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AN UPDATE ON AYURVEDIC HERB HENNA (*LAWSONIA INERMIS* L.): A REVIEW

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ABSTRACT: *Lawsonia inermis* L. is a much branched glabrous shrub or small tree, cultured for its leaves although stem bark, roots, flowers and seeds have also been used in traditional medicine. It has been traditionally reported in use of headache, hemicranias, lumbago, bronchitis, boils, ophthalmia, syphilitis, sores, amenorrhea, scabies, diseases of the spleen, dysuria, bleeding disorder, skin diseases, diuretic, antibacterial, antifungal, anti-amoebiasis, astringent, anti-hemorrhagic, hypotensive and sedative effect. The plant is reported to contain Lawsone, Esculetin, Fraxetin, Isoplumbagin, Scopoletin, Betulin, Betulinic acid, Hennadiol, Lupeol, Lacoumarin, Laxanthone, Flavone glycosides, two pentacyclic triterpenes. The plant is reported to contain carbohydrates, proteins, flavonoids, tannins and phenolic compounds, alkaloids, terpenoids, quinones, coumarins, xanthenes and fatty acids. The plant has been reported to have analgesic, hypoglycemic, hepatoprotective, immunostimulant, anti-inflammatory, antibacterial, antimicrobial, antifungal, antiviral, antiparasitic, antitrypanosomal, antidermatophytic, antioxidant, antifertility, tuberculostatic and anticancer properties. It is now measured as a valuable source of exclusive natural products for growth of medicines against various diseases and also for the development of industrial products. This review gives a bird's eye vision mainly on the pharmacognostic characteristics, traditional uses, phytochemistry and pharmacological actions of the plant.

INTRODUCTION: Medicinal plants are part and parcel of human society to combat diseases, from the dawn of civilization¹. There exists a plethora of knowledge, information and benefits of herbal drugs in our ancient literature of Ayurvedic (Traditional Indian Medicine), Siddha, Unani and Chinese medicine.

According to the World Health Organization, 2003 about 80 % of the population of developing countries being unable to afford pharmaceutical drugs rely on traditional medicines, mainly plant based, to sustain their primary health care needs². Herbal medicines are in great demand in the developed as well as developing countries for primary healthcare because of their wide biological and medicinal activities, higher safety margins and lesser costs³.

The traditional medicinal methods, especially the use of medicinal plants, still play a vital role to cover the basic health needs in the developing

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countries. In recent years there has been a phenomenal rise in the interest of scientific community to explore the pharmacological actions of herbs or to confirm the claims made about them in the official books of Ayurveda⁴. *Lawsonia inermis* Linn (Family: Lythraceae) is a much branched glabrous shrub or small tree (2-6 m in height), cultivated for its leaves although stem bark, roots, flowers and seeds have also been used in traditional medicine. This plant is a worldwide known cosmetic agent used to stain hair, skin and nails⁵.

The present attempt is to review and compile updated information on various aspects of *L. inermis* Linn. a plant used all over the world. This plant is commonly known as Henna and abundantly available in tropical and subtropical areas. Ancient history of India describes its diverse uses and also plays appreciable role in Ayurvedic or natural herbal medicines⁶.

Plant profile:



FIG. 1: *LAWSONIA INERMIS* LINN.

Synonym: *Lawsonia alba* Lam.

Vernacular names⁷:

English: Henna, Samphire, Cypress shrub.

Sanskrit: Mendhi, Mendika, Timir.

Arabic: Alhenna, Hinna.

French: Alcana d'orient.

Greek: Kypros.

Gujrat: Medi.

Hindi: Hena, Mhindi.

Marthi: Mendhi, Mendi.

Tamil: Alvanam, Aivani.

Telugu: Goranta, Kormmi.

Scientific classification⁸:

Kingdom: Plantae

Subkingdom: Viridiaeplantae

Division: Tracheophyta

Subdivision: Spermatophytina

Class: Magnoliopsida

Order: Myrtales

Family: Lythraceae

Genus: *Lawsonia*

Species: *inermis*

Description: It is much branched, deciduous, glabrous, sometime spinescent shrub or small tree with grayish brown bark, attaining a height of 2.4-5 m. It is cultivated as a hedge plant throughout India, and as a commercial crop in certain states of India for its dye. Leaves are 1.3-3.2 by 0.6-1.6 cm, elliptic or broadly lanceolate, acute or obtuse, often mucronulate, base tapering; petioles very short. Flowers are numerous, less than 1.3 cm. across fragrant, white or rose-colored, in large terminal pyramidal paniced cymes; pedicels short, slender. Calyx 3-5 mm, long broadly campanulate; lobes 2.5-3 mm, long, suborbicular or subreniform, undulate. Stamens 8, inserted in pairs on the calyx-tube. Capsules 6 mm, diameter; hlobose, slightly veined outside, supported by the persistent calyx and tipped with the style. Seed capsules are red, globose, about the size of a pea, with numerous tiny pyramidal, brown pitted seeds⁹.

Habitat: Henna, a traditional product with religious associations, has been widely used over the centuries for medical and cosmetic purposes in

Africa, Asia, the Middle East and many other parts of the world. Henna is a finely ground brown or green powder originating from dried leaves of the plant *Lawsonia inermis* which is grown in dry tropical and subtropical zones, including North Africa, India, Sri Lanka, and the Middle East¹⁰.

Propagation: by seeds¹¹.

Chemical constituents: Table 1¹².

TABLE 1: CHEMICAL CONSTITUENTS

S no.	Plant Parts	Chemical constituents
1.	Leaves	2-Hydroxy-1,4-napthoquinone, 1,4-dihydroxynaphthalene, 1,4-napthoquinone, 1,2-dihydroxy-glucyloxynaphthalene, luteolins, apigenin, and their glycosides, esculetin, fraxetin, scopletin, β -sitosterol, tannin, gallic acid, glucose, mannitol, fat, resin and mucilage.
2.	Barks	napthoquinone, isoplumbagin, triterpenoids-Hennadiol, aliphatics (3-methylnonacosan-1-ol)
3.	Flowers	essential oil (0.02 %) rich in ionones (90 %), β -ionones.
4.	Roots	24 β -ethylcholest-4-en-3 β -ol
5.	Seeds	Linoleic acid, Arachidic acid, Stearic acid, Palmitic acid
6.	Whole plant	Laxanthone I, Laxanthone II, Laxanthone III, n-Triacontanol,

Traditional uses^{13, 14, 15}:

1. It is used for the treatment of epilepsy and jaundice, and for dyeing grey hair.
2. It is used as a remedy for malignant ulcers.
3. The Ayurvedic Pharmacopoeia of India indicated the use of leaves in dysuria, bleeding disorder, prurigo and other obstinate skin diseases.
4. The leaf is used in vulnerary, diuretic, headache, hemicranias, lumbago, bronchitis, boils, ophthalmia, syphilitis, sores, amenorrhoea, scabies, and spleen diseases and favours the growth of the hair.
5. The bark is given in jaundice and enlargement of the spleen, also in calculous affections and as an alternative in leprosy and obstinate skin diseases.
6. It is used as medicinal plant because of its attributed antibacterial, antifungal, antiamoebiasis, astringent, antihemorrhagic, hypotensive and sedative effect.

Medicinal importance¹⁶:

1. It is used for antidiarrheal.
2. It is used for antidyenteric.
3. It is used for astringent.
4. It is used for emmenagogue.
5. It is used for liver tonic.

6. It is used for antifungal.

Ethnobotanical Uses¹⁷:

1. Henna leaf has an orange-red dye and leaf paste or powder is widely used for decorating hands, nails and feet with patterns.
2. **Flowers** are very fragrant and used to extract a perfume, which is used as base for local scents. 3. An infusion of the flowers is a valuable application to bruises. Decoction of the flowers is describes as an emmenagogue.
3. **Seeds** are deodorant. Powered seeds with real ghee (clarified butter) are effective against dysentery.
4. The bark is applied in the form of a decoction to burns and scalds. It is given internally in a variety of affections, such as jaundice, enlargement of the spleen, calculus, as an alternative in leprosy and obstinate skin affections.
5. **Root** is considered as a potent medicine for gonorrhoea and herpes infection. Root is astringent may be pulped and used for sore eyes. Pulped root may also be applied to the heads of children for boils.
6. The root is supposed to be useful in treatment of hysteria and nervous disorders.

TABLE 1: PRELIMINARY PHARMACOLOGICAL ACTIVITIES OF ALCOHOLIC EXTRACTS OF *L. INERMIS* L.

S No.	Activity	Plant part/ Extract	Dose/ Model	Standard drug	Result
1.	Antiviral Activity	Fruits / Ethanol	swiss mice and chick embryo models	Sembiki forest virus	It exhibiting 100 to 65 % activities after 10 to 25 days of virus challenge ¹⁸
2.a.	Wound Healing Activity	Leaves / Ethanol	200 mg/kg/day / excision, incision and dead space wound models.	Topical application, Oral treatment	The extract-treated animals showed 71% reduction in the wound area when compared with controls which was 58% ¹⁹
2.b.	Wound Healing Activity	Leaves / Chloroform	200 mg/kg/day / excision, incision and dead space wound models.	Topical application, Oral treatment	The effects of chloroform extracts against the primary invaders of burnt wounds was investigated ²⁰
3.	Protein Glycation Inhibitory Activity	Leaves / Ethanol	1500µg/mL, 1000µg/mL and 1000µM / the model system of bovine serum albumin and glucose.	AGE fluorescence intensity	The alcoholic extract, showed significant inhibition of Advanced Glycated End Products (AGEs) formation ^[21] .
4.a	Anti Diabetic Activity	Leaves / Ethanol	800 mg/kg / alloxan induced model	Glimepride	The extract decreased the glucose concentration after the 14th day and decreased total cholesterol and triglyceride concentration ²² .
4.b	Anti Diabetic Activity	Leaves / Methanol	800 mg/kg / alloxan induced model	Glimepride	It show the inhibitory effect of glucose utilization, and use as hypoglycemic agents ²³ .
5.	Nootropics activity	Leaves / Acetone	Elevated plus maze and passive shock avoidance paradigms.	Haloperidol	The leaves of <i>Lawsonia inermis</i> possess a potential for exploring a nootropic principle ²⁴ .
6.a	Antimicrobial Activity	Leaves / Methanol	Staphylococcus aureus Staphylococcus epidermidis	Tetracycline, Ampicillin	The presence of anthraquinones in the plant leaves are commonly known to possess antimicrobial activity ²⁵ .
6.b	Anti-Microbial Activity	Leaves / Alcoholic	Staphylococcus aureus Staphylococcus epidermidis	Tetracycline, Ampicillin	Alcoholic extracts had the highest antibacterial activity against β-hemolytic streptococci ²⁶
6.c	Anti-Microbial Activity	Leaves, Seeds / Ethanol	Staphylococcus aureus, Escherichia coli and Pseudomonas aeruginosa.	Tetracycline, Ampicillin	Omani henna does possess in-vitro antibacterial activity against a wide spectrum of bacterial strains and <i>C. albicans</i> ²⁷ .
7.a	Antibacteria-l Activity	Leaves / Ethyl acetate	E.coli ATCC 8739, S.aureus 6538 / Cup plate Model	Tetracycline	Ethyl acetate extract of <i>Lawsonia inermis</i> was found to be the most active one against all bacteria in the test system ²⁸ .
7.b	Antibacteria-l Activity	Leaves, seed / Ethanol	<i>Pseudomonas aeruginosa</i> (NCTC 10662)	Tetracycline	The highest susceptibility was against <i>P. aeruginosa</i> with henna samples obtained from Al-sharqyia region ²⁹ .
8.	Trypsin Inhibitory Activity	Leaves / Ethanol		Aralast	<i>Lawsonia inermis</i> alcoholic extract and lawsone have shown a significant Trypsin inhibitory effect ³⁰ .
9.a.	Cytotoxic Activity	Leaves / Chloroform	microculture tetrazolium salt assay	<i>Escherichia coli</i>	It displayed the cytotoxic effects against liver (HepG2) and Human breast (MCF-7) with IC50 values of 0.3 and 24.85 µg/ml ³¹ .
9.b.	Cytotoxic	Leaves /	Ames mutagenicity	<i>Escherichia coli</i>	Lawsone exposure inhibited the growth of

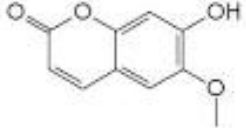
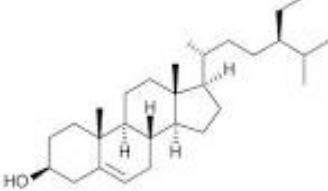
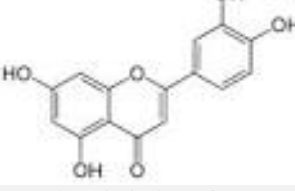
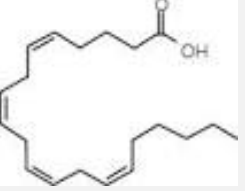
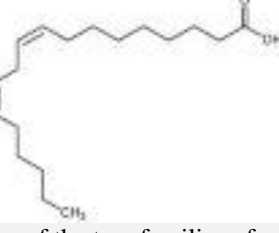
	Activity	Ethanol	assay		both Csa and Csb strains in a dose-dependent manner ³² .
10.a.	Antioxidant Activity	Leaves / Ethanol	200 and 400 mg/kg / 1, 1-diphenyl-2-picryl- hydrazyl model.	Ascorbic acid	The doses were effective in increasing the hepatic glutathione reductase, superoxide dismutase and catalase activities ³³ .
10.b.	Antioxidant Activity	Leaves / Methanol	200 and 400 mg/kg / 1, 1-diphenyl-2-picryl- hydrazyl model.	Ascorbic acid	In effect of different concentrations of methanolic extract of henna in comparison with synthetic antioxidant ³⁴ .
10.c	Antioxidant Activity	Leaves / Methanol	200 and 400 mg/kg / 1, 1-diphenyl-2-picryl- hydrazyl model.	Ascorbic acid	It was shown extraction method has significant effect on phenolic compound and antioxidant activity of Henna extract ³⁵ .
10.d	Antioxidant Activity	Leaves / Methanol	free radical scavenging assay	Ascorbic acid	It resulted in the isolation of seven compounds; three have been isolated for the first time from the genus, namely <i>p</i> -coumaric acid ³⁶ .
11.	Anticorrosi-n Activity	Leaves / Ethanol	SEM/EDS		Maximum inhibition efficiency (92.06 %) is obtained at 1.2 g/l henna extract ³⁷ .
12.	Anti-Inflammatory, Analgesic And Antipyretic Activity	Leaves / Chloroform	500 mg/kg	Ibuprofen	The isolated compound was found to possess significant anti-inflammatory, analgesic, and antipyretic activity ³⁸ .
13.	Tuberculost-atic Activity	Leaves / Ethanol	5 mg/kg	<i>Mycobacterium tuberculosis</i>	The growth of <i>Tubercle bacilli</i> from sputum and of <i>Mycobacterium tuberculosis</i> H37Rv was inhibited by 6 µg/ml of the herb ³⁹ .
14.a	Hepatoprote-ctive Activity	Bark / Alcohol	carbon tetrachloride induced model	Silymarine	Pretreatment of rats with the extract also inhibited the peroxidation of microsomal lipids in a dose-dependent manner ⁴⁰ .
14.b	Hepatoprote-ctive Activity	Leaves / Ethanol	CCl4-induced liver toxicity.	Silymarine	The effects of the extract on hexobarbitone-induced sleep, BSP clearance, and on certain biochemical parameters indicated its protective role ⁴¹ .
15.	Immunomo-dulatory Activity	Leaves / Methanol	mice lethality test, indirect hemagglutination test.	Levamisole	The immuomodulatory profile was studied using an <i>in vitro</i> immunoassay, the lymphocyte transformation assay ⁴² .
16.	Anticarcino-genic Activity	Leaves / chloroform	microculture tetrazolium salt (MTT) assay	Sulforaphane	The extract displayed the cytotoxic effects against HepG2 and MCF-7 with IC ₅₀ -value of 0.3 and 24.85 µg ml ⁻¹ ⁴³ .
17.	Molluscicid-al Activity	Seed / ethanol	Biomphalaria alexandrina snails	indomethacin	Highest toxicity was observed in the seed of <i>Lawsonia inermis</i> ⁴⁴ .
18.	Antifungal Activity	Bark / Ethanol	Microsporium gypseum and Trichophyton mentagrophytes	Voriconazole	The extract was found to possess fungistatic nature at its maximum inhibitory dilution of 1:30 (W/V) against both the test pathogens ⁴⁵ .
19.	Antitrypano-somal Activity	Leaf/ Methanolic extract	<i>invitro</i> activity against <i>Trypanosoma brucei</i>	<i>Trypanosoma brucei</i> / 8.3 mg/ml of blood in mice	The treatment tends to ameliorate the disease condition, but did not affect the level of parasitaemia and pack cell volume ⁴⁶ .
20.	Antisickling Activity	Leaves/ Aqueous extract	sickle cells counting	pentoxifylline	It found to inhibit sickling and to increase the oxygen affinity of HbSS blood ⁴⁷ .
21.	Abortifacie-nt	Roots /	Ovariectomized		The methanol extract effectiveness as an

Activity	Methanol extract	rats.	abortant due to its maternal and foetal toxic effects ⁴⁸
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Chemical Review ⁴⁹: The principal colouring matter of henna is lawsone, 2-hydroxy-1:4 naphthaquinone (C₁₀H₆O₃, m.p.190° decomp.) besides lawsone other constituents present are gallic acid, glucose, mannitol, fats, resin (2 %), mucilage and traces of an alkaloid. Leaves yield hennatannic acid and an olive oil green resin, soluble in ether and alcohol. Flowers yield an essential oil (0.01-0.02 %) with brown or dark brown colour, strong fragrance and consist mainly

of α - and β - ionones; a nitrogenous compound and resin. Seeds contain proteins (5.0 %), carbohydrates (33.62 %), fibers (33.5 %), fatty oils (10- 11 %) composed of behenic acid, arachidic acid, stearic acid, palmitic acid, oleic acid and linoleic acid. The unsaponified matter contains waxes and colouring matter. The root contains a red colouring matter. Phytochemicals reported in *L. inermis* L. are listed in **Table 2** with their structures.

TABLE 2: PHYTOCHEMICAL STRUCTURES PRESENT IN *L. INERMIS* L.

S no.	Chemical Name	Chemical Structure
1.	Scopoletin	 <p>Scopoletin is a coumarin found in root of plants in the genus scopolia.</p>
2.	β -sitosterol	 <p>β-Sitosterol is one of several phytosterols (plant sterols) with chemical structures similar to that of cholesterol. It is used in treating hypercholesterolemia</p>
3.	Luteolins	 <p>Luteolin can be found in <i>Terminalia chebula</i>. It is most often found in leaves, but it is also seen in rinds, barks, clover blossom, and ragweed pollen</p>
4.	Arachidic acid	 <p>Arachidonic acid (AA, sometimes ARA) is a polyunsaturated omega-6 fatty acid 20:4(ω-6). It is the counterpart to the saturated arachidic acid found in peanut oil</p>
5.	Linoleic acid	 <p>Linoleic acid belongs to one of the two families of essential fatty acids; it is an unsaturated n-6 fatty acid.</p>


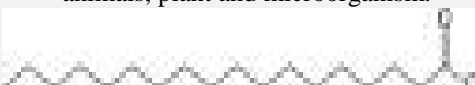
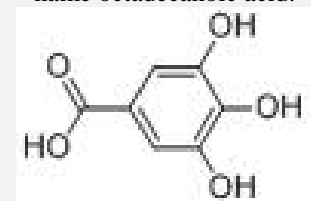
6.	Palmitic acid		Palmitic acid also called hexadecanoic acid, is the most common fatty acid found in animals, plant and microorganism.
7.	Stearic acid		Stearic acid is the saturated fatty acid with an 18-carbon chain and has the IUPAC name octadecanoic acid.
8.	Gallic Acid		Gallic acid is a trihydroxybenzoic acid, a type of phenolic acid, a type of organic acid, also known as 3,4,5-trihydroxybenzoic acid, found in gallnuts, sumac etc.

TABLE 3: PHYSICAL ANALYSIS OF *L. INERMIS* L. ⁵⁰
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Parameters	Values
Alcohol soluble extractive value	3.8 % w/w
Aqueous extractive value	5.0 % w/w
Loss on drying	4.5 % w/w
Total ash	14.60 % w/w
Acid insoluble ash	4.50 % w/w
Water soluble ash	3.0 % w/w
Swelling Index	Absent
Foaming Index	Less than 100
Ph 1% solution	7.22
Ph 10% solution	7.53
Extractive value	Hot Extraction (w/w)
Methanol	12.34 %
Aqueous	15.50 %

TABLE 4: PRELIMINARY PHYTOCHEMICAL TEST OF ALCOHOLIC EXTRACTS OF *L. INERMIS* L. ⁵⁴⁻⁵⁵

Phytochemical Tests	Results
Test for Alkaloids	-
Test for Glycosides	+
Test for Carbohydrates	+
Test for Saponins	-
Test for Fats & oils	-
Test for Volatile oils	-
Test for Tannis & phenolic compounds	+
Test for Protein	-
Test for Gums & mucilage	+
Test for Steroids	-

Morphological characters: The leaf of *Lawsonia inermis* L. is short, smooth, compound, ovate-lanceolate, acute, symmetrical, entire, pinnate, opposite, sweet smelling, characteristics or bitter in taste and varies in length, Lawsonia is mainly present in the marginal vein or petiole in large quantity ^{56, 57, 58}. Fig. 1 shows a photograph of *Lawsonia inermis* L.

Microscopic characters:

1. The leaf of *Lawsonia inermis* L. is short and smooth. The midrib is distinct from the lamina. It is broadly shallow on the adaxial side and convex on the abaxial side.
2. It also consists of unicellular covering trichome. Diacytic stomata are present on both the surface.
3. The leaf of *Lawsonia inermis* L. is dorsiventral as oblong palisade cells are present below the upper epidermis and absent on lower epidermis.
4. Tannin is seen in some of the cells. The vascular strand is single, small, collateral and hemispherical in shape.
5. It consists of a thick horizontal band of xylem and a fairly wide band of phloem. Xylem elements are narrow, angular, thin walled and somewhat diffuse.
6. The lamina is uniformly flat with even surface. Both adaxial and abaxial epidermal layers are thin and distinct.
7. The mesophyll tissue is differentiated into palisade and spongy parenchyma.
8. Some of the epidermal cells are smaller and have dark tannin content. The stomata are present on both surfaces.

9. Stomata are abundant. The stomata are Dicytic type. Each stoma is surrounded by two subsidiary cells the long axis of which is perpendicular to the long axis of stoma pore. The stomata are elliptical with wide opening.

10. The surface of the petiole is even and smooth. The epidermal layer is thin and very distinct.

11. The ground tissue is homogeneous and parenchymatous, the cells are thin walled and compact⁵⁹. **Figure 2** shows a photograph of T.S of *Lawsonia inermis* L. leaf.

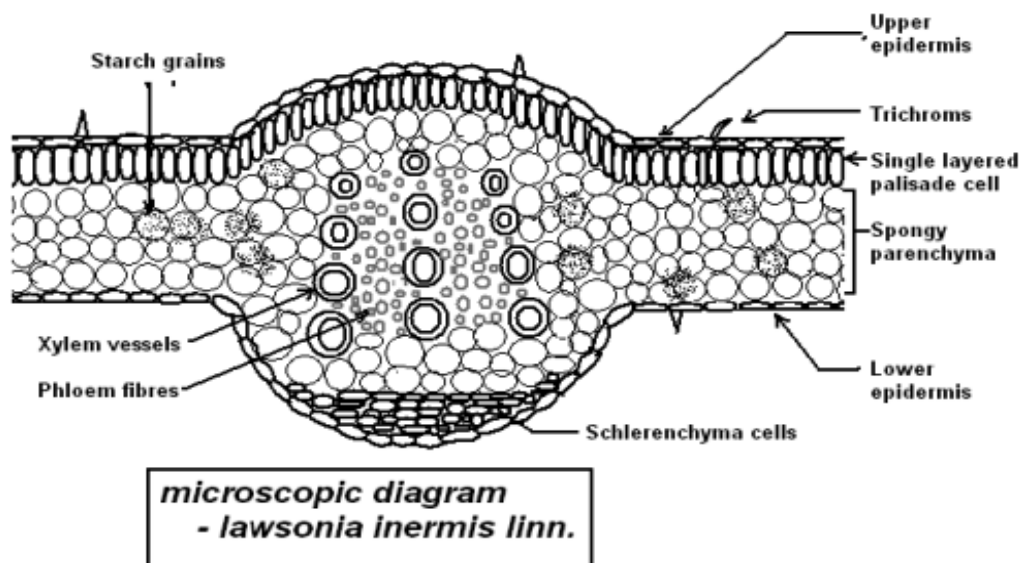


FIG. 2. PHOTOGRAPH OF T.S OF *LAWSONIA INERMIS* L. LEAF

CONCLUSION: The widespread survey of literature exposed that *L. inermis* L. is highly regarded as a universal solution in the herbal medicine with diverse pharmacological activity range. This versatile medicinal plant is the unique resource of various types of chemical compounds, which are responsible of the various activities of the plant. Hence extensive investigation is needed to develop their therapeutic utility to combat diseases. As the global scenario is now altering towards the use of non-toxic plant products having traditional medicinal use, development of modern drugs from *L. inermis* should be emphasized for the organize of various diseases. Further evaluation needs to be carried out on *L. inermis* L. in order to discover the concealed areas and their practical clinical applications, which can be used for the benefit of the mankind.

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REFERENCES:

1. Bandyopadhyay U, Biswas K, Chattopadhyay I and Banerjee RK: Biological activities and medicinal properties of neem (*Azadirachta indica*). *Currnt Sci* 2002; 82(11): 1336-1345.
2. Goyal BR, Goyal RK and Mehta AA: Phyto-Pharmacognosy of *Archyranthes aspera*: A Review. *Pharmacog Rev* 2008; 1: 1.
3. Padma TV: India Ayurveda. *Nature* 2005; 436-486.
4. Kasture SB, Une HD, Sarveiyal VP and Pal SC: Nootropic and anxiolytic activity of saponins of *Albizia lebbeck* leaves. *Pharmacology, Biochemistry and Behavior* 2001; 69: 439-444.
5. Hanna R, Maciej JN, Lapinsky L and Adamowicz L: Molecular structure and infra-red spectra of 2-hydroxy-1,4-naphthaquinone; Experimental matrix isolation and theoretical Hatree-Fock and post Hatree-Fock study. *Spec Act* 1998; 54: 1091-103.
6. Lavhate MS and Mishra SH: A review: nutritional and therapeutic potential of *Ailanthus excelsa*. *Pharmacog Rev* 2007; 1(1): 105-113.
7. Kirtikar KR and Basu BD: 2005. Indian Medicinal Plants. Second edition. International book distributors, Dehradun, vol-II, 1076-1086.
8. The PLANTS Database, database (version 5.1.1) 2000. National Plant Data Center, NRCS, USDA. Baton Rouge, LA 70874-4490 USA.
9. Dev S: 2006. A selection of prime Ayurvedic Plant Drugs, Ancient- modern concordance. Anamaya Publishers, New Delhi, 276-279.
10. Jallad KN and Jallad CE: Lead exposure from the use of *Lawsonia inermis* (Henna) in temporary paint-on-tattooing

- and hair dyeing. Science of the Total Environment 2008; 397: 244-250.
11. Nadkarni KM: 1982. Indian Materia Medica, Vol. 1. Popular Book Depot, Bombay, India, 730-773.
 12. Nayak BS, Isitor G, Davis EM and Pillai GK: The evidence based wound healing activity of *Lawsonia inermis* Linn. Phytotherapy Research 2007; 21: 827-831.
 13. Khare CP: Indian Medicinal Plants: An Illustrated Dictionary. Springer 2007; 366.
 14. Gogte VM: Ayurvedic Pharmacology and Therapeutic uses of Medicinal plants 2000; 686-687.
 15. Abdulmoneim MA: Evaluation of *Lawsonia inermis* Linn. (Sudanese Henna) leaf extract as an antimicrobial agent. Research Journal of Biological Sciences 2007; 2: 417-423.
 16. www.crescentbloom.com
 17. Chetty KM: 2008. Flowering plants of Chittoor, Edn 1, Andhra Pradesh, pp. 132.
 18. Khan MM, Ali A, Jain DC, Bhakuni RS, Zaim M and Thakur RS: Occurrence of some antiviral sterols in *Artemisia annua*. Plant Sci 1991; 75(2): 161-165.
 19. Nayak BS, Isitor G, Davis EM and Pillai GK: The evidence based wound healing activity of *Lawsonia inermis* Linn. Phytotherapy Research 2007; 21(9): 827-831.
 20. Muhammad HS and Muhammad S: The use of *Lawsonia inermis* Linn. (Henna) in the management of burn wound infection. African Journal of Biotechnology 2005; 4: 934-937.
 21. Sultana N, Choudhary MI and Khan AJ: Protein glycation inhibitory activities of *Lawsonia inermis* and its active principles. Enzyme Inhib Med Chem 2009; 24(1): 257-261.
 22. Syamusudin I and Winarno H: The effect of Inai (*Lawsonia inermis* Linn) leaves extract on blood sugar level: An experimental study. Research Journal of Pharmacology 2008; 2: 20-23.
 23. Arayne MS, Sultana N, Mirza AZ, Zuberi MH and Siddiqui FA: In vitro hypoglycemic activity of methanolic extract of some indigenous plants. Pak J Pharm Sci 2007; 20(4): 268-273.
 24. Iyer MR, Pal SC, Kasture VS and Kasture SB: Effect of *Lawsonia Inermis* on memory and behavior mediated via monoamine neurotransmitters. Indian Journal of Pharmacology 1998; 30: 181-185.
 25. Abdulmoneim MA: Evaluation of *Lawsonia inermis* Linn. (Sudanese Henna) leaf extract as an antimicrobial agent. Research Journal of Biological Sciences 2007; 2: 417-423.
 26. Al-Rubaiy KK, Jaber NN, Al-Mhaawe BH and Alrubaiy LK: Antimicrobial Efficacy of Henna Extracts. Oman Medical Journal 2008; 23(4): 253-6.
 27. Habbal OA, Al-Jabri AA, El-Hag AH, Al-Mahrooqi ZH and Al-Hashmi NA: In vitro antimicrobial activity of *Lawsonia inermis* Linn (henna). A pilot study on the Omani henna. Saudi Med J 2005; 26(1): 69-72.
 28. Awadh ANA, Julich WD, Kusnick C and Lindequist U: Screening of Yemeni medicinal plants for antibacterial and cytotoxic activities. Journal of Ethnopharmacology 2002; 74: 173-179.
 29. Ghosh A, Das BK, Roy A, Mandal B and Chandra G: Antibacterial activity of some medicinal plant extracts. J Nat Med 2008; 62(2): 259-262.
 30. Yogisha S, Samiulla DS, Prashanth D, Padmaja R and Amit A: Trypsin inhibitory activity of *Lawsonia inermis*. Fitoterapia 2002; 73: 690-691.
 31. Endrini S, Rahmat A, Ismail P and Taufiq YYH: Comparing of the Cytotoxicity Properties and Mechanism of *Lawsonia inermis* and *Strobilanthes crispus* extract against several cancer cell lines. Journal Medical Science 2007; 7: 1098-1102.
 32. Sauriasari R, Sano K, Horita M, Wang B and Ogino K: Cytotoxicity of lawsone and cytoprotective activity of antioxidants in catalase mutant *Escherichia coli*. Toxicology 2007; 235: 103-111.
 33. Dasgupta T, Rao AR and Yadava PK: Modulatory effect of henna leaf (*Lawsonia inermis*) on drug metabolising phase I and phase II enzymes, antioxidant enzymes, lipid peroxidation and chemically induced skin and forestomach papillomagenesis in mice. Molecular and Cellular Biochemistry 2003; 245: 11-22.
 34. Prakash D, Suri S, Upadhyay G and Singh BN: Total phenol, antioxidant and free radical scavenging activities of some medicinal plants. International Journal of Food Sciences and Nutrition 2007; 58: 18-28.
 35. Khodaparast H, Hosein M and Zinab D: Phenolic Compounds and Antioxidant Activity of Henna Leaves Extracts (*Lawsonia Inermis*) World Journal of Dairy & Food Sciences 2007; 2(1): 38-41.
 36. Mikhael BR, Badria FA, Maatooq GT, and Mohamed MA: Antioxidant and Immunomodulatory Constituents of Henna Leaves. Z. Naturforsch 2004; 9: 468-476.
 37. Ostovari A, Hoseinie SM, Peikari M, Shadizadeh SR and Hashemi SJ: Corrosion inhibition of mild steel in 1 M HCl solution by henna extract: A comparative study of the inhibition by henna and its constituents Corrosion Science 2009; 51: 1935-1949.
 38. Ali BH, Bashir AK and Tanira MO: Antiinflammatory, antipyretic and analgesic effects of *Lawsonia inermis* L. (henna) in rats. Pharmacol 1995; 51: 356-363.
 39. Sharma VK: Tuberculostatic activity of henna *Lawsonia inermis* Linn. Tubercle 1990; 71: 293-296.
 40. Ahmed S, Rahman A, Alam A, Saleem M, Athar M and Sultana S: Evaluation of the efficacy of *Lawsonia alba* in the alleviation of carbon tetrachloride induced oxidative stress. Journal of Ethnopharmacol 2000; 69: 157-164.
 41. Anand KK, Singh B, Chand D and Chandan BK: An evaluation of *Lawsonia alba* extract as hepatoprotective agent. Tubercle 1990; 71(4): 293-195.
 42. Mikhael BR, Badria FA, Maatooq GT and Amer MM: Antioxidant and immunomodulatory constituents of henna leaves. Zeitschrift fuer Naturforschung Section C Journal of Biosciences 2004; 59: 468-476.
 43. Endrini S, Rahmat A and Patimah Ismail P: Anticarcinogenic Properties and Antioxidant Activity of Henna (*Lawsonia inermis*) Journal of Medical Sciences 2002; 2(4): 194-197
 44. Singh A and Singh DK: Molluscicidal activity of *Lawsonia inermis* and its binary and tertiary combinations with other plant derived molluscicides. Indian J Exp Biol 2001; 39(3): 263-268.
 45. Singh VK and Pandey DK: Fungitoxic studies on bark extract of *Lawsonia inermis* against ringworm fungi. Hindustan Antibiot Bull 1989; 31(1- 2): 32-35.
 46. Wurochekke AU, Chechet G and Nok AJ: *In-vitro* and *In-vivo* antitrypanosomal brucei infection in mice. J Med Sci 2004; 4(3): 236-239.
 47. Chang H and Suzuka SE: Lawsone (2-OH-1, 4-naphthoquinone) derived from the henna plant increases the oxygen affinity of sickle cell blood. Bio-Chem Biophys Res 1982; 107: 602-608.
 48. Aguwa CN: Toxic Effects of the Methanolic Extract of *Lawsonia inermis* Roots. International J Crude Drug Res 1987; 25: 241-245.
 49. Nadhkarni KM.: Textbook of Pharmacognosy, Srishti Book Distributors, 2004: pp. 214.
 50. Karnick CR: Pharmacopoeial Standards of Herbal Plants, Indian Books Center, 2002: pp. 215.

51. Trease GE and Evans WC: Pharmacognosy, Bailliere Tindall, 1983: pp. 300- 244.
52. Indian Pharmacopoeia, Ministry of Health and Family Welfare, Government of India, Controller of Publication, 1996: A53-54.
53. Becket AH and Setnlake JB: Practical Pharmaceutical Chemistry, CBS Publication, 1983: pp. 333-336.
54. Mukherjee PK: Quality Control of Herbal Drug, Business Horizons, 2002: pp. 187-191.
55. Agrawal SS and Paridhavi M: Herbal Drug Technology, Universities Press (India) Private Ltd., 2007: pp. 625-638.
56. Nadhkarni KM: Textbook of Pharmacognosy, Srishti Book Distributors, 2004: pp. 214.
57. Kokate CK: Practical Pharmacognosy, Vallabh Prakashan, 2001: pp. 218.
58. Panigrahi AK and Sahu A: Glossary of Useful and Economical Important Plants, New Central Book Agency, 1998: pp. 60.
59. Bhattacharjee SK and De LC: 2003. Medicinal Herbs and Flowers, Aavishkar Publishers and Distributors, 2003: pp. 366.

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