normal study, Bronchitis, chronic bronchitis.

Excluding criteria in the case of Eraippu Erumal

- 1. Facial puffiness
- 2. Abdominal distension
- 3. Pedal oedema
- 4. Hepatomegaly
- 5. Haemoptysis

- 6. Haematemesis
- 7. Orthopnoea
- 8. Cyanosis
- 9. Evening rise of temperature
- 10. Sputum for AFB positive
- 11. Mantoux positive
- 12. clubbing
- 13. Albuminuria
- 14. Increased blood urea and serum creatinine
- 15. Status Asthmaticus.
- 16. High fever.

Line of treatment

The drug sadamanil ver chooranam was administered internally in a dose of 1gm three times a day with hot water after food to each patient.

The duration of treatment varied from patient to patient

Diet and medical advice for Eraippu Erumal

- 1. Intake of hot water and hot foods were advised.
- 2. Advised to avoid chill water.
- 3. Advised to avoid factors which cause digestive disturbances.
- 4. Advised to avoid allergic factors.
- 5. Advised to avoid smoking and snuff.

- 6. Advised to take bath strictly in hot water.
- 7. Advised to take dinner before 8 p.m.
- 8. Advised to avoid stress.

Observation

The results were assessed on the basis of symptomatic relief obtained by the patient and clinically by daily examination in the In – patient department and subsequent visits in out-patient department.

Out of 30 cases 17 were males and the remaining cases were female patients, 9 patients had evidence of this particular disease in their family, 21 cases had history of allergy.

Almost all the patients were Labourers and farmers of poor socio economic status. Among the male patients most of them were chronic smokers.

The clinical improvements were recorded for every 5 days for the Out - patients. The clinical investigations were done for the patient before and after the treatment and the prognosis was noted.

No adverse effects were encountered during the study and there were no known contra –indications.

Result:

Among 30 cases 24 cases 80% showed good response, 6 cases 16.5% showed fair response and 1 case (3.5%) showed poor response.

TABLE ILLUSTRATING THE IMPROVEMENT AND THEIR PERCENTAGES

SI.No	Result	No.of. Patients	Percentage
1	Good	23	80%
2	Fair	6	16.5%
3	Poor	1	3.5%
4	Total	30	100

DISCUSSION

Humeral pathology ascribes Eraippu Eximal to the morbid condition of the kapha humour.

Thus the affected Kapha humour manifests as clinical symptoms like difficulty in breathing, cough with expectoration, wheezing, sneezing, chest tightness etc.,

The drug Sadamanjil Ver Chooranam selected for this study possesses bitter taste (kaippu) and hot property (veppa veeriyam). It also has expectorant and anti-spasmodic actions. Kaippu has the tendency to mitigate the harmful effects of the vitiated Kapha humour.

- ÁÕÒĐÅÒ ¾É¢ÔÀ;¼ø °¢Ò¾ ÁÕÒĐÅ;í,î ÍÕì,õ

All these factors seem to neutralise the vitiated kapha humour. This explanation is arrived on the basis of the analysis of the Gunapadam aspect of the drug which correlates with that of the pharmacological analysis and the clinical assessment.

Bio chemical analysis shows the presence of starch, amino acid and unsaturated compound. Their presence augments the therapeutic value of this drug by providing indispensable nutrition.

Pharmacological analysis shows that this drug has got significant

Anti spasmodic and Anti – histaminic activity.

In the clinical assessment of the 30 cases selected, 80% cases showed good response, 16.5% cases showed fair response, 3.5% cases showed poor response.

The improvement was proved by the alleviation of the signs and symptoms present before the treatment. During the clinical trail, the patients showed no adverse reactions and had no contra indications.

SUMMARY

The drug sadamanjil ver chooranam has been taken to establish the efficacy in treating Eraippu Erumal. The dose of sadamanjil ver chooranam is 1 gram thrice daily with hot water after food.

A brief description pertaining to its botanical aspect, phyto chemical constituents and Gunapadam aspect has been done.

A review of literatures about the drug and its significance in medicine since ancient period has been done.

Collected information from various literatures has been referred.

Bio chemical analysis shows the presence of starch, Amino acids & unsaturated compounds.

Pharmacological analysis shows that the drug has got significant anti- spasmodic activity and anti – histamine activity.

From the clinical assessment, it is inferred that sadamanjil ver chooranam posses remarkable efficacy in treating Eraippu Erunal and the drug has no contra- indications and has no adverse reactions.

CONCLUSION

It is concluded that the drug sadamanjil ver chooranam is an effective drug in relieving the severity of the symptoms in Eraippu Erumal and it has no adverse reactions.

INTRODUCTION

The siddha system of medicine constitutes one of the ancient Medical Practices. Siddhars have classified the source of materials for drugs as vegetable i.e. plant section, metal and mineral section and Animal section.

The drug Amaiodu (Turtle shell) is used for medicinal purposes in siddha system from the period of Bohar. Much references can be seen in his manuscripts especially in Bohar.

Amaiodu is used for perunkazhichal, seethakazhichal (dysentery), Girhani (sprue), Moolam(piles), manthram (Infants diarrhoea)

In this dissertation study, the author has taken **Amaiodu Purpum** for the treatment of **Eraippu Erumal**.

AIM AND OBJECTIVE

The main aim and objective of this dissertation work is to do a scientific review of Amaiodu parpam and its efficacy in treating Eraippu Erumal.

Eraippu erumal is one of the common respiratory disorders affecting millions of people all over the world. Amaiodu parpam has been suggested as an effective drug for Eraippu Erumal in ANUBOGA VAIDYA NAVANEETHAM PART III (PAGE NO:115). This study is aimed at the specific target of exploring the medicinal aspects of the Amaiodu parpam. So far no scientific analysis has been done in this regard. The modern science is growing every day by leaps and bounds. It is imperative to use this knowledge to bring out the efficacy of this drug.

The author has in depth dealt with

- 1. Zoological aspect
- 2. Gunapadam aspect
- 3. Bio-chemical analysis
- 4. Pharmacological analysis
- 5. Clinical assessment of Amaiodu parpam in this dissertation study.

REVIEW OF LITERATURES

ZOOLOGICAL ASPECT

ஆமை (TURTLE)

Zoological Name: Chelonia Turtle

Tamil name : ஆமை

English name : Turtle

Hindi : Kachakru

Gujarathi : Kachoo

Malayalam : Lisk, Kurakura, Kulifpaun.

- Indian Materia Medica A.K. Nadkarni Vol III Print 1993

Scientific Classification

Kingdom : Animalia

Phylum : Chordata

Class : Sauropsida

Order : Testudines Linnaeus, 1758

Suborders : Cryptodira pleurodira

Family : Chelonidae

- Outlines of zoology

J.Arthur thomson M.A., L.L.B P.No: 686-690

8th Edition October 1929

TURTLE

Zoological Aspect

Introduction

Turtles are found throughout the world on all continents and in all oceans except Antarctica. There are 247 species of turtles far lower than Snakes and Lizards, which belong to 75 genera in 13 families.

Turtles appeared in the fossil record over 200 million years ago, and were on earth long before mammals and other forms of present day reptiles. They occur in terrestrial, freshwater aquatic, semi aquatic and marine environments. They range in size from 11cm to 185cm and one species can weigh close to a ton, making (The leather back sea turtle) it the world's largest reptile.

Turtles are reptiles of the order Testudines (all living turtles belong to the crown group chelonia), most of whose body is shielded by a special bony or cartillagenous shell developed from their ribs.

The order Testudines includes both extant and extinct species, the earliest turtles being known from the early Triassic period, making turtles one of the oldest reptile groups, and a much more ancient group than the lizards and Snacks. About 300 species are alive today.

Classification of Chelonia:

1. Athecae

Vertebra and ribs free from carapace. Skull without descending processes from parietals.

Sphargidae, Leathery – skinned Turtles, with flexible carapace. Spargis coriacea, the only living species, the largest modern Chelonian, sometimes measuring 6ft in length. It is a widely, but now sparsely distributed in intertropical seas, and is said to be herbivorous.

2. Thecophora

Dorsal vertebrae and ribs fused in the carapace. Parietals prolonged downwards, including the following and other families.

Chelonidae, marine turtles, with fin like feet, and partially ossified carapace. They occur in intertropical seas and bury their soft- shelled eggs on sandy shores. The green turtle is much esteemed as food; the hawk's bill turtle furnishes much of the commercial tortoise shell.

Testudinae, land tortoises, with convex perfectly ossified carapace, and feet adapted for walking. They are found in the warmer regions of both the old and the new world, but not in Australia. In diet they are vegetarian. The common tortoise and the nearly estimated giant tortoises of the mascarene and Galapogas islands are good representatives. The latte may reach the age of the 150 years.

- Outlines of Zoology

J. Arthur Thomson M.A, L.L.D P. No. 686 690. 8th Edition October 1929

Physical Description

Turtles vary widely in size, although marine turtles tend to be relatively big animals. The largest chelonian is a marine turtle, the great leather back sea turtle, which can reach a shell length of 200m and can reach a weight of over 900kg.

Fresh water turtles are smaller, with the largest species being the Asian soft shell turtle, Pelochelys Cantorii which has been reported to measure upto 200cm or 80inch (Das 1991). This dwarfs even the better known alligator snapping turtle, the largest chelonian in North America, which attains a shell length of upto 80cm and a weight of about 76kg.

Giant tortoises of the genera Geochelone, Meiolania and others were relatively widely distributed around the world into prehistonric times and are known to have existed in North and South America, Australia and Africa.

They became extinct at the same time as the appearance of Man, and it is assumed that humans hunted them for food. The only surviving giant tortoises are on the segchelles and Galapagos Islands and can grow to over 130cm in length, and weight about 300kg.

The largest ever chelonian was Archelon ischyros, a late Cretaceous sea turtle known to have been upto 4.6m long.

The smallest turtle is the speckled padloper tortoise of South Africa. It measures no more than 8cm in length and weighs about 140gm. Two other species of small turtles are the Amercian mud turtles and musk

turtles that live in an area that ranges from Canada to South America. The shell length of many species in this group is less than 13cm in length.

Neck Folding:

Turtles are broken down into two groups according to how they evolved a solution to the problem of withdrawing their neck into their shell

- The Cryptodira which can draw their neck in while folding it under their spine.
- 2. The Pleurodira which fold their neck to the side.

Head:

Most turtles and tortoises have eyes placed on the upper sides of their heads. Species of turtles that spend most of their life on land have their eyes looking down at objects in front of them.

Some aquatic turtles, such as snapping turtles and soft shelled turtles, have eyes closer to the top of the head. These species of turtles can hide form predators in shallow water where they lie entirely submerged except for their eyes and nostrils. Sea turtles posses glands near their eyes that produce salty tears that rids their body of excess salt taken in from the water they drink.

Turtles are thought to have exceptional night vision due to the unusually large amount of rod cells in their retinas. Normal daytime vision is marginal at best due to their colour blindness and poor visual acuity. In addition to daytime vision problems, turtles have very poor pursuit movement abilities, which is most likely due to the fact that pursuit

movement abilities are normally reserved for predators that hunt quick moving pray.

Turtles have a rigid beak. Turtles use their jaws to cut and chew food. Instead of teeth, the upper and lower jaws of the turtle are covered by horny ridges. Carnivorous turtles are covered by horny ridges. Carnivorous turtles usually have knife- sharp for slicing through their prey Herbivorous turtles have serrated edged ridges that help them cut through tough plants. Turtles use their tongues to swallow food, but they cann't unlike most reptiles, stick out their tongues to catch food.

Shell

The upper shell of the turtles is called the carapace. The lower shell that encases the belly is called the plastron. The carapace and plastron are joined together on the turtles sides by bony structures called bridges. The inner layer of a turtle's shell is made up of about 60 bones that included portions of the backbone and the ribs, meaning the turtles cannot crawl out of its shell.

In most turtles, the outer layer of the shell is covered by horny scales called scutes that are part of its outer skin or epidermis scutes are made up of a fibrous protein called keratin that also makes up the scales of other reptiles. These Scutes overlap the seams between the shell bones and add strength to the shell. Some turtles do not have horny scutes.

The shape of the shell gives helpful clues to how the turtle lives.

Most tortoises have a large dome, shaped shell that makes it difficult for

predators to crush the shell between their jaws. One of the few exceptions is the African pancake tortoise which has a flat, flexible shell that allows it to hide in rock crevices. Most aquatic turtles have flat, streamlined shells which aid in swimming and diving. American snapping turtles and Musk turtles have small, cross – shaped plastrons that give them more efficient leg movement for walking along the bottom of ponds and streams.

Tortoises have rather heavy shells in contrast to aquatic and soft shelled turtles which have lighter shells that help them avoid sinking in water and swim faster with more agility. These lighter shells have larger spaces called fontanelles between the shell bones. The shell of a leather back turtle is extremely light because they lack scutes and contain many fontanelles.

The colour of a turtle's shell may vary shells are commonly coloured brown, black or olive green. In some species shells may have red, orange, yellow or grey marking and these marking are often spots, lines or irregular blotches one of the most colourful turtles is the eastern painted turtle which included a yellow plastron and a black or olive shell with red markings around the rim.

Skin and Moulting

The outer layer of the shell is part of the skin, each scute (or plate) on the shell corresponding to a single modified scale. The remainder of the skin is composed of skin with much smaller scales, similar to the skin of other reptiles.

Turtles and terrapins do not moult their skins all in one go, as snakes do, but continously, in small pieces. When kept in aquaria, small sheets of dead skin can be seen in the water (often appearing to be thin piece of plastic) when it has been sloughed off, often when the animal deliberately runs itself against a piece of wood or stone. Tortoises also shed skin, but a lot of dead skin is allowed to accumulate into thick knobs and plates that provide protections to parts of the body outside the shell.

This scutes on the shell are never moulted and as they accumulate over time, the shell becomes thicker. By counting the rings formed by a stack of smaller, older scutes on top of the larger, newer ones, it is possible to estimate the age of a turtles, if you know how many scutes are produced in a year.

Limbs

Terrestrial tortoises have short, sturdy feet. Tortoises are famous for moving slowly, in part because of their heavy shell but also because of their relatively inefficient sprawling gait that they have, with the legs being bent, as with lizards rather than being straight and directly under the body as is the case with mammals.

The amphibious turtles normally have limbs similar to those of tortoises except that the feet arewebbed and often have long claws. These turtles swim using all four feet. Large turtles tend to swim less than smaller ones, and the very big species, such as aligator snapping turtles, hardly swim at all, preferring to simply walk the bottom of the river or lake. Male

turtles tend to have particularly long claws, and these appear to be used to stimulate the female while matting.

Sea turtles are almost entirely aquatic and instead of feet they have flippers. Compared with fresh water turtles, sea turtles have very limited mobility on land, and apart from the dash from the nest to the sea as hatch lings, male sea turtles normally never leave the sea. Females must come back onto land to lay eggs. The back flippers are used to dig the burrow and then fill it back with sand once the eggs have been deposited.

Anatomy

Parts of shell

There are two parts to the shell of a turtle; the upper portion is called the carapace and the bottom half is called the plastron. Both shells are actually made of many fused bones. The carapace is the fusion of about 50 bones the ribs and vertebrae. The plastron is the fusion of bones including the clavicles (or collar bones), bones between the clavicles and portions of the ribs. A bony bridge joins the carapace and the plastron along the side of the turtle. Some turtles have a movable joint usually in the plastron, which acts as a "hinge" and allows the turtle to pull the carapace and plastron together tightly, while the turtle retracts its body into the shell. Shells have a blood and nerve supply, so bleeding and pain can result if the shell is injured.

Scutes

The shells are covered with a layer of keratin (same type of material that makes up our fingernails or horse's hooves). The keratin is arranged in patches called scutes or shields. The carapace usually has 38 scutes, and the plastron, twelve to fourteen. The names and numbers of the scutes roughly correspond to the adjacent bones and body portions. The scutes, however, do not precisely overlap the bones. Instead, they are staggered which helps give the shell more rigidity. Some aquatic turtles such as soft – shelled sea turtles may have fewer bones in their carapaces and the scutes are replaced by leathery skin.

Scute patterns

Different species of turtles have scutes of different patterns and designs, and there are often individual differences among members of the same species.

Shell shape

The shell shapes of turtles differ with each species, and are often related to habitat. Most aquatic turtles are generally flatter, allowing them to move faster through the water. Tortoises, on the other hand, have carapaces that are dome – shaped.

Shell growth

As a shell grows, the number of scutes generally does not change, but their size does. In some turtles, old scutes are shed and replaced by larger, new ones. In other species, including box turtles, tortoises, and

wood turtles, scutes enlarge in diameter as new keratin is laid down. The growth rings in scutes have been used be some experts to help determine the age of a turtle. Age estimation based on growth layers, however, can be erroneous for several reasons:

- Some turtles produce multiple growth zones per year.
- Growth is determined by changes in the environment (season) so age determination by examination of growth rings would be more accurate in wild turtles, than those kept in environments which do not change significantly.
- Growth layers may wear with age, so older turtles may be estimated to be younger than they really are.

Anatomy and Physiology

The following facts apply to most species of turtles kept in captivity.

- ❖ Both the pelvic and pectoral girdles are contained entirely within the rib cage which is fused to the protective shell. The shell is a vascular bony structure which should be included when calculating drug dosages from the animal's weight.
- Sexual dimorphism exists in many species. Male tortoises have a concave plastron and male aquatic turtles usually have very long toe nails on their front feet. The tail is relatively larger in males than in females but this does not always hold true.
- Turtles lack teeth but most possess a sharp beak called a tomium.

- Turtles can live a long time and tortoises generally live longer than aquatic species. The documented record is 152 years by a Seychelles island tortoise. Some species can be aged by growth rings on the scutes. This does not hold true for many aquatic species which periodically shed their scutes.
- ❖ Turtles lack a diaphragm and since they are housed in a shell most have little or no abdominal breathing component. Most pressure changes allowing for lung expansion are accomplished by muscles in the pockets surrounding the fore and hind limbs. Aquatic species can also respire through their skin and the mucus membranes of the throat and cloaca.
- Turtles have paired kidneys and a cloacal opening for the urogenital and gastrointestinal tracts.
- Like most other reptiles, the heart has three chambers.
- All turtles lay eggs and most bury them in the earth. Some species may lay several clutches per year and females of certain species can store sperm for several years.
- ❖ The gastrointestinal tract is standard in that it includes a simple S-shaped stomach, liver, gall bladder, pancreas, spleen, small and large intestine.
- Sea turtles possess special salt glands in their head behind each eye which allow them to drink seawater.

www.wikipedia.org

Chemical Constituents:

- Calcium compounds make up about half of the tortoise plastron and turtle carapace.
- Collagen, a fibrous protein makes up about 7% of the tortoise plastron,
- There are also small amounts of fats, magnesium, trace minerals, such as zinc and vitamins, including vitamin D in the tortoise shells.
 As with other natural calcium sources, there are small amounts of lead but not enough to be of concern.

Medicinal uses of Turtle shell (in China)

The calcium content of the plastron, when used in the dosages recommended by the Chinese texts, contributes a significant amount several hundreds mgs- compared to the currently recommended nutrition levels of about 1gm of calcium.

According to the report in pao zhi, the raw tortoise shell is mainly used for treating vertigo, tinnitus, deafness, headache and convulsions, whereas processed i.e. vinegar treated tortoise shell is appropriate for treating Night sweating, weakness of back and legs, Insomnia, Heart palpitations and other disorders due to deficiency of liver and kidney.

Tortoise shell gelatin is especially used for treating Impotence, Low back pain, and Uterine breading. Turtle shell gelatin is made as a medicinal product and is also used to treat uterine bleeding; it is also used for hemoptysis associated with Tuberculosis, but is not indicated for the kidney deficiency symptoms of back pain and impotence.

Tortoise shell is also essential in the treatment of late stage Rheumatois arthritis. It is possible that gelatin polypeptides (fragments with partial digestion) contribute to inhibition of bleeding. So useful in treatment of uterine breeding associated with uterine fibroids.

The bone disease – rickets, which is due to impaired deposition of bone calcium in children, has been treated in china with shell formulas,

Recently interest has developed in the ability of ingested collagen to inhibit Arthritis, and for its ability to inhibit angiogenesis as a means of inhibiting tumor growth.

In Parkinson's disease

The treatment of difficult and recalcitrant diseases with Chinese herbs, formulas for treating parkinson's disease, frequently include tortoise shells and othe gelatins.

In Aplastic Anaemia

Gelatins from tortoise, turtle, antler or donkey skin are prescribed in some formulas for the treatment of Aplastic Anaemia.

Another example is the use of Buxue Tang (blood nourishing decoction) plus Buxue san (blood nourishing powder) used in a study of treatment for Aplastic Anaemia. The decoction includes Turtle and Tortoise shell.

In thrombocytopenia

The treatment of difficult and recalcitrant diseases with Chinese

herbs, formulas listed for treating this disorder i.e. Thrombocytopenia

include ciyhus Cangxue fang, which is comprised of Tortoise shell, oyster

cell, and herbs to clear heat (Phellodendron, Lycium, Gardenia), stop

bleeding and tonify deficiency (Lycium, Eucommics)

- From www.wikipedia.org

Vaccine from Tortoise

This is a cure recommended for consumption. The report of the

commission appointed in Germany to examine the efficiency of Dr.

Friedman's vaccine for treatment of Tuberculosis says;"The vaccine is

valuable in the antituberculosis struggle as having given surprising results

after 1 or 2 injections". The vaccine is composed of the pure cultures of the

tubercle bacilli of the tortoise.

Indian Materia Medica – A.K.Nadkarni

Vol III P; no : 154, Print 1993

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REVIEW OF LITERATURES

GUNAPADAM ASPECT

ஆமை ஒரு சீவப்பொருள். இத "உபரசம்" எனப்படும் தாதுப்பொருளின் பிரிவின் கீழ் வருகிறது. தாதுப்பொருள் 4 பிரிவுகளை உடையது.

- 1. உலோகம் 11
- 2. காரசாரம் 25
- 3. பாடாணம் 64
- 4. உபரசம் 120.

ஆக மொத்தம் தாதுப்பொருள்கள் 220 ஆகும் இப்பிரிவில் "உபரசம்" என்ற பிரிவில் ஆமை வருகிறது. இங்கு உபரசம் என்ற சொல்லால் வழங்கப்படும் இரசம் என்ற சரக்கு சிவம் தனித்தும், பல்லுயிர்களில் கலந்தும் இருப்பது போன்று தனித்தும், ஒவ்வாரு சரக்கிலும் அதனதன் தன்மைக்கேற்ப " சரக்கிற் கலந்திடு சீவன்" எனக் கலந்தும் இருப்பதாகக் கூறப்படுகிறது.

ஆகையால் இரசம் என்ற சொல் சிவமென வழங்கப் படுவதாலின், அச்சிவம் கலவா உயிர் இல்லாதது போல இரசம் கலவா சரக்கு இல்லை. 120 உபரசச் சரக்குகளில் இரசமானது காணமுடியாத குறைந்த அளவில் காணப்படுவதால் "உபரசம்" என்றாயிற்று.

உபரசத்தின் வகைகளாவன

	கண்டு6	ிகாள் ளு	நபரசத் தி ன்	வகையைச்	சொல்வேன்
<u>ഉ</u>	₋ண்டுநா	கரவண்	ரடு கருவன	ர்டாமை	

- போகா் காரசாரத்துறை.

ஆமை

CHELONIA TURTLE, TORTOISE

ஆமை மேல் ஓடு பொருந்தியுள்ள ஓர்வகை நீர்வாழ் பிராணி. நீர் வற்றிய காலத்து இது மணலில் புதைந்து கிடக்கும்.

> - T.V.சாம்பசிவம் பிள்ளை மருத:துவ அகராதி முதற்பகுதி ப.எ: 367.

வேறு பெயர்கள்:

- ❖ கூர்மம்
- 🌣 கூனன்
- 💠 கமடம்
- ❖ கமடாதரி
- ❖ கச்சபம்.
 - குணபாடம் தாதுசீவ வகுப்பு ப.எ : 434. நான்காம் பதிப்பு 1992
- 🌣 கூன்னன்
- 🌣 கச்சவம்.
 - தமிழக சித்த வைத்திய குருகுலம் "ராஜயோகி"
 A.P.செல்வராஜன் R.I.M.P, R.H.M.P ப.எ.102.
 இரண்டாம் பதிப்பு ஆகஸ்ட் 2001

"ஆமையுடைய பெயர்தனையே அறையக்கேளு

அந்தரங்கத்தி யானியாம் நீர்ப் பரிசையாகும்

ஊமையாந் தனித்தாம் மோதரன் தான்

உந்தங்க மூடியே மவுனியாகும்

பாமையாம் பஞ்சாங்கக் கூர்மையுடன் உள்ளோன்

பாரமாங் கச்சனாங் கூர்மராசன்

வாமையால் ரசப்பிர திட்டை யுள்ளோன்

வகுத்த தெல்லாம் ஆமையுட பேருமாமே".

- போக முனிவா் நிகண்டு - 1200

பதிப்பாசிரியர் : எஸ்.பி.இராமச்சந்திரன்

ப.எ : 17.

பதிப்பு மே 1992

- **≭** அந்தரங்கத்தியானி
- ★ நீர்ப்பரிசை
- ★ தனித்தது
- ★ மோதிரன்
- ≭ அங்கமுடி
- ∗ மௌனி
- \star பஞ்சங்கம் கூர்மையுள்ளோன்
- ∗ கச்சன்
- **★** கூர்மைராசன்
- \star மால் பிரதிஷ்டையுள்ளோன்.

வகைகள்:

1. நிலத்தாமை - Land Tortoise - Testudinidae

2. நீராமை - Water Tortoise - Emididae

3. கடலாமை - Sea Tortoise - Trionycidae

Marine turtle

4. பறையாமை - Black Tortoise - Batagurellitti

- T.V.சாம்பசிவம்பிள்ளை மருத்துவ அகராதி - முதல் பாகம் ப.எ: 367.

ஆ. 1. நீர் ஆமை

2. நில ஆமை

என இருவகைப்படும்.

முதலில் கூறப்பட்ட நீா் ஆமையே மருத்தவத்திற்கு பயன்படுகிறது. இது அதிகமாக தென்னிந்தியக் கடற்கரையிலும், மன்னாா் குடாவிலும் கிடைக்கிறது நில ஆமை கருமை நிறம் உடையது.

கடலாமை திருஷ்டியினால் குஞ்சு பொரிக்கும் (திரீட்சணம்) ஜீவப் பொருள்களில் ஒன்றாகக் கருதப்படுகிறது.

இதனை,

"கடலிலே திரியுமாமை கரையேறி முட்டை யி ட'டுக் கடலிலே திரிந்தபோத குஞ்சான வாறு பொல". என்று சிவவாக்கியார் பாடலால் உணரலாம்.

> - குணபாடம் தாதுசீவ வகுப்பு ப.எ. 434 நான்காம் பதிப்பு 1992

பயன்படும் உறுப்பு

ஓடு தோல்

கறி முட்டை

கொழுப்பு பித்தநீர்

ஆமை ஒடு

பொதுகுணம்

"வல்லை கவுசையுடன் மாந்த சரம் பேதி

தொல்லையுறும் மூலந் தொலைந்துபோம் - எல்லையிலாத்

தீமைதரு முட்சூடுந் தீவிரமா யோடிவரும்

அமை முது கோட்டா லறி."

குணம் : (இ - ள்)

ஆமையோட்டினால் வல்லை, கவுசை என்னும் கட்டிகள், பாலருக்கு காணகின்ற மாந்தம் , சுரம் , பேதி , மூலரோகம், தேகவெப்பம் முதலியவைகள் தீரும் என்க.

செய்கை

உதரவாதஹரகாரி தாதுவர்த்தினி

கபஹரகாரி பசித்தூண்டி

சுரஹரகாரி

- சித்த வைத்திய பதார்த்த குணவிளக்கம்

தூது-சீவ வர்க்கம்

ப.எண் : 193. பதிப்பு 1997

ஆமையோட்டு சுத்தி

பூநீறு, கற்சுண்ணாம்பு சமனெடை சேர்ந்து அதற்கு எண்மடங்கு நீர் விட்டுக் கலக்கி தெளிவெடுத்து , அதில் ஆமை ஓட்டைச் சிறு சிறு துண்டுகளாய் நறுக்கி இட்டு எண்ணெய்க் கசிவு நீங்கும் வரை எரித்தெடுத்து, சுத்த நீரில் கழுவி எடுக்கச் சுத்தியாகும்.

உபயோகிக்கும் முறைகள்

ஆமை ஓடு , சிறப்பாக குழந்தைகள் நோய்க்குப் போடும் குடிநீராகவும் , கருக்காகவும் , பற்பமாகவும் , மாத்திரையாகவும் கையாளப்படுகின்றது.

ஆமையோட்டுக் கருக்குக் குடிநீர்

"கூனன் முதுகு வசம்புள்ளிக் கூறும் ஓம நுணாவேலி மானே பொடுதலை வாரடையும் வளரும் வெற்றிலை-யின் காம்பும் கானே கிராம்பு கருக்கியிட்டுக் காய்ச்சிச் சவலைப் பிள்ளகைக்குக் தானே கொடுக்கப் போர் மாந்தம் தங்கா தோடிப்போய்விடுமே".

(பொ- ள்) ஆமை ஓட்டைச் சுட்டதூள், சுட்ட வசம்பு, வெள்ளுள்ளி, ஓமம், நுணா இலை , வேலிப்பருத்தி, பொடுதலையிலை, வெற்றிலைக் காம்பு, கிராம்பு இவைகள் வகைக்கு வராகணெடை ஒன்று (4.2கிராம்) எடுத்துச் சட்டியிலிட்டுக் கருக்கி விதிப்படி குடிநீர் செய்து, காலை , மாலை இரு வேளையும் 3 - 5 நாள் கொடுக்கப் "போர் மாந்தம்" முதலிய மாந்த நோய்கள் நீங்கும்.

2. ஆமையோட்டுக் கருக்கு

520மிகி வரை, தேனிலாவது , தாய்ப்பாலிலாவது கொடுக்க, மாந்தம், கணம், அஜீரணபேதி, மந்தவாந்தி, கணமாந்தம் தீரும்.

3. ஆமையோட்டு பற்பம்

65மிகி - 130மிகி வரை தாய்ப்பால், பசும்பாலில் தர, குழந்தைகளின் மாந்தம், கண மாந்த பேதி தீரும்.

4. ஆமையோட்டு மாத்திரை

உளுந்தளவு மாத்திரை - 65மி.கி முலைப்பால், பால், வெந்நீர், மாந்தக் குடிநீரில் கொடுக்க, உப்புமாந்தம், கணமாந்த சுரம், பசியின்மை, அசீரணபேதி முதலியன தீரும்.

> - குணபாடம் தாதுசீவ வகுப்பு (ப.எ: 435 - 436) நான்காம் பதிப்பு 1992

5. ஆமையோட்டு சத்து

ஆமை ஒட்டினின்று தயாரித்து சாலவித்தையில் உபயோகிக்கும் ஓர் பொடி மருந்து.

> - T.V.சாம்பசிவம்பிள்ளை மருத்துவ அகராதி முதல் பாகம் ப.எண்: 367

ஆமைஓடு சேரும் பிறமருந்துகள்

1. உத்தாமணி மாத்திரை

குழந்தைகளுக்கு வேளைக்கு 1 மாத்திரை (கடலைபிரமாணம்) வீதம் தினம் இருவேளை தேன் (அ) முலைப்பாலில் கொடுத்துவர, பேதி, வாந்தி, அசீரணம், நெஞ்சுக்கபம் தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் ப.எ. 148.

2. குழந்தைகளுக்கு மாந்தக் கியாழம்

குழந்தைகளுக்கு வேளைக்கு 1-2 தேக்கரண்டி தினம் 3 முதல் 4 வேளை கொடுத்துவர, சுரம், கபத்தினுபரி தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம் ப.எ.86.

3. மஹாபூபதி பற்பம்

1-2 குன்றிஎடை (130 - 260 மிகி) நெய், வெண்ணெய் இவற்றில் தினம் இரவேளை உண்ண மார்புநோய், ஈளை, ஷயம் தீரும்.

- சிகிச்சாரத்ன தீபம் என்னும் வைத்திய நூல் ப.எ. 227.

பதிப்பு 1991

4. ஆமையோட்டு பற்பம்

½ - 1 குன்றி வரை (65 - 130மிகி) நெய்யில் தர மார்பு நோய் , அசீரணம் தீரும்.

- சிகிச்சாரத்ந தீபம் என்னும் வைத்திய நூல் ப.எ. 216

பதிப்பு 1991

5. மாந்த மாத்திரை

பயறு (அ) உளுந்தளவு மாத்திரையை தேன் , இஞ்சி சுரசம் (அ) மாந்த கியாழத்தில் தினம் ஒரு வேளை ஒரு நாள் விட:டு ஒரு நாள் மூன்று முறை கொடுக்க, சுரம் , மாந்தம் சலதோஷம் தீரும்.

- கண்ணுசாமி பரம்பரை வைத்தியம்ப.எ.130.

6. மாந்த மாத்திரை

பயறளவு மாத்திரையை தாய்ப்பாலில் காலையில் ஒரு வேளை கொடுக்க பேதியாகும் . இதனால் சுரம், சீதளம் , வலி , மாந்தம் தீரும்.

> - சித்த வைத்திய பதார்த்த குண விளக்கம் (தாது - சீவ வர்க்கம்) ப.எண்: 194. பதிப்பு 1998

7 . சண்டமாருத குழம்பு

50 - 100மிலி ஒருவேளை 5 நாட்களுக்கு கொடுத்து 5 நாள் இடைவெளி விட்டு நோய் தீரும் வரை கொடுக்க , மேகம், வாதம், சர்மநோய் தீரும்.

> ் தமிழக சித்த வைத்திய குருகுலம் "இராஜயோகி" A.P.செல்வராஜன் R.J.M.P., R.H.M.P ப.எ: 436. 2ம் பதிப்பு ஆகஸ்ட் 2001

ஆமையின் பிற உறுப்புகளின் மருத்துவ பயன்

அ) ஆமைக்கறி

குணம்

"ஆமைக் கறியருசி யையம்பித் துண்டுமூலஞ் சேமக் கனல்சூதி காமேக - நாமரத்தப் பேதி கிராணிமலப் பேதியகற் றும்பசியுந் தாதுமே ருங்கொடுக்குந் தான்"

பொருள்

ஆமைக்கறி , அரோசகம், கபம் , பித்தம், வன்மூலம், உட்சூடு , கா்ப்பமேகம்இ , இரத்தாதிசாரம், சீதக் கிரகணி, மலபேதி இவைகளை நீக்கும். பசி , வீாியம், அழகு இவற்றை தரும்.

கறியைப் பாகப்படி சமைத்துண்ண மேலே கூறப்பட்ட பிணிகள் நீங்கும்.

ஆமைக்கறியுடன் ,சீந்தில் உப்பு கூட்டிச் சமைத்து அருந்த உடலில் உண்டாகின்ற காயங்கள் நீங்கும்.

இதனை,

"காயத்தி லேபிறந்த காயத்தைப் போக்கிமிக்க மாயத்தைப் போலே மயக்குதலா - லோயா அமுதவல்லி யுப்புடனே யாக்கியமை யான கமடமெனு மாமையதி கம்" என்ற தேரன் பொருட்பண்பு நூலில் கூறப்பட்ட வெண்பாவால் உணர்க.

- குணபாடம் தாது சீவ வகுப்பு ப.எ. 437.

நான்காம் பதிப்பு 1992

ஆமைக்கறியில் அரோசகம், கபநோய்கள், பித்த உஷ்ணம், உள்மூலம், உட்சூடு, இரத்ததாதிசாரம் (இரத்தப்போக்கு) , மலபேதி இவைகளை நீக்கி பசி,வீரியம், அழகு இவைகளை உண்டாக்கும்.

> - தமிழக சித்த வைத்திய குருகுலம் ப.எ. 102. இரண்டாம் பதிப்பு ஆகஸ்ட் 2001

ஆமைக்கொழுப்பு

1. செய்கை

- 💠 உடற்றேற்றி
- 💠 காமம் பெருக்கி
- 💠 தாது வெப்பகற்றி.

இதனை உருக்கி நெய்யாக்கி, ஒன்று (4மிலி) முதல் இரண்டு (8மிலி) தேக்கரண்டி வரை அருந்திவர , கண்டமாலை, பாண்டு , நுரையீரலைப் பற்றிய பிணிகள் நீங்கும்.

இதனால் இரத்தமூலம் நீங்கும் என்பதை " ஊணாமை நெய் மூலமாட்டும் " என்ற அடியால் தெளிக.

ஆமைத்தோல்:

பெண்ணாமைத் தோலினால் பெருஞ்சிரங்கு, குழி விரணங்கள் நீங்கும் எனக் கூறப்பட்டுள்ளது.

இதனை,

"ஆனைச் சிரங்குமுத லான ரண நோயை யூனிற் படாம லொதுக்குமே - மானப் பிளிறின் குரல்காட்டும் பெண்ணாமை யான துனியின் புறவோட்டுத் தோல்". என்ற வெண்பாவால் உணரலாம்.

வழக்கிலின்மையின் இதனை உபயோகிக்கும் வகை தெரியவில்லை.

உலர்ந்த தோலைப் பொடித்து நெருப்பிலிட்டுப் புகை பிடிக்க மூலம் நீங்குமென்பர்.

ஆமை முட்டை

முட்டையைப் பொரித்து குழந்தைகளுக்குக் கொடுக்க கக்குவான், வலி முதலிய நோய்களுக்குக் கொடுத்தல் பழக்கம்.

ஆமை - பித்த நீர்

இதனை நசியமிட வலி நீங்குமென்றும்,

கடிவாயில் பூச , விடமிறங்குமென்றும், மேலே பூச , தொண்டை வலி நீங்குமென்றும் கூறப்பட்டுள்ளன.

ஆமை - இரத்தம்

குருதியை அருந்துவதினால் புப்புச சம்பந்தப்பட்ட பிணிகள் நீங்கும். இது தூத்துக்குடியில் வழக்கிலிருப்பதைக் காணலாம்.

- குணபாடம் தாது சீவ வகுப்பு

- நான்காம் பதிப்பு 1992 பக்க எண் 437

MATERIALS AND METHODS

(Amaiodu parpam)

Collection of the drug

Amaiodu was collected from Gopalan Asan raw drug store, Nagercoil after identification.

Purification of the raw drug

Equal amount of pooneeru and karchunnam were taken in a mudpot. They are mixed with water at the ratio of 1:8. Then the cut pieces of Amaiodu were put in that mixture & boiled well till the impurities of the drug are removed. After that the drug (Amaiodu) becomes soft and white. The drug taken out then and washed out in clean water &dried in sunlight.

Preparation of Amaiodu Parpam

Drugs required

Amaiodu (purified) – 87.5gm. (2.5 palam)

Rubbed Adathoda leaves – 700gm. (20 palam)

Method of preparation

Adathoda leaves were rubbed in a kalvam to make a karkam.

Then the purified cut pieces of Amaiodu were buried in the Adathoda leaves karkam. Two mud seelai were made on that karkam. Then it was dried well. After that it was subjected to pudam with 1600gms, Of varatti (8 veesai). Parpam obtained was made into a fine powder by grinding in the kalvam.

Dose:-

250mg two times a day with honey after meals.

Route of administration:-

Enteral.

The prepared Amiodu Parpam was used for the treatment of Eraippu Erumal was analysed by the following methods.

- 1. Bio chemical analysis.
- 2. Pharmacological analysis
- 3. Clinical Study.

BIO - CHEMICAL ANALYSIS

BIO - CHEMICAL ANALYSIS OF AMAIODU PARPAM

PREPARATION OF THE EXTRACT

100mgs of parpam is weighed accurately & placed into a clean beaker and added a few drops of concentrated hydrochloric acid and evaporated it well. After evaporation cooled the content and added a few drops of conc. Nitric acid and evaporated it well. After cooling the content add 20ml of distilled water and dissolved it well. Then it is transferred to 100ml volumetric flask and made up to 100ml with distilled water. Mix well filter it. Then it is taken for analysis.

Qualitative Analysis

S.no	Experiment	Observation	Inference
1.	Test for calcium 2ml of the above prepared extract is taken in a clean test tube. To this add 2 ml of 4% ammonium oxalate solution.	A white precipitate is formed.	Indicates the presence of calcium.
2.	Test for sulphate: 2ml of the extract is added to 5% barium chloride solution.	A white precipitate is formed.	Indicates the presence of sulphate.

3.	Test for chloride The extract is treated with silver nitrate solution.	A white precipitate is formed.	Indicates the presence of chloride.
4.	Test for carbonate The substance is treated with concentrated Hcl.	A brisk effervescence is formed.	Indicates the presence of carbonate.
5.	Test for zinc The extract is added with potassium ferro cyanide.	A white precipitate is formed.	Indicates the presence of zinc.
6.	Test for iron Ferric The extract is treated with concentrated glacial acetic acid and potassium ferro cyanide.	No blue colour is formed.	Absence of ferric iron.
7.	Test of iron: Ferrous: The extract is treated with concentrated Nitric acid and ammonium thio cynate.	Blood red colour is formed.	Indicates trace of ferrous is present.
8.	Test for phosphate The extract is treated with ammonium molybdate and concentrated nitric acid.	Yellow precipitate is formed.	Indicates the presence of phosphate.
9.	Test for albumin The extract is treated with Esbach's reagent.	No yellow precipitate is formed.	Absence of albumin.

10.	Test for Tannic acid	No blue black	Absence of
	The extract is treated with ferric	precipitate is	Tannic
	chloride.	formed.	acid.
11.	Test for unsaturation	It does not	Absence of
	Potassium permanganate solution is	get	unsaturated
	added to the extract.	decolourised.	compound.
12.	Test for the reducing sugar 5ml of benedict's qualitative solution is taken in a test tube and allowed to boil for 2 mts and added 8-10 drops of the extract and again boil it for 2 mts.	No colour change occurs.	Absence of reducing sugar.
13.	Test for amino acid: One or two drops of the extract is placed on a filter paper and dried it well. After drying, 1% ninhydrin is sprayed over the same and dried it well.	No violet colour is formed.	Absence of amino acid.

Inference

The given sanple of Amiaiodu Parpam contains calcium, Sulphate, Chloride, Carbonate, Zinc, Phosphate and Trace of ferrous iron.

PHARAMACOLGICAL ANALYSIS

ANTI SPASMODIC EFFECT OF AMAIODU PARPAM ON ISOLATED RABBIT ILEUM

Aim

To find out the anti – spasmodic effect of Amaiodu parpam on isolated rabbit ileum.

Preparation of the Test drug

100 mg of the Amaiodu parpam was dissolved in 10 ml of water and boiled for 15 minutes. The filterate was used for the experiments.

Solutions required

Acetylcholine – 1mg/ml. Atropine – 0.5mg/ml

Test drug (Amaiodu parpam) – 10mg/ml.

Nutrient Solutions

Tyrode – 1 to 2 liters.

Tissue used

Rabbit ileum.

Apparatus Required

Student's organ bath, sherrington rotating drum.

Procedure

A Rabbit was starved for 48hours and was allowed water ad – libtum. It was sacrificed by a blow on the head and by carotid bleeding. The abdomen was quickly opened and the ileo - caecal junction was found out. A small piece of ileal portion was cut, removed and placed in a dish

containing warm aerated Tyrode solution. The Lumen of the ileum was gently rinsed out by pushing Tyrode solution into it, 3 cms length segment was cut from this part of ileum and was tied with thread on both ends without closing the lumen and the tissue was mounted in the organ bath containing T grade solution maintained at 37°c bubbled with air by an oxygen tube.

First the drum was allowed to run for 1 minute from the base line.

Drugs were given to study the inhibiting effect of Actyl-choline. 0.2ml (1 mg / ml) of Acetylcholine was added and allowed to run the drum for 30 seconds. Thus the tissue was standardised and then the drum was stopped and the Acetyl – choline was washed out.

Again the Tyrode solution was added to the organ bath till the lever comes to the base line. The drum was allowed to run for 1 minute.

To the organ bath 1 ml of test drug and 0.2 ml (1mg / ml) Acetyl – choline was simultaneously added and the drum was allowed to run for 30 seconds. The response was recorded. Then the drum was stopped and the Acetyl – choline solution and the test drug solutions were washed out. Then the above experiment was done for 0.2ml dose of Acetyl – choline. The drum was allowed to run for 30 seconds. The response was recorded.

Then 0.2ml of Atropine and 0.2ml of Acetyl –Choline was added and the drum was allowed to run for 30 seconds. There is no elevation in the graph. Then 0.2ml of Acetyl –choline was added to standardise the tissue. Then the tracing was labelled and fixed.

Inference

From the graph, it is inferred that the test drug doesn't antagonise the effect of Acetyl-choline when added together. So the drug **does not possess Anti – spasmodic activity**.

ANTI HISTAMINE STUDY OF AMAIODU PARPAM ON ISOLATED GUINEA PIG ILEUM

Aim

To study the anti – histamine effect of Amaiodu parpam on isolated Guinea pig ileum.

Preparation of the Test drug

100mg of Amaiodu parpam was dissolved in 10ml of water and boiled for 15 minutes. The filterate was used for the experiments.

Solutions required

Histamine - 1 in 100000 strength,

Anti Histamine (Pheniramine maleate 22.75 mg / ml)

Test drug - (Amaiodu parpam) 10 mg/ ml.

Nutrient Solution

Tyrode - 1 to 2 litres.

Tissue Used

Guinea Pig Ileum.

Apparatus Required

Student's Organ bath.

Sherrinton rotating drum.

Procedure

An overnight fasted Guinea pig weighing about 400 grams was sacrificed by a blow on the head and by carotid bleeding. The abdomen was suddenly opened and ileocacal junction was found out. A small piece of ileal portion was cut and placed in a dish containing warm aerated Tyrode solution. The Lumen of the ileum was gently rinsed out by pushing Tyrode solution into it, 3cms length segment was cut from this part of ileum, and was tied with thread on both ends without closing the lumen and the tissue was mounted in the organ bath containing Tyrode solution maintained at 37°C and bubbled with air by an oxygen tube.

First the drum was allowed to run for 1 minute from the base line.

Drugs were given to study the inhibiting effect of Histamine. 0.2 ml (10mg / ml) of Histamine was added and allowed to run the drum for 30 seconds.

Thus the tissue was standardised and then the drum was stopped and the Histamine was washed out.

Again the Tyrode solution was added to the organ bath till the lever comes to the base line. The drum was allowed to run for 1 minute.

To the organ bath 1 ml of the test drug and 0.2ml (10 mg / ml) Histamine was simultaneously added and the drum was allowed to run for 30 seconds. The response was recorded. Then the drum was stopped and the histamine solution and test drug solutions washed out. Then the above experiment was done for 0.2ml dose of Histamine. The drum was allowed to run for 30 seconds. The response was recorded.

Then 0.2ml of Anti – histamine and 0.2 ml of histamine was added and the drum was allowed to run for 30 seconds. There is no elevation in the graph. Then 0.2 ml of histamine was added to standardise the tissue. Then the tracing was labelled and fixed.

Inference:

From the graph, it is inferred that the test drug doesn't antagonise the effect of Acetyl choline, when added together. So the drug **doesnot** possess Anti – Histaminic action.

CLINICAL ASSESSMENT

Eraippu Erumal is a common disease and vexes mankind constantly. To assess the therapeutic efficacy of Amaiodu parpam, Eraippu Erumal patients of both sexes were selected and treated in both Inpatient and Out patient department of Gunapadam, Government Siddha Medical College hospital, Palayamkottai.

For all cases full clinical data were recorded and they were dignosed on the basis of Siddha basic pricriples i.e, Envagai thervugal etc. All the patients were undergone for hematological and radiological investigations in the corresponding sections of the hospital.

Clinically patients were selected as 'Eraippu Erumal' having the cardinal symptoms namely,

- Tightness of chest
- Dyspnoea
- Expiratory Wheezing Rhonchi
- Cough with or without expectoration
- H/O previous attacks before etc.

To exclude Pulmonary tuberculosis & other Respiratory infections, patients with the following symptoms are omitted from the study.

- Evening rise of Temperature
- Haemoptysis
- Expectoration of large quantities of foul yellow coloured sputum.
- Emaciation of body
- Acute pyrexial illness with respiratory symptoms.

Respiratory system of patients was thoroughly examined clinically and exact cases of Eraippu Erumal were selected for the study.

Accordingly 30 patients were selected and studied both in I.P and O.P wards.

Of those 20 patients were treated as Out patients and 10 were treated as In-Patients.

Patients who could not stay for enough number of days in I.P ward were requested to come to out patient Department after their discharge from in patient Department and their progress were watched without fail.

Drug and Dosage

The test drug Amaiodu parpam was given to the patients at the dose of 250mg two times a day with honey.

Diet and Medical Advice

- All patients were advised to avoid food which increases Kapha.
- They were strictly advised not to take meat, fish, chicken, Lemon, bitter guard, Leaves of agathi, Pumpkin and those vegetables which have cooling effects on body. (e.g. chow chow, Raphanus)
- They were also advised not to take curd, butter milk, cold water for drinking.
- They were advised to avoid exposure to cold air, dust and sleeping in bare floor.
- The male patients who were having the habit of smoking were strictly advised to abstain from smoking.

TABLE ILLUSTRATING THE IMPROVEMENT AND THEIR
PERCENTAGES

SI.No	Result	No.Of Patients	Percentage
1	Good relief	18	60%
2.	Fair relief	09	30%
3.	Poor relief	03	10%
	Total	30	100

From the above table it can be observed that 60% of patients receiving the drug Amaiodu parpam had good relief, 30% had fair relief, and 10% had poor relief.

DISCUSSION

The basic abnormality in Eraippu Erumal is derangement of Kapha humour.

Thus the affected Kapha humour reflects the clinical symptoms like cough with expectoration scanty sputum and difficulty in breathing. The above signs and symptoms were partially relieved by administration of the drug Amaiodu parpam. The possible explanation is given below and the drug acts in such a way that.

Amaiodu has Expectorant action.

According to the basic principle of siddha, all these factors tend to the Pharmacological and clinical trails.

Bio-chemical analysis showed that it contains calcium, Sulphate, Chloride, Carbonate, Zinc, Phosphate, Ferrous Iron & Mercury.

They provide valuable nutritional support.

Pharmacological studies show that the Amaiodu parpam has not get significant Anti – Histamine and Anti Spasmodic Activity.

In the clinical assessment, 30 cases were selected, 18 cases showed good response, 9 cases showed fair response and 3 cases showed poor response.

During the course of study no adverse reactions were observed and there were no contra indications.

SUMMARY

Amaiodu parpam was taken for this study to establish its efficacy in Eraippu Erumal, as the important aspect of this dissertation work.

The dose of Amaiodu parpam is 250mgm two times a day with honey.

A review of literatures about the drug and its significance in medicine were collected.

In Bio-chemical analysis the drug has got Calcium, Sulohate, Carbonate, zinc, Ferrous Iron , Phosphate and Mercury.

Pharmacological analysis shows no significant anti-histamine and Anti spasmodic activity.

From the clinical study the drug had a moderate response in Eraippu Erumal.

CONCLUSION

It is concluded that the drug Amaiodu parpam has moderate effect in the treatment of Eraippu Erumal for reducing its severity and difficulties.

1.Name:	Mr.Thangavel	Age:	75	Sex:	M	Ward:		MPGII	I.P.No	2359	Occupa	tion:	Working in Rice m	nill
	servai													
Date of A	dmission	23.10.2006	j [Date of D	ischarg	e 01	1.11.00	6			No.of Days T	reated	11 Days	
Drug	Sadamanjil Ve	er Choorana	m 1 gm T	DS with	hot wate	r		Dia	ignosis	Eı	aippu Erumal			
Complain	ts & Duration		Investig	ations B	efore Tr	eatment		Investig	gations Af	ter Trea	tement		Response	
			BLOOD	-				BLOOD)-					
	h expectoration		TC -		9200Ce	ells/cu.mr	n	TC -		9000	Cells/cu.mm			
•	ire to dust, cold		DC - I	60%	L 34	% E	06%	DC -	P 60%		6% E 04%			
difficulty in	n breathing since	2 years	ESR 1	⁄ ₂ hr 2	20mm /	1 hr 40	mm.	ESR ¹	½ hr 20	mm 1	hr 40mm			
			Hb-		74%			Hb-	74%	6				
Family Hi	story		Bl.Sugai	•	83mgs9	%		Bl.Suga	ır 83r	ngs%				
					25mgs ⁹	%		Bl. Urea	a 25r	ngs%				
No signific	o significant family history presen			Cholester	ol 299ı	mgs%		Serum (Cholestero	1 295	imgs%		Good	
			URINE -					URINE	-				0 000	
			Albumin	-	Nil			Albumir)-	NII				
			Sugar -		Nil			Sugar -		Nil				
			Deposits	3 -	NAD			Deposit	s -	NAD				
			CHEST	X – RAY		Chronic Bronchitis	8	CHEST	X – RAY	Nor	mal			
			SPUTIM	FOR AF	B 1	Negative		SPUTIN	I FOR AFE	3				
			MANTO	UX	1	Negative		MANTO	OUX					
Days			1	3		6	g)	12	15				
Cough	ough			Р)	R	F	₹						
Breathles	sness		++	+		+	-							
Sput	um C	olour	W	V		W	V	V						
-	Q	uantity	10ml	8n	nl	5ml	2r							
Rhonchi			Р	Р)	R	F	₹						

2.Name:	Mr.Dhanusulal	Age:	85	Sex:	М	Ward	:	MPGI	l I.P	.No	1983	Occupa	tion:	Farmer	
Date of Ad	mission	28.8.06		Date of D	Dischar	ge 1	1.9.06		,		No.	of Days Tr	eated	15 Days	
Drug	Sadamanjil Ve	er Choorana	ım 1 gm 🛚	TDS with	hot wat	er		D	iagnosi	S	Eraipp	ou Erumal			
Complaint	s & Duration		Investig	gations E	Before ⁻	Freatmer	ıt	Inves	tigation	s Afte	r Treaten	nent		Response	
	expectoration	difficulty in	BLOOD)-				BLO	D-						
breathing s	ince 1 year.		TC -		90000	Cells/cu.m	nm	TC -			8500Cel	ls/cu.mm			
			DC -	P 60%	L 3	6% E	04%	DC -	P 60°	% I	L 36%	E 04%			
			ESR	½ hr 3	30mm	1 hr 6	88mm	ESR	⅓ hr	15m	nm 1 hr	30mm			
			Hb-		68%			Hb-		68%	-				
Family His	tory		Bl.Suga	ır	79mg	s%		Bl.Su	gar	80mg	gs%				
			Bl. Urea	ì	32mg	s%		Bl. Ur	ea	32m	gs%				
	family history p			Cholester	ol 15	7mgs%			n Choles	terol	157mg	s%		Fair	
His father h	as similar episo	ode	URINE	-				URIN	E -						
			Albumin	-	Nil			Albun			NII				
			Sugar -		Nil			Sugai			Nil				
			Deposit			oithelial c	ells	Depos			NAD				
				X – RAY		Normal			ST X – R						
				I FOR AF	FB	Negative			IM FOR	AFB					
			MANTO			Negative	-	MAN				1			
Days			1	3		6		9	12		15				
Cough			Р	F		R	ŀ	₹	R		R				
Breathless			++	+		++		+	+		+				
Sputu	m	olour	W	V		W	_	N	W		W				
-	··· Qı	uantity	10ml	10		8ml		ml	2ml		2ml				
Rhonchi			Р	F)	R	F	₹	R		R				

3.Name: Shann	nugathammal Age:	60	Sex:	F	Ward	:	FPGI	I.P.	.No	2417	Occupa	tion:	Housewife	
Date of Admission			Date of D	Discha	rge 1	4.11.0	6			No	of Days Tr		16 Days	
Drug Sada	amanjil Ver Choorana	m 1 gm ⁻	TDS with	hot wa	ter			iagnosi	S	Eraip	pu Erumal			
Complaints & Du	uration	Investi	gations E	Before :	<u>Treatmen</u>	ıt	Inves	tigations	s Afte	r Treaten	nent		Response	
		BLOOD)-				BLO	D-						
	ctoration, difficulty	TC -		90000	Cells/cu.m	ım	TC -			9000Ce	lls/cu.mm			
in breathing since		DC -	P 58%		10% E		DC -	P 559	% I	43%	E 02%			
exposure to cold	air, dust etc.,	ESR	½ hr ′	10mm	1 hr 2	22mm	ESR	½ hr	10n	nm 1 hr	22mm			
		Hb-		71%			Hb-		70%					
Family History		Bl.Suga	ır	80mg	s%		Bl.Su	gar	85mg	gs%				
		Bl. Urea		20mg			Bl. Ur		20m					
No significant fam	nily history present		Cholester	ol 15	7mgs%			n Choles	terol	157mg	s%		Poor	
		URINE	-				URIN	E	ı					
		Albumir		Nil			Albun			NII				
	-	Sugar -		Nil			Suga			Nil				
	-	Deposit		NAD	1 -		Depo			NAD				
		CHEST	X – RAY	•	Chronic		CHES	ST X – R	AY	Chroni				
	-				Bronchit					Bronch	nitis			
	_		I FOR AF	-В	Negative			IM FOR	AFB					
		MANTO			Negative		-	<u>roux</u>			1		T	1
Days		1		3	6		9	12		15				
Cough		P	F		P		₹	R		R				
Breathlessness		++		-	+		+	+		-				
Sputum	Colour	W	V		W		٧	W		W				
-	Quantity	<u>15ml</u>		ml	5ml		ml	3ml		2ml				
Rhonchi		P	F	,	R	<u> </u>	₹	R		R				

4.Name: F	Petchiammal	Age:	70	Sex:	F	Ward	:	FPGI	I.P.	.No	2497	Occupa	tion:	Housewife	
Date of Adı	mission	10.11.06		Date of	Discha	rge 2	28.11.0	6	•		No	of Days Tr		19 Days	
Drug	Sadamanjil V	er Choorana	m 1 gm	TDS with	n hot wa	ıter			iagnosi	S	Eraip	pu Erumal			
Complaints	s & Duration		Invest	igations	Before	Treatmer	nt	Inves	tigation	s Afte	r Treater	ment		Response	
			BLOO	D-				BLO	D-						
	ith expectorati		TC -		8600	Cells/cu.m	nm	TC -			8000Ce	lls/cu.mm			
	ifficulty in brea	thing since	DC -	P 50%	L	44% E	06%	DC -	P 509	% I	44%	E 06%			
4 years			ESR	1⁄₂ hr	15mm	1 hr 3	35mm	ESR	½ hr	10m	nm 1 hr	20mm			
			Hb-		64%			Hb-		64%					
Family Hist	tory		BI.Sug	ar	132m	ngs%		Bl.Su	gar	110n	ngs%				
			Bl. Ure		22mg			Bl. Ur		20mg					
significant fa	amily history p	resent		Choleste	rol 14	48mgs%			n Choles	terol	148mg	js%		Good	
			URINE	-				URIN	E						
			Album		Nil			Albun			NII				
			Sugar		Nil			Suga			Nil				
			Depos	its -	1-2pt HPF	uscells mg	ıs%	Depo	sits -		NAD				
			CHES	T X – RA	Y	Normal		CHES	ST X – R	AY					
			SPUTI	M FOR A	.FB	Negative	Э	SPUT	IM FOR	AFB					
			MANT	OUX		Negative	e	MAN	roux						
Days			1		3	6		9	12		15	18			
Cough			Р		Р	Р	I	R	R		R	R			
Breathless			+++		++	++		-+	+		+	+			
Sputur	m	Colour	W		W	W		N	W		W	W			
•	Q	uantity	20m		8ml	18ml	_)ml	5ml		3ml	2ml			
Rhonchi			Р		<u>P</u>	Р		Р	R		R	R			<u> </u>

5.Name: M	lr.Krishnan	Age:	64	Sex:	М	Ward	:	MPGI	I I.P.	.No	2538	Occupa	tion:	Farmer	
Date of Adm	nission	15.11.06		Date of	Discha	rge 2	7.11.0	6	•		No	o.of Days T		13 Days	
Drug	Sadamanjil Ve	er Choorana	m 1 gm	TDS with	hot wa	iter		D	iagnosi	S	Erai	ppu Erumal			
Complaints	& Duration				Before	Treatmen	t		tigations	s Afte	r Treate	ment		Response	
			BLOO	D-				BLOC)D-						
	expectoration (TC -		1000	0Cells/cu.r	nm	TC -			9000Ce	ells/cu.mm			
	ice 6 months,		DC -	P 70%			02%	DC -	P 709	% I	28%				
exposure to	cold air, dust e	etc.,	ESR	1⁄₂ hr	15mm	1 hr 4	2mm	ESR	½ hr	10m	m 1 h	r 20mm	1		
			Hb-		71%			Hb-		72%					
Family Histo	ory		Bl.Sug	ar	88mg	js%		BI.Su	gar	80mg	js%				
			Bl. Ure	a	18mg	js%		Bl. Ur	ea	20mg	js%				
No significan	nt family histor	y present	Serum	Choleste	rol 14	46mgs%		Serun	n Choles	terol	150m	gs%		Good	
			URINE	-				URIN	E -					0000	
			Album	== '='	Nil			Albun			NII				
			Sugar		Nil			Sugar			Nil				
			Depos			us cells		Depos			NAD				
			CHES	T X – RA	Y	Chronic		CHES	ST X – R	AY	Norm	al			
						bronchiti									
				M FOR A	.FB	Negative			IM FOR	AFB					
			MANT	OUX		Negative	1	MAN		-		1	<u> </u>	T	T
Days			1		3	6		9	12		15				
Cough			Р		<u>P</u>	р	F	₹	R						
Breathlessn			++		++	+		+	+						
Sputum	1	olour	W		W	W		٧	W						
•	Qı	uantity	15m		<u>Oml</u>	8ml	-	ml	2ml						
Rhonchi			Р		P	Р	<u> </u>	₹	R						

6.Name: Mr.Ash	iok	Age:	55	Sex:	М	War	d :	MPGI	l I.P.	No	254	5	0	ccupa	tion:	Fa	rmer		
Date of Admission		1.06	l e	Date of	Discha		27.11.0		<u>,</u>		·				reated			Days	
Drug Sada	manjil Ver Ch	oorana	m 1 gm	TDS with	hot wa	ater			iagnosis	S	Е	raipp	ou E	rumal					
Complaints & Du	ıration		Invest	igations	Before	Treatme	ent	Inves	tigations	s Afte	r Tre	atem	nent				Resp	onse	
			BLOO	D-				BLO	D-										
Cough with expec			TC -		7500	Cells/cu.	mm	TC -			750	0Cel	ls/cı	u.mm	1				
breathlessness, c			DC -	P 68%	L	28% E	E 04%	DC -	P 65%	6		2%	Е	03%					
since 1 year on ex	cposure to col	d air,	ESR	½ hr	37mm	1 hr	65mm	ESR	½ hr	15n	nm '	1 hr	3	0mm					
dust etc			Hb-		68%			Hb-		68%									
Family History			BI.Sug		75m			Bl.Su		78m	gs%								
			Bl. Ure		24m			Bl. Ur		24m									
No significant fam	ily history pre	sent		Choleste	rol 1	64mgs%			n Cholest	terol	16	4mgs	s%_				F	air	
			URINE					URIN	E										
			Album		Nil			Albun			NII								
			Sugar		Nil			Suga			Nil								
			Depos		NAD			Depo			NAC								
			CHES	T X – RAY	Y	Chroni		CHES	ST X – R/	AΥ	No	rmal							
						Bronch													
				M FOR A	FB	Negati			IM FOR	AFB									
			MANT			Negati		MAN							1,	_			
Days			1		3	6		9	12		15								
Cough			Р		Р	Р		₹	R										
Breathlessness			++		+	+		+	+										
Sputum	Colou		W		W	W		N	W										
	Quanti	ty	10m		ml	5ml		ml	2ml										
Rhonchi			Р		P	Р		₹	R										

7.Name: Mrs.Chit	ra	Age:	39	Sex:	F	Ward :		FPGI	I.P	.No	2553	Occupat	ion:	House wife	
Date of Admission		.11.06		Date of			8.11.0)6			No	of Days Tr	eated	13 Days	
Drug Sadam	nanjil Ver C	hoorana	m 1 gm	TDS with	hot wa	ter		D	iagnosi	is	Eraip	pu Erumal			
Complaints & Dur	ation		Investi	gations	Before	Treatment		Inves	tigation	s Afte	r Treater	nent		Response	
			BLOO	D-				BLO	DD-						
Cough with expector			TC -		9200	Cells/cu.mr	m	TC -			9200Ce	lls/cu.mm			
breathlessness since		•	DC -	P 60%			5%	DC -	P 60	% I	_ 35%	E 05%			
exposure to cold ai	r, dust etc.	,	ESR	1⁄₂ hr	14mm	1 hr 3	0mm	ESR	½ hr	10m	m 1 hr	20mm			
			Hb-		70%			Hb-		70%					
Family History			Bl.Sug	ar	83mg	ıs%		Bl.Su	gar	83mg	gs%				
	significant family history prese				47mg	ıs%		Bl. Ur	ea	47mg	gs%				
No significant family	significant family history prese				rol 14	l6mgs%		Serur	n Choles	sterol	146mg	s%			
	o significant family history prese							URIN	E -					Good	
			Albumi	n-	Nil			Albun	nin-		NII				
			Sugar -	-	Nil			Suga	r -		Nil				
			Deposi	ts -	few e	pithelia cel	ls	Depo	sits -		few epith HPF	nelial cells			
			CHEST	ΓX – RA`	Y	Chronic Bronchitis	s	CHES	ST X – R	AY	Norma	I			
			SPUTI	M FOR A	FB	Negative		SPU1	IM FOR	AFB					
			MANT	OUX		Negative		MAN	TOUX						
Days			1		3	6	Ç	9	12		15				
Cough	ough				Р	R	F	3	R						
Breathlessness			++		+	+	-	+	+						
Sputum	Colo	ur	W	١	W	W	V	٧	W						
-	Quan	tity	10ml	8	ml	5ml	2r	ml	2ml						
Rhonchi			Р		Р	R	F	₹	R						

8.Name: Mr	rs. Pitchammal	Age:	65	Sex:	F	Ward	: t	FPGII	I.P	.No	2590)	00	cupa	tion:	Н	ouse wi	fe	
Date of Admi	ission 23	3.10.2006	•	Date of	Discha	rge	01.11.0	6	•			No.c	•		reated		11 D	ays	
Drug S	Sadamanjil Ver (Choorana	m 1 gm	TDS with	hot wa	ter		D	iagnosi	is	Er	aipp	u Er	umal					
Complaints 8	& Duration		Investi	gations	Before	Treatme	nt	Inves	tigation	s Afte	er Trea	tem	ent				Resp	onse	
			BLOOD)-				BLOC)D-										
	xpectoration che		TC -		9600	Cells/cu.i	mm	TC -			9000	Cells	s/cu	.mm					
	iculty in breathir	ng since		P 62%	L ;	34% E	04%	DC -	P 62	% I	L 34	1%	Е	04%					
one month			ESR	½ hr	45mm	1 hr	72mm	ESR	½ hr	15m	nm 1	hr	30)mm					
			Hb-		70%			Hb-		70%									
Family Histo	ry		BI.Suga	ar	88mg	js%		Bl.Su		88mg	gs%								
			Bl. Urea		27mg			Bl. Ur		27mg							_		
significant fan	nily history pres	ent		Choleste	rol 15	9mgs%			n Choles	sterol	159)mgs	%				Go	od	
			URINE	-	•			URIN	E -										
			Albumir		Nil			Albun			NII								
			Sugar -		Nil			Sugai			Nil								
			Deposit		NAD	1		Depos			NAD								
				X – RA		Normal			T X – R										
				M FOR A	FB	Negativ			IM FOR	AFB					_				
_			MANTO		•	Negativ		MAN		-		1			1				1
Days			1		3	6		9	12		15								
Cough			<u>P</u>		P	R		R											
Breathlessne		0111	++ W	-	+ //	+ W		+ N											
Sputum	Cole Quai		vv 15ml		ov Oml	5ml		ml .											
Rhonchi	Quai	iiiiy	P		P	R		R											
Milononi			•		•	11		`											

9.Name:	Alagumuthu	Age:	691	Sex:	М	Ward	: k	MPGI	l I.P	.No	2593		Occup	ation:	Cod	olie		
Date of Ad		23.10.2006	;	Date of	Discha	rge	01.11.0	6	•		ı		Days 7		•	11 Days		
Drug	Sadamanjil V	er Choorana	m 1 gm	TDS with	hot wa	ter		D	iagnosi	S	Era	aippu	Eruma					
Complaint	s & Duration		Investi	gations I	Before	Treatme	nt	Inves	tigation	s Afte	r Treat	teme	nt			Respons	9	
	expectoration,		BLOO)-				BLOC)D-									
	reathlessness s		TC -		10800	0Cells/cu	.mm	TC -			96000	Cells/	cu.mm					
1 -	xposure to cold	air, dust		P 72%		26% E		DC -	P 72	% I	_ 26							
etc.,			ESR	½ hr	15mm	1 hr	32mm	ESR	⅓ hr	15m	nm 1	hr	32mm					
			Hb-		69%			Hb-		69%								
Family His	story		Bl.Suga	ar	80*m	gs%		BI.Su	gar	82mg	gs%							
			Bl. Urea	-	66mg	js%		Bl. Ur		42mg								
significant f	family history p	resent		Choleste	rol 16	9mgs%			n Choles	terol	169	ngs%	0			Good		
			URINE	-				URIN	E									
			Albumii		Nil			Albun			NII							
			Sugar -		Nil			Sugai			Nil							
			Deposit		NAD	T		Depos			NAD							
				X – RA)		Normal			T X – R									
				M FOR A	FB	Negativ			IM FOR	AFB								
_			MANTO			Negativ		MAN						1,	1			
Days			1		3	6		9	12		15							
Cough			Р		Р	R		₹										
Breathless		2-1	++		+	+		+				_						
Sputu	ım —	Colour	W 10ml		<i>N</i>	W		N										
Rhonchi		uantity	10ml P		ml P	3ml R		ml R				+						
KIIOHCIII			Г		ı	Г	Г	`				L_						

10.Name: Muthuve	Age:	40	Sex:	M Ward:	:	MPGI	I I.P.	.No	2422	Occupat	ion: Farr	ner	
Date of Admission	31.10.06	D	ate of Di	scharge 2	7.11.0	6	•	•	No.	of Days Tr		8 Days	
Drug Sadam	anjil Ver Choorana	m 1 gm T[OS with he	ot water		D	iagnosi	S	Eraipp	ou Erumal			
Complaints & Dura	tion	Investiga	ations Be	fore Treatmer	nt	Inves	tigations	s Afte	r Treaten	nent		Response	
Cough with expector		BLOOD-				BLOC)D-						
breathlessness since		TC -		9600Cells/cu.n	nm	TC -			9000Cel	ls/cu.mm			
exposure to dust, co	old air etc	DC - P	60%	L 36% E	04%	DC -	P 60°	% L	36%	E 04%			
		ESR ½	∕₂ hr 5r	mm 1 hr 1	1mm	ESR	½ hr	5mr	n 1 hr	11mm			
		Hb-		72%		Hb-		70%					
Family History		Bl.Sugar		114mgs%		BI.Su	gar	114n	ngs%				
		Bl. Urea		26mgs%		Bl. Ur	ea	26m	gs%				
No significant family	history present	Serum Cl	holestero	l 159mgs%			n Choles	terol	159mg	s%		Good	
		URINE -				URIN	E -						
		Albumin-		Nil		Albun			NII				
		Sugar -		Nil		Sugai			Nil				
		Deposits		NAD		Depos			NAD				
		CHEST >		Normal			ST X – R						
		SPUTIM					IM FOR	AFB					
_		MANTOL		Negative		MAN							
Days		1	3	6	9		12		15	18	21	24	27
Cough		Р	Р	R	R	-	R		R	R	R	R	R
Breathlessness		+++	+++		+-	-	++		++	+	+	+	+
Sputum	Colour	W	W	W	V		W		W	W	W	W	W
-	Quantity	20ml	20m		15		15ml		10ml	5ml	5ml	3ml	2ml
Rhonchi		Р	P	R	R	₹	R		R	R	R	R	R

1. Name:	Mr.Raman		Ag	e: 60	Sex:	M	0.1	P.No	o 385	86	No	. of	Days T	reated		35	Da	ys	F	rom	28.06.06
Drug	Sadamanjil ver	chooranam 1gm T	hree ti	mes a day with	Hot water			Di	iagnosis	5	Eraippu	Εrι	ımal			Occi	ıpati	on:	(Coolie	
Complaint	s & Duration	Respirat	ory Sy	stemExaminat	ion								Investi	gations	3						Doononoo
Complaint	S & Duration	Before Treatme	nt	After Treatm	ent				Before	Trea	atment					Aft	er Tr	eatme	nt		Response
				0		BLO	OD-	-						BLO	OD-						
		Cough	Р	Cough	R	TC -				9000	0Cells/cu.	mm		TC -				9000	Cells	cu.mn	n
						DC -	F	Р	52%	L	46%	Е	02%	DC	Р	52%	L	46%	E	029	%
		Coutum		Chutum		ESR		⅓ h	nr 5mr	n	1 hr	12	mm	ESR		⅓ hr	5m	m	1 hr	10m	m
		Sputum	W	Sputum	W	Hb-				69%				Hb-			69	%			
Cough with	expectoration					BI.St	ıgar	•		75m	gs%			BI.St	ıgar		70	mgs%			
difficulty in	breathing since	Breathlessness		Breathlessnes		BI. U	rea			23m	gs%			BI. U	rea		25	mgs%			Good
one month		Dieatillessiless	++	Dieatillessiles	+	Seru	m C	hole	esterol	15	53mgs%			Seru	m C	holester	ol	153	mgs%	ı	Good
						URIN	IE -							URIN	1E -						
						Albu		-		Nil				Albui	min-			Nil			
						Suga	ır -			Nil				Suga	ır -			Nil			
		Rhonchi	Р	Rhonchi	R	Depo	sits	; -		NAD)			Depo	osits	-		NAD			
		Knonchi		Knonchi		CHE	ST)	X – I	RAY		Normal			CHE	ST 2	X – RAY	,				
						SPU	TIM	FO	R AFB		Negative		•	SPU	TIM	FOR A	FB			•	
Ì						MAN	ΤΟΙ	UX			Negative			MAN	TOI	UX					

2. Name:	Krishnammal		Ag		Sex:	F	0.P.N	No :	39052	No). O	f Days T	reated		49	Da	ys		From	30.	06.06
Drug	Sadamanjil ver	chooranam 1gm T	Three tir	mes a day with	Hot water		[Diagno	sis	Eraippu	ı Er	rumal			Occu	pati	on:		House	wife	
Complaint	o P Duration	Respirat	ory Sy	stemExaminat	ion							Investig	gations	3							Doononoo
Complaint	s & Duration	Before Treatme	nt	After Treatm	ent			Befo	ore Tre	atment					Afte	er Tr	eatme	nt			Response
				Ob		BLO	OD-						BLO	OD-	•						
		Cough	Р	Cough	R	TC -			850	0Cells/cu	.mr	n	TC -				8000	Cell	s/cu.m	m	
						DC -	Р	58%	L	35%	Е	07%	DC	Р	58%	L	38%	E	0	4%	
		Consideration		Cmustum		ESR	1/2	hr	15mm	1 hr	3	2mm	ESR		⅓ hr	10r	nm	1 hr	20	mm	
		Sputum	W	Sputum	W	Hb-			71%)			Hb-			71	%				
Cough with	n expectoration,					Bl.Su	igar		70m	ıgs%			BI.Su	ıgar	•	80	mgs%				
	Breathlessness	Breathlessness		Breathlessnes		Bl. Ui	rea		19m	ıgs%			BI. U	rea		20	mgs%				Cood
since 10 da	ıys	Dieatillessiless	++	Dieatillessiles	s +	Seru	m Cho	lestero	ol 2	23mgs%			Seru	m C	Cholester	ol	230	mgs%	6		Good
						URIN	IE -						URIN	IE -							
						Albur	nin-		Nil				Albur	min-	-		Nil				
						Suga	r -		Nil				Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits -		NAE)			Depo	sits	; -		NAD				
		Knonchi	-	Knonchi	K	CHE	ST X -	- RAY		Chronic	Bro	nchtitis	CHE	ST 2	X – RAY		Nor	mal			
						SPU	TIM FO	OR AFI	В	Negative)		SPU	TIM	FOR AF	В			•		
						MAN	TOUX			Negative	,		MAN	TO	UX						

 $P-Present \quad R-Reduced \ W-Whitish \quad Y-Yellowish$

3. Name:	Mrs. Lakshmi		Age	e: 65	Sex:	F	0.P.I	No 3	9726		No.	of I	Days Tı	reated		32	Da	ys		From	04	.07.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water		I	Diagno	sis	Е	raippu	Eru	mal			Occu	patio	on:		Hous	ewife	
Complaint	o 9 Durotion	Respirat	ory Sys	stemExaminati	on							ı	nvestig	gations	3							Baananaa
Complaint	s & Duration	Before Treatme	nt	After Treatme	ent			Befo	re Tre	atm	ent					Afte	er Tr	eatme	nt			Response
				Count		BLO	OD-							BLO	OD:	-						
		Cough	Р	Cough	R	TC -			910	0C	ells/cu.r	nm		TC -				900	0Cel	ls/cu.	mm	1
						DC -	Р	70%	L	2	8%	Е	02%	DC	Р	68%	L	30%	E		02%	
		Constitution		Consideration		ESR	1/2	hr 1	2mm		1 hr	25r	mm	ESR		⅓ hr	10n	nm	1 hr	. 2	:0mm	1
01 .41		Sputum	W	Sputum	W	Hb-			68%	ó				Hb-			68	%]
	expectoration					BI.Su	gar		100	mgs	%			BI.St	ıgaı	ſ	95	/mgs%)			
Breathlessr		Breathlessness		Breathlessnes		Bl. Ui	rea		17m	ngs%	6			BI. U	rea		19	mgs%				Good
since 2 yea	o dust cold air	Dieatillessiless	++	Dieatillessiles	+	Seru	n Cho	lesterol	1	65m	ıgs%			Seru	m C	Cholester	ol	165	mgs	%		Good
Sirice 2 yea	113					URIN	E-							URIN	1E -							
						Albur			Nil					Albui	min	<u> </u>		Nil				
						Suga	r -		Nil					Suga	ır -			Nil				
		Rhonchi	P	Rhonchi	R	Depo	sits -		2-36	epith	elial cell	ls		Depo	osits	S -		NAD				
		KIIOIICIII	Г	KIIOIICIII	K	CHE	ST X -	- RAY		Ν	ormal			CHE	ST	X – RAY						
						SPU	ΓIM F	OR AFE	3	N	egative	ĺ	•	SPU	TIM	FOR AF	В			•	•	
						MAN	TOUX			N	egative	ĺ	•	MAN	ΤO	UX				•	•	

4. Name:	Tamilselvi		Age		Sex:	F	0.P.N	No	63716		No.	of [Days T	reated		29	Da	ys		Fron	m 07	'.11.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water			Diagno	osis	Era	aippu E	Erui	mal			Occi	ıpati	on:		Hou	sewife	
Complaint	o 9 Durotion	Respirat	ory Sys	stemExaminati	on							li	nvestig	ations	5							Baananaa
Complaint	s & Duration	Before Treatme	nt	After Treatme	ent			Bef	ore Tre	atme	nt					Aft	er Tr	eatme	ent			Response
				Count		BLO	OD-							BLO	OD-							
		Cough	Р	Cough	R	TC -			880	00Cells	s/cu.n	nm		TC -				850	0Ce	lls/cu	ı.mm	1
						DC -	Р	68%	L	23%	6	Е	09%	DC	Р	72%	L	26%	Е		02%	
		Constitution		Consideration		ESR	1/2	hr	10mm	1	hr	18r	nm	ESR		⅓ hr	5m	m	1 h	r	10mm	
		Sputum	W	Sputum	W	Hb-			719	6				Hb-			72	:%				
	expectoration					BI.St	ıgar		85n	ngs%				BI.St	ıgar		82	mgs%				
	breathing on	Breathlessness		Breathlessnes		BI. U	rea		26n	ngs%				BI. U	rea		25	mgs%				Cood
since 6 yea	o cold air, dust	Dieatillessiless	++	Dieatillessiles	+	Seru	m Cho	lester	ol 1	80mgs	s%			Seru	m Cł	nolester	ol	180)mgs	8%		Good
Since 6 yea	115					URIN	IE -							URIN	IE -							
						Albui	min-		Nil					Albui	min-			Nil				
						Suga	ar -		Nil					Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits -		NAI)				Depo	sits	-		NAD				
		Knonchi	P	Knonchi	K	CHE	ST X -	- RAY	,	Bror	nchitis			CHE	ST X	- RAY	,	No	rmal			
						SPU	TIM F	OR AF	B	Neg	ative			SPU	TIM	FOR A	-B					
						MAN	TOUX			Neg	ative			MAN	TOU	ΙX						

P-Present R-ReducedW-Whitish Y-Yellowish

5. Name:	Mrs. Alima		Ag	e: 65	Sex:	F	0.1	P.No	399	64	No	. of	Days T	reated		40	Da	ys		From	05.	07.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tii	mes a day with	Hot water			Di	iagnosis	;	Eraippu	Eru	mal			Occi	ıpati	on:		House	wife	
Complaint	s & Duration	Respirate	ory Sy	stemExaminat	tion								Investig	ations	3							Baananaa
Complaint	S & Duration	Before Treatmen	nt	After Treatm	ent				Before	Trea	atment					Aft	er Tr	eatm	ent			Response
				Carrell		BLO	OD-	•						BLO	OD-	•						
		Cough	Р	Cough	R	TC -				9000	0Cells/cu.	mm		TC -				850	00Cel	lls/cu.r	nm	
						DC -	F	Р	65%	L	30%	Е	05%	DC	Р	65%	L	32%	6 E		03%	
		Cautum		Chutum		ESR		⅓ h	r 16m	ım	1 hr	35	mm	ESR		⅓ hr	10r	mm	1 hr	- 20)mm	
		Sputum	W	Sputum	W	Hb-				68%				Hb-			68	3%				
Cough with	h expectoration,					BI.St	ıgar	•		88m	gs%			BI.St	ıgar		80)mgs%	, 0			
Breathlessr	ness since 1 ½	Breathlessness		Breathlessnes		BI. U	rea			23m	gs%			BI. U	rea		23	3mgs%	, 0			Good
months		Dieatillessiless	++	Dieatillessiles	+	Seru	m C	hole	esterol	14	48mgs%			Seru	m C	holester	ol	15	0mgs	%		Good
						URI	<u> 1E -</u>							URIN	1E -							
						Albu	min-	-		Nil				Albui	min-			Nil				
						Suga	ır -			Nil				Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits	; -		1NA	D			Depo	sits	-		NAD	1			
		KIIOIICIII	-	KIIOIICIII	K	CHE	ST)	X – I	RAY		Normal			CHE	ST 2	X – RAY	,					
						SPU	TIM	FOI	R AFB		Negative			SPU	TIM	FOR A	FB					
						MAN	ΤΟI	UX			Negative			MAN	TOI	UX						

6. Name:	Velayudham		Age	e: 58	Sex:	M	O.P.N	o 4	0157	No	o. of	Days T	reated		49	Da	ys	F	rom (6.07.06
Drug	Sadamanjil vei	chooranam 1gm T	Three tir	mes a day with	Hot water		D	iagnos	sis	Eraippi	ı Eru	mal			Occi	ıpati	on:	C	oolie	
Complaint	o ^o Durotion	Respirat	ory Sys	stemExamina	tion						I	Investiç	gations	•						Baananaa
Complaint	s & Duration	Before Treatme	nt	After Treatm	nent			Befor	re Trea	atment					Aft	er Tr	eatm	ent		Response
				0		BLOO	D-						BLO	OD-						
		Cough	Р	Cough	R	TC -			880	0Cells/cu	.mm		TC -				800	00Cells/	cu.mm	
						DC -	Р	63%	L	23%	Е	04%	DC	Р	65%	L	32%	6 E	03%	
		Cm.utuum		Constant		ESR	1/2 h	ır 3	0mm	1 hr	60	mm	ESR		½ hr	15r	nm	1 hr	30mm	ı
		Sputum	W	Sputum	W	Hb-			68%				Hb-			70)%			
	expectoration,					Bl.Sug	jar		83m	gs%			Bl.Su	ıgar		85	mgs%	6		
	breathing on	Breathlessness		Breathlessne		Bl. Ure	ea		33m	gs%			Bl. Ui	rea		32	mgs%	6		0
	cold air, dust	Dreamlessness	++	Dreatillessile	+	Serum	Chol	esterol	15	51mgs%			Seru	m Cl	holester	ol lo	15	2mgs%		Good
etc., Since	o monuis					URINE	<u> </u>						URIN	IE -						
						Album	in-		Nil				Albur	min-			Nil			
						Sugar	-		Nil				Suga	ır -			Nil			
		Rhonchi	Р	Dhanahi	R	Depos	its -		occu	It puscells	;		Depo	sits	_		NAD)		
		Knonchi		Rhonchi	K	CHES	T X –	RAY		chronic	bronc	hitis	CHE	ST X	(– RAY	•	No	ormal		
						SPUT	IM FO	R AFB	3	Negative	9		SPU	ТІМ	FOR A	FB				
						MANT	OUX			Negative	9		MAN	ΤΟι	JX					

P-Present R-ReducedW-Whitish Y-Yellowish

7. Name:	Gomathy		Age	e: 57	Sex:	F	0.P.N	No 4	1125		No. o	of Days T	reated		26	Da	ys		From	11.0	07.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water		I	Diagnos	sis	Erai	ippu E	rumal			Occu	pati	on:		Housev	vife	
Complaint	o 9 Durotion	Respirat	ory Sys	stemExaminat	ion							Investi	gations	3							Beenenee
Complaint	s & Duration	Before Treatme	nt	After Treatm	ent			Befor	re Tre	atmer	nt				Afte	er Tr	eatme	nt			Response
				0		BLO	OD-						BLO	OD	-						
		Cough	Р	Cough	R	TC -			840	0Cells	s/cu.m	m	TC -				800	Cell	s/cu.m	m	
						DC -	Р	60%	L	36%)	E 04%	DC	Р	60%	L	36%	E	()4	
		Consideration		Construe		ESR	1/2	hr 1	4mm	1	hr	30mm	ESR		⅓ hr	10r	nm	1 hr	20r	mm	
0		Sputum	W	Sputum	W	Hb-		•	66%)			Hb-			68	3%				
	expectoration					BI.Su	ıgar		166	mgs%			BI.St	ıgaı	r	13	0mgs%	, 0			
	breathing on cold air, dust	Breathlessness		Breathlessnes		BI. U	rea		29m	ıgs%			BI. U	rea		28	mgs%				Good
etc. Since 3		Dieatillessiless	++	Dieatillessiles	+	Serui	m Cho	lesterol	1	65mgs	%		Seru	m C	Cholester	ol	165	mgs%	Ď		Good
etc. Since c	5 1110111115					URIN	E-						URIN	IE -	•						
						Albur	nin-		Nil				Albu	min	-		Nil				
						Suga	r -		Nil				Suga	ar -			Nil				
		Rhonchi	P	Rhonchi	R	Depo	sits -		2-36	pithelia	al cells		Depo	osits	S -		NAD				
		Knonchi	Г	Knonchi		CHE	ST X -	- RAY		Norn	nal		CHE	ST	X – RAY						
						SPU	TIM F	OR AFB	}	Nega	ative		SPU	TIM	I FOR AF	В					
						MAN	TOUX			Nega	ative		MAN	ITO	UX						

8. Name:	Jafar		Ag		Sex:	M	O.P.N	No 4	41489	1	o. of	Days T	reated		31	Da	ys		From	13.07.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water			Diagno	sis	Eraip	ou Eru	ımal			Occi	ıpati	on:		coolie	
Complaint	o 9 Durotion	Respirat	ory Sys	stemExaminati	ion							Investi	gations	5						Boomenas
Complaint	s & Duration	Before Treatme	nt	After Treatme	ent			Befo	re Tre	atment					Aft	er Tr	eatme	nt		Response
				Oh		BLO	OD-						BLO	OD-						
		Cough	Р	Cough	R	TC -			970	0Cells/c	u.mm	1	TC -				9000	Cel	ls/cu.mr	n
						DC -	Р	70%	L	27%	Е	03%	DC	Р	65%	L	30%	E	05	%
		Cmartana		Constitute		ESR	1/2	hr 2	2mm	1 h	4 r	nm	ESR		⅓ hr	2m	m	1 hr	4mr	n
		Sputum	W	Sputum	W	Hb-			78%)			Hb-			78	3%			
	expectoration					BI.Su	ıgar		89m	ıgs%			BI.Su	ıgar		90	mgs%			
Breathlessr		Breathlessness		Breathlessnes		BI. U	rea		22m	ıgs%			BI. U	rea		25	mgs%			Cood
etc since 1	o cold air, dust	Diedillessiless	++	Dieatillessiles	+	Seru	m Cho	lestero	ol 1	51mgs%			Seru	m Cl	holester	ol	151	mgsʻ	%	Good
elc since i	HOHUI					URIN	IE -						URIN	IE -						
						Albui	min-		Nil				Albui	min-			Nil			
						Suga	ır -		Nil				Suga	ır -			Nil			
		Rhonchi	Р	Rhonchi	R	Depo	sits -		NAE)			Depo	sits	-		NAD			
		Knonchi		Knonchi	Γ.	CHE	ST X -	- RAY		Bronch	itis		CHE	ST X	(– RAY	7	Nor	mal		
						SPU	TIM FO	OR AFI	В	Negati	/e	•	SPU	TIM	FOR AF	FB			•	
						MAN	TOUX		•	Negati	/e	•	MAN	ΤΟΙ	JX				•	

P – Present R – Reduced W- Whitish Y - Yellowish

9. Name:	Mrs. Shankari		Ag	e: 45	Sex:	F	O.P.	No	4175	В	No	o. of	Days T	reated		47	Da	ys		From	14.	7.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day wit	h Hot wate	r		Diagr	nosis		Eraippu	ı Erı	umal			Occu	pati	on:		House	wife	
Complaint	s & Duration	Respirat	ory Sys	stemExamin	ation								Investig	gations	S							Bassanas
Complaint	s & Duration	Before Treatme	nt	After Treat	ment			Be	fore T	rea	tment					Afte	er Tr	eatme	nt			Response
				Carrela		BLO	OD-							BLO	OD	-						
		Cough	Р	Cough	R	TC -			90	000	Cells/cu	.mm	1	TC -				850	0Cel	s/cu.m	ım	
						DC -	Р	56%	% I	_	40%	Е	04%	DC	P	58%	L	40%	E		02	
		Considerate		Constitute		ESR	1/2	hr	13mm	1	1 hr	28	3mm	ESR		⅓ hr	10r	nm	1 hr	20	mm	
		Sputum	W	Sputum	W	Hb-			68	3%				Hb-			68	%		•		
Cough with	expectoration					BI.St	ugar		88	Bmg	s%			BI.St	uga	r	87	mgs%				
Breathlessr	ness since one	Breathlessness		Breathlessn	000	BI. U	rea		20)mg	s%			BI. U	rea		20	mgs%				Fair
month		Dieatillessiless	+++	Dieatillessii	++	Seru	m Ch	oleste	erol	18	5mgs%			Seru	m (Cholester	ol	180	mgs ^c	%		Ган
						URI	NE -							URIN	NE -							
						Albu	min-		N	l				Albu	min	-		Nil				
						Suga	ar -		N	l				Suga	ar -			Nil				
		Rhonchi	P	Rhonchi	R	Depo	osits -		F	ew p	ous cells /	/ HPI	F	Depo	osits	3 -		NAD				
		Knonchi	P	Knonchi		CHE	ST X	– RAY	Y		Normal			CHE	ST	X – RAY						
						SPU	TIM F	OR A	νFΒ		Negative)		SPU	TIN	FOR AF	В					
						MAN	ITOU	Κ			Negative	•		MAN	ITO	UX						

10. Name: Jebas		Ag		Sex:		O.P.N	lo 41	781	N	o. of	Days T	reated		48	Da	ys	F	rom	14.07.06
Drug Sadamanjil v	er chooranam 1gm 1	Three tii	mes a day wit	h Hot water			Diagnosi	is	Eraipp	u Eru	mal			Occu	ıpati	on:	C	Coolie	
Complaints & Duration	Respirat	tory Sy	stemExamin	ation						I	nvesti	gations	5						Beenenee
Complaints & Duration	Before Treatme	ent	After Treat	ment			Before	Tre	atment					Aft	er Tr	eatm	ent		Response
			0		BLOC	D-						BLO	OD-						
	Cough	Р	Cough	R	TC -			940	0Cells/cu	ı.mm		TC -				90	00Cells	/cu.mm	
					DC -	Р	64%	L	32%	Е	04%	DC	Р	64%	L	329	6 E	04%	, 6
	Constant		Consideration		ESR	1/2	hr 10	mm	1 hr	22	mm	ESR		½ hr	10r	nm	1 hr	20mr	n
0 1 111 1 11	Sputum	W	Sputum	W	Hb-			71%)			Hb-			71	%			
Cough with expectoration					Bl.Sug	gar		80m	gs%			BI.Su	ıgar		88	3mgs%	6		
chest tightness, breathless			Breathlessn	000	Bl. Ur	ea		17m	gs%			BI. U	rea		20	mgs%	6		Cood
during winter season since	breatiliessiless	++	Dieatillessii	+	Serun	n Chol	lesterol	1:	50mgs%			Seru	m C	holester	ol	15	0mgs%		Good
1 year					URINI	E-						URIN	IE -						
					Album	nin-		Nil				Albui	min-			Nil			
					Sugar	-		Nil				Suga	ır -			Nil			
	Rhonchi	Р	Rhonchi	R	Depos	sits -		NAC)			Depo	sits	-		NAD)		
	KIIOIICIII	-	KIIOIICIII	^	CHES	T X –	RAY		Bronchi	tis		CHE	ST >	(– RAY	,	N	ormal		
					SPUT	IM FC	R AFB		Negativ	е		SPU	TIM	FOR AF	FB				
					MANT	TOUX			Negativ	е		MAN	TOL	JX					

P-Present R-ReducedW-Whitish Y-Yellowish

11. Name:	Adhimoola	am	Age	e: 50	Sex:	М	0.P.I	No 4	42918		No	. of	Days T	reated		43	Da	ys		From	20.	07.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water			Diagno	sis		Eraippu	ιErι	umal			Occu	pati	on:		Farmer		
Compleint	a 9 Dunation	Respirat	ory Sys	stemExaminat	ion								Investi	ations	3							Deemanas
Complaint	s & Duration	Before Treatme	nt	After Treatm	ent			Befo	re Tre	eat	ment					Afte	er Tr	eatme	nt			Response
				0		BLO	OD-							BLO	OD	-						
		Cough	Р	Cough	R	TC -			970	000	Cells/cu.	.mm)	TC -				950	0Cel	ls/cu.m	m	
						DC -	Р	60%	L		36%	Е	04%	DC	P	62%	L	36%	E	0.	2%	
		Constant		Constitute		ESR	1/2	hr 5	5mm		1 hr	12	2mm	ESR		⅓ hr	5m	m	1 hr	12	mm	
0		Sputum	W	Sputum	W	Hb-			699	%				Hb-			70	1%				
	expectoration,					Bl.St	ıgar		98r	ng	s%			BI.St	ıga	r	90	mgs%				
Breathless		Breathlessness		Breathlessnes		BI. U	rea		18*	'nį	js%			BI. U	rea		20	mgs%				Good
'	o dust since 4	Dieatillessiless	++	Dieatillessiles	99 +	Seru	m Cho	lestero	ol ʻ	191	mgs%			Seru	m (Cholester	ol	190)mgs'	%		Good
years						URIN	IE -							URIN	۱E -							
						Albui	min-		Nil					Albui	min	-		Nil				
						Suga	ır -		Nil					Suga	ar -			Nil				
		Rhonchi	P	Rhonchi	R	Depo	sits -		Oc	cul	t pus cells	S		Depo	osits	s -		NAD				
		Knonchi		Knonchi	^	CHE	ST X -	- RAY			Normal			CHE	ST	X – RAY						
						SPU [*]	TIM F	OR AF	В		Negative			SPU	TIN	I FOR AF	В					
						MAN	TOUX	(Negative			MAN	ITO	UX						

12. Name:	Subbulakshi		Age		Sex:	F	O.P.N	No	45526		No.	of [Days T	reated		43	Da	ys		Fron	n 03	.08.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tir	nes a day with	Hot water			Diagno	osis	Er	aippu	Erui	mal			Occi	ıpati	on:		Hous	sewife	
Complaints	o O Duration	Respirat	ory Sys	stemExamina	tion							li	nvestig	gations	5							Beenenee
Complaints	s & Duration	Before Treatme	nt	After Treatm	nent			Befo	ore Tre	atme	ent					Aft	er Tr	eatme	ent			Response
				Carrela		BLO	OD-							BLO	OD-							
		Cough	Р	Cough	R	TC -			940	00Ce	lls/cu.r	nm		TC -				900	0Ce	lls/cu	.mm	
						DC -	Р	55%	L	43	3%	Е	02%	DC	Р	52%	L	46%	E		102%	
		Constant		Constant		ESR	1/2	hr	30mm		1 hr	62r	nm	ESR		½ hr	15r	nm	1 hr	r	30mm	
		Sputum	W	Sputum	W	Hb-			68%	, 0				Hb-			70	%				
Cough with	expectoration					BI.Su	igar		74n	ngs%				BI.St	ıgar		73	mgs%				
	oreathing since	Breathlessness		Breathlessne	00	BI. U	rea		25n	ngs%				BI. U	rea		22	mgs%				Cood
4 years	· ·	Dieatillessiless	++	Dieatillessile	+	Seru	m Cho	lestero	ol 1	87mg	gs%			Seru	m Cl	holester	ol	190	Omgs	%		Good
_						URIN	IE -							URIN	IE -							
						Albur	nin-		Nil					Albui	min-			Nil				
						Suga	r -		Nil					Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits -		occ	ult pu	s cells			Depo	sits	_		NAD				
		Knonchi	P	Knonchi	K	CHE	ST X -	- RAY		No	rmal			CHE	ST X	(– RAY	•					
						SPU	TIM FO	OR AF	В	Ne	gative			SPU	TIM	FOR A	FB					
						MAN	TOUX			Ne	gative			MAN	ΤΟΙ	JX						

P-Present R-ReducedW-Whitish Y-Yellowish

13. Name:	Annamalai		Ag	e: 70	Sex:	М	O.P.I	No	47089	9	No.	. of	f Days T	eated		43	Da	ys		Fro	n 11	.08.06
Drug	Sadamanjil ver	chooranam 1gm T	Three tir	mes a day with	Hot water			Diagn	osis		Eraippu	Er	umal			Occi	ıpati	on:		Lab	ourer	
Complaint	s & Duration	Respirat	ory Sy	stemExaminat	ion								Investig	ations	3							Bassanas
Complaint	S & Duration	Before Treatme	nt	After Treatm	ent			Bef	fore Tr	rea	tment					Aft	er Tr	eatme	ent			Response
				Carrel		BLO	OD-							BLO	OD.	-						
		Cough	Р	Cough	R	TC -			98	300	Cells/cu.i	mn	n	TC -				900	0Ce	lls/cu	ı.mm	1
						DC -	Р	66%	6 L	-	30%	Е	04%	DC	Р	60%	L	36%	, E		04%	
		Chutum		Coutum		ESR	1/2	hr	40mm	1	1 hr	85	5mm	ESR		½ hr	201	nm	1 h	r	40mm	
Б	D (I. I	Sputum	-	Sputum	-	Hb-			75	5%				Hb-			75	5%				
, ,	Breathlessness					BI.Su	ıgar		91	mg	s%			BI.St	ıgar	r	90)mgs%				
, chest tight		Breathlessness		Breathlessnes		BI. U	rea		19	mg	s%			BI. U	rea		20)mgs%				Good
•	dust since 3	Diedillessiless	++	Dieatillessiles	+	Seru	m Cho	olester	rol	15	3mgs%			Seru	m C	Cholester	ol	153	3mgs	%		Good
years						URIN	IE -							URIN	IE -	•						
						Albur	nin-		Ni	l				Albui	min	-		Nil				
						Suga	r -		Ni	l				Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits -		N/	٩D				Depo	sits	3 -		NAD				
		KIIOIICIII	-	KIIOIICIII	I.V.	CHE	ST X -	- RAY	1		Normal			CHE	ST	X – RAY	'					
						SPU	ΓΙΜ F	OR AF	FB		Negative			SPU	TIM	FOR A	-B					
						MAN	TOUX	(Negative		·	MAN	ΤO	UX						

14. Name:	Chandraseka	ar	Ag	e: 65	Sex:	M	O.P.	No	50687	,	No	. of	f Days T	reated		27	Da	ys		From	30	.08.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tii	nes a day with	Hot water			Diagn	nosis		Eraippu	Er	umal			Occu	ıpati	on:		Labo	ırer	
Complaint	o ^o Duration	Respirat	ory Sy	stemExamina	tion								Investig	ations	3							Baananaa
Complaints	s & Duration	Before Treatme	nt	After Treatn	nent			Be	fore Tr	eat	tment					Aft	er Tr	eatme	ent			Response
				Oh		BLO	OD-							BLO	OD-	•						
		Cough	Р	Cough	R	TC -			84	00	Cells/cu.	mn	n	TC -				850	0Cel	ls/cu.	mm	1
						DC -	Р	65%	6 L		28%	Е	07%	DC	Р	65%	L	30%	, E		05%	
		Curretures		Curretura		ESR	1/2	hr	15mm		1 hr	28	8mm	ESR		⅓ hr	15r	mm	1 hr	. 2	5mm	
0 1 11		Sputum	Υ	Sputum	W	Hb-			75	%				Hb-			75	5%				
	expectoration,					BI.Su	ıgar		21	1m	gs%			BI.St	ıgar	•	19	00mgs	%			
Breathlessn		Breathlessness		Breathlessne		BI. U	rea		22	mg:	s%			BI. U	rea		20)mgs%)			Fair
since 1 yea	cold air dust et	Dieatillessiless	+++	Dieatillessile	++	Seru	m Cho	oleste	rol	198	3mgs%			Seru	m C	Cholester	ol	198	8mgs	%		Fall
Since i yea						URIN	IE -							URIN	IE -							
						Albui	min-		Nil					Albui	min-	-		Nil				
						Suga	ır -		+					Suga	ar -			+				
		Rhonchi	Р	Rhonchi	Ь	Depo	sits -		N/	۱D				Depo	osits	; -		NAD				
		Knonchi		Knonchi		CHE	ST X -	– RAY	<u> </u>		Normal			CHE	ST	X – RAY						
						SPU	TIM F	OR A	FB		Negative			SPU	TIM	FOR AF	В]
						MAN	TOU	(Negative			MAN	ITO	UX]

P-Present R-Reduced W-Whitish Y-Yellowish

15. Name:	Mariappan		Ag	e: 60	Sex:	M	O.P.	No	6162	8	No	. of	Days T	reated		27	Da	ıys		From	26.	10.06
Drug	Sadamanjil ver	chooranam 1gm T	Three ti	mes a day with	Hot water			Diag	nosis		Eraippu	Εrι	umal			Occu	ıpati	on:		Farmer		
Complaint	s & Duration	Respirat	ory Sy	stemExaminat	ion								Investig	ations	3							Beenenee
Complaints	s & Duration	Before Treatme	nt	After Treatm	ent			Be	efore T	rea	atment					Aft	er Tr	eatm	ent			Response
				Count		BLO	OD-							BLO	OD-	-						
		Cough	Р	Cough	R	TC -			86	30C	Cells/cu.	mm	1	TC -				800	00Cel	ls/cu.m	m	
						DC -	Р	56%	% I	_	40%	Е	04%	DC	Р	56%	L	40%	6 E	04	4%	
		Chutum		Coutum		ESR	1/2	hr	20mm	1	1 hr	42	2mm	ESR		⅓ hr	10r	mm	1 hr	201	mm	
0 1 31		Sputum	W	Sputum	W	Hb-			71	1%				Hb-			71	l%				
	expectoration					BI.Su	gar		75	5%				BI.St	ıgar	ſ	72	2%				
Breathlessn		Breathlessness		Breathlessnes		BI. U	rea		19	9%				BI. U	rea		19	9%				Good
etc since 1	cold air, dust	Dieatillessiless	++	Dieatillessiles	+	Serui	n Cho	oleste	erol	15	3%			Seru	m C	Cholester	ol	15	3%			Good
elc since i	yeai					URIN	E-							URIN	IE -	•						
						Albur			Ni	I				Albui	min-	-		Nil				
						Suga	r -		Ni	I				Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	R	Depo	sits -		1-	2рι	uscells			Depo	sits	S -		NAD				
		KIIOIICIII	-	Kilolicili	K	CHE	ST X	– RA	Υ.		Normal			CHE	ST	X – RAY	'					
						SPU	ΓIM F	OR A	\FB		Negative		•	SPU	TIM	FOR AF	FB					
						MAN	TOU	(Negative		·	MAN	TO	UX						

16. Name:	Balasubram	an ian	Age	e: 62	Sex:	M	O.P.	No (61737		No.	of D	ays Ti	reated		28	Da	ys		From	26	.10.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tin	nes a day wit	h Hot water			Diagno	osis	Era	ippu I	Erur	nal			Occu	pati	on:		House	wife	
Complaint	o 9 Duration	Respirate	ory Sys	temExamina	ation							lr	nvestig	ations	5							Beenenee
Complaint	s & Duration	Before Treatmen	nt	After Treati	ment			Befo	ore Tre	atmer	nt					Afte	er Tr	eatme	nt			Response
				0		BLO	OD-							BLO	OD-							
		Cough	Р	Cough	R	TC -			860	0Cells	s/cu.n	nm		TC -				800	0Cel	ls/cu.r	nm	
						DC -	Р	64%	L	32%	0	Е	04%	DC	Р	60%	L	36%	E		04%	
		Current		Carretrona		ESR	1/2	hr	10mm	1	hr	30n	nm	ESR		⅓ hr	10r	nm	1 hr	2	Omm	
		Sputum	W	Sputum	W	Hb-			71%	, 0				Hb-			71	%				
Cough with	expectoration					BI.Su	ıgar		70n	ngs%				BI.Su	ıgar		75	mgs%				
	breathing on cold air, dust	Breathlessness		Breathlessne	nee.	BI. U	rea		23n	ngs%				BI. U	rea		20	mgs%				Good
etc. Since 2		Dieatillessiless	++	Dieatillessile	+	Seru	m Cho	olestero	ol 1	54mgs	s%			Seru	m C	holester	ol	154	lmgs'	%		Good
etc. Since 2	z years					URIN	IE -							URIN	IE -							
						Albui	min-		Nil					Albur	min-			Nil				
						Suga	ır -		Nil					Suga	ır -			Nil				
		Rhonchi	Р	Rhonchi	В	Depo	sits -		1-2	ouscells	S			Depo	sits	-		NAD				
		Knonchi	Ρ	Knonchi	R	CHE	ST X ·	- RAY		Norn	mal			CHE	ST)	K – RAY						
						SPU	TIM F	OR AF	В	Nega	ative			SPU	TIM	FOR AF	В					
						MAN	TOU	(Nega	ative			MAN	ΤΟΙ	JX						

 $P-Present \quad R-Reduced \ W-Whitish \quad Y-Yellowish$

17. Name:	Sornam		Ag	e: 55	Sex:	F	0.P.N	No	61854	N	0. 0	f Days T	reated		27	Da	ys	F	rom	29.10.06
Drug	Sadamanjil ver	chooranam 1gm T	Three tii	mes a day with	Hot water		1	Diagno	osis	Eraipp	u Er	rumal			Occu	ıpati	on:	ŀ	Housew	rife
Complaint	s & Duration	Respirat	ory Sy	stemExaminati	on							Investig	ations	5						Baanan
Complaint	S & Duration	Before Treatme	nt	After Treatme	ent			Befo	ore Tre	atment					Aft	er Tr	eatme	nt		Respons
				Carrel		BLO	OD-						BLO	OD-						
		Cough	Р	Cough	R	TC -			112	00Cells/d	cu.m	nm	TC -				9000	Cells	s/cu.mi	n
						DC -	Р	68%	L	20%	Е	12%	DC	Р	68%	L	28%	E	04	%
		Cnutum		Coutum		ESR	1/2	hr	3mm	1 hr	6	imm	ESR		½ hr	3m	m	1 hr	6m	m
0 1 31		Sputum	W	Sputum	W	Hb-			71%	,)			Hb-			70	%			
	expectoration					Bl.Su	gar		94n	ngs%			BI.Su	ıgar	•	94	mgs%			
	reathlessness	Breathlessness		Breathlessnes	_	Bl. Ur	ea		27m	ngs%			BI. U	rea		25	mgs%			Cood
	re to dust since	Dreatillessiless	+++	Dreatillessiles	S +	Serur	n Cho	lestero	ol 1	94mgs%			Seru	m C	holester	ol	190	mgs%	1	Good
1 year						URIN	E -						URIN	IE -						
						Albur	nin-		Nil				Albur	min-	-		Nil			
						Suga	r -		Nil				Suga	ır -			Nil			
		Dhanahi	Р	Dhanah!	D	Depo	sits -		2-3	ouscells			Depo	sits	; -		NAD			
		Rhonchi		Rhonchi	R	CHES	ST X -	- RAY		Bronchi	tis		CHE	ST 2	X – RAY	·	Nor	mal		
						SPUT	TIM FO	OR AF	В	Negativ	е		SPU	TIM	FOR AF	FB				
						MAN	TOUX			Negativ	е		MAN	TOI	UX					

18. Name:	Annamalai		Age	e: 60	Sex:		M	0.P.N	lo 6	2185	1	lo. c	of Days T	reated		25	Da	ys		From	30	.10.06
Drug	Sadamanjil ver	chooranam 1gm T	hree tin	nes a day	with Hot wa	ter			Diagnos	is	Eraip	ou E	rumal			Occu	pati	on:		House	ewife	
Complaint	o 9 Durotion	Respirat	ory Sys	stemExam	ination								Investi	gations	3							Baananaa
Complaint	s & Duration	Before Treatme	nt	After Tre	atment				Befor	e Tre	atment					Afte	er Tr	eatme	nt			Response
				0			BLOC	D-						BLO	OD-	•						
		Cough	Р	Cough	R		TC -			700	0Cells/d	u.m	m	TC -				750	0Cel	ls/cu.r	nm	1
							DC -	Р	64%	L	30%	ı	E 06%	DC	Р	64%	L	30%	E		06%	
		C		0			ESR	1/2	hr 15	5mm	1 h	r 2	28mm	ESR		⅓ hr	15r	nm	1 hr	. 2	8mm	1
		Sputum	Υ	Sputum	W	1	Hb-			81%)			Hb-			81	%]
Cough with	expectoration						Bl.Sug	gar		91m	ıgs%			BI.St	ıgar	•	98	mgs%				
Breathlessr	ness, chest	Breathlessness		Breathles	enose		Bl. Ure	ea		32m	ıgs%			BI. U	rea		32	mgs%				Good
tightness si	nce 2 months	Dieatillessiless	++	Dicallics	+		Serum	n Cho	lesterol	1:	56mgs%			Seru	m C	Cholester	ol	15	mgs	%		Good
							URINE	E -						URIN	IE -							
							Album	in-		Nil				Albui	min-	-		Nil				
							Sugar	-		Nil				Suga	ar -			Nil				
		Rhonchi	Р	Rhonchi			Depos	sits -		NAD)			Depo	osits	; -		NAD				
		Knonchi	Р	Knonchi	R		CHES	T X -	- RAY		Chroni	c bro	nchitis	CHE	ST	X – RAY		No	rmal			
							SPUT	IM FO	OR AFB		Negati	ve		SPU	TIM	FOR AF	B					
							MANT	OUX			Negati	ve		MAN	ITO	UX						

P-Present R-Reduced W-Whitish Y-Yellowish

19. Name:	Jeyalaxmi		Ag	e: 30	Sex:	F	O.P.N	lo 6	2329	N	o. of	Days T	reated		25	Da	ys	F	rom	30.1	10.06
Drug	Sadamanjil ver	chooranam 1gm T	Three tir	nes a day with	Hot water		D	Diagnos	sis	Eraipp					Occu	patio	on:	ŀ	Housev	vife	
Commisint	a 9 Duration	Respirat	ory Sy	stemExamina	tion							Investiç	gations	3							Deemanaa
Complaint	s & Duration	Before Treatme	nt	After Treatm	ent			Befor	re Trea	atment					Afte	r Tr	eatme	nt			Response
				6		BLO	OD-						BLO	OD)-						
		Cough	Р	Cough	R	TC -			800	0Cells/cu	ı.mm		TC -				8000	Cells	s/cu.m	m	
						DC -	Р	64%	L	32%	Е	04%	DC	P	6 4%	L	32%	Е	04	4%	
		Constitute		Constitute		ESR	1/2	hr 7	mm	1 hr	15	mm	ESR		½ hr	5mi	m	1 hr	101	mm	
		Sputum	W	Sputum	W	Hb-		•	68%		•		Hb-			68	%		•		
	expectoration					Bl.Su	ıgar		92m	gs%			BI.Su	ıga	r	90	mgs%				
	oreahlessnesss	Breathlessness		Breathlessne		BI. Ui	rea		20m	gs%			BI. U	rea		22	mgs%				Fair
dust since	e to cold air,	Dieatillessiless	++	Dieatillessiles	+	Serur	m Chol	lesterol	15	59mgs%			Seru	m (Cholester	ol	160	mgs%)		raii
dust since .	3 1110111115					URIN	IE -						URIN	IE -	-						
						Albur	nin-		Nil				Albur	min	 -		Nil				
						Suga	r -		Nil				Suga	ar -			Nil				
		Rhonchi	P	Rhonchi		Depo	sits -		Few	epithelial	cells		Depo	osite	s -		NAD				
		Knonchi		Knonchi	R	CHE	ST X -	RAY		Normal			CHE	ST	X – RAY						
						SPU	TIM FC	R AFE	3	Negativ	Э		SPU	TIN	I FOR AF	В					
						MAN	TOUX			Negativ	Э		MAN	ITO	UX						

20. Name:	Shankarasuk	bu	Age	e: 63	Sex:	M	O.P.N	lo 6	52937	No	. of	Days T	reated		34	Da	ys	F	rom	02.11.06
Drug	Sadamanjil ver	chooranam 1gm T	Three tin	nes a day with	Hot water		D	Diagno	sis	Eraippu	ιErι	umal			Occu	pati	on:	la	abourer	
Complaint	o 9 Duration	Respirat	ory Sys	temExamina	tion							Investig	gations	3						Beenenee
Complaint	s & Duration	Before Treatme	nt	After Treatn	nent			Befo	re Tre	atment					Afte	er Tr	eatme	nt		Response
				0		BLO	OD-						BLO	OD-	•					
		Cough	Р	Cough	R	TC -			105	00Cells/ci	u.mı	m	TC -				9000	Cells	/cu.mr	n
						DC -	Р	50%	L	35%	Е	15%	DC	Р	58%	L	36%	E	06	%
		C		C		ESR	1/2	hr 4	1mm	1 hr	10)mm	ESR		⅓ hr	4m	m	1 hr	10m	ım
		Sputum	W	Sputum	W	Hb-			70%	,			Hb-			70	%			
	expectoration					BI.St	ıgar		91m	gs%			BI.Su	ıgar		92	mgs%			
	ness breathness	Breathlessness		Breathlessne		BI. U	rea		17m	gs%			BI. U	rea		18	mgs%			Good
	e to dust, since	Dieatillessiless	++	Dreatillessile	+	Seru	m Chol	lestero	1 10	64mgs%			Serui	m C	holester	ol	165	mgs%		Good
young age						URIN	IE -						URIN	1E -						
						Albui	min-		Nil				Albur	min-			Nil			
						Suga	ır -		Nil				Suga	ır -			Nil			
		Dhanah!	Р	Dhanahi		Depo	sits -		NAC)			Depo	sits	-		NAD			
		Rhonchi		Rhonchi	R	CHE	ST X -	RAY		Normal			CHE	ST 2	X – RAY					
						SPU	TIM FC	R AFE	3	Negative	1		SPU	TIM	FOR AF	В				
						MAN	TOUX			Negative	!		MAN	TOI	UX					

P-Present R-Reduced W-Whitish Y-Yellowish

Mild -+ Moderate - ++ Severe - +++
P - Present R - Reduced
W- Whitish Y - Yellowish