Pharmacological Review of Mimusops Elengi Linn

PunamVilas kamble¹, Santosh Waghmare², Ashwini andhale³, Dr. H.V.Kambale⁴

^{1, 3, 4} Dept of pharmacology

²Dept of pharmaceutical chemistry

^{1, 2, 3, 4}Loknete Shri DadapatilPharate College of Pharmacy ,Mandavganpharata.

Abstract- Mimusopselengi is Indian native plant and is used for a long time in the history of the medicine. Plant was well studied in majority of the world because of its high potential medicinal value. Traditionally all different part of this plant, namely leaf, root, fruit, seed, bark and flower are used to cure various kinds of disorders. Mimusopselengi Linn. is traditionally used as antianxiety, cytotoxic, antimicrobial, antioxidants etc. Also, conventionally, Alstoniascholaris Linn. is used in treating cancer, psychosis, nociception etc. The present study was designed to appraise neuropharmacological and cytotoxic potential of M. elengi and A. scholaris leaves. Hole cross and hole board tests were conducted for assessing sedative effect, thiopental sodium induced sleeping time test for hypnotic property, elevated plus maze (EPM) test for anxiolytic potential, tail suspension test for antidepressant effect and finally brine shrimp lethality bioassay for evaluating cytotoxic potential of both extracts. In both hole cress and hole board tests, M. elengi leaves (MEL) displayed greater locomotion reduction compared to A. lower of Mimusopselengi were reported for treatment of various human ailments in traditional system of medicine. Pharmacological activities like antimicrobial, antifungal, antioxidant and free radical scavenging, anti-inflammatory analgesic, antipyretic, antiurolithiatic, cytotoxic, diuretic, neuroprotective, antiamnesic. cognitive enhancing, antihyperglycemic, antihyperlipidemic, hypotensive, antiulcer, anthelmintic, antitumor, wound healing, larvicidal activities have been scientifically evaluated for various parts of this plant. A number of phytochemical constituents have been identified in this plant that may be responsible for its pharmacological activities.

Keywords- elengilenn, taxonomy, chemical constituents, pharmacological activity.

I. INTRODUCTION

Molsari (Mimusopselengi L.) is a large glabrous evergreen Indian origin tree attaining a height of 12-15 m distributed in peninsular region, western and eastern ghats and cultivated in the plains e.g., tropical forests in south Asia and in India. Plant is cultivated for its ornamental appearance, elegant look, shade and for fragrant flowers.1 The Plant has vast description in Unani literature as Molsari. It is described as large tree like mauwah, kherni and cheeku. This tree gives characteristic cool shade and fragrant flowers.

TAXONOMY:

- Kingdom: Plantae
- Order: Ericales
- Family: Sapotaceae
- Genus: Mimusops
- Species: Elengi
- Botanical Name: Mimusopselengi
- Vernacular name: English :bullet wood, Spanish cherry, Hindi mul sari, Tamil magadam.
- Part use: steam ,bark, leaves, flower , fruit ,seed
- Distribution: memusopselengi tree is a native of Western pininsula.
- Traditional medicinal use: the bark is used for cardiotonic ,stomachic ,tonic, flower is cooling, fragrant flower.



Fig no 1: Mimusopelengi

II. CHEMICAL COMPOSITION

Bark of M. elengi contains tannin, some caoutchouc, wax, coloring matter, starch and ash forming inorganicsalts[4]. Saponin was isolated from the ethanolic extract of the bark, which on hydrolysis yielded β -amyrin and bassic acid. Hexane soluble fraction of the alcoholic extract yielded taraxerone, taraxerol, α -spinasterol, sodium ursolate and betulinic acid, where as hexane insoluble fraction yielded β -D-glucoside of β -sitosterol and the aqueous extract, gave quercitol. Other pentacyclic triterpenoids betulic acid (2 -167), lupeol (4- 167), taraxerol (3-167) and ursolic acid (3-167). Fatty acid ester of α -spinasterol(3-167) was also isolate from bark[8]. The petroleum ether extracts of stem bark yielded α -spinasterol and taraxerol, the same was also isolated from wood portion of M. elengi along with meso-inositol.

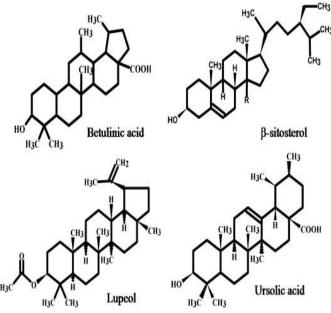


Fig no 2: Chemical constituents of M.elengi.

III. BIOLOGICAL ACTIVITY

M.elengi: The leaf extract showed in vitro antibacterial activity against Bacillus anthracis, Bacillus mycoides, Bacillus pumilus, Bacillus subtilis, Salmonella paratyphii, Staphylococcus albus, Vibraechlorae, and andXanthomonasmalvacearum, the inhibition was significant against Xanthomonascampestris and Bacillus anthracis.

IV. PHARMACOLOGICAL ACTIVITIES

• Antioxidant activity:

Saaha et al., evaluated the Antioxidant potential of the methanol extract of the leaves of Mimusopselengi by using 1, 1-diphenyl-2- picrylhydrazyl (DPPH) scavenging assay, reducing power and total antioxidant capacity. The extract showed significant activities in all antioxidant assays compared to the reference antioxidant ascorbic acid in a dose dependent manner. In DPPH scavenging assay the IC50 value of the extract was found to be 43.26μ g/ml while theIC50 value of the reference standard ascorbic acid was 58.92μ g/ml. Total antioxidant activity was also found to increase in a dose dependent manner. M. elengi extract also showed strong reducing power.

In-vitro anti-inflammatory activities:

Kar et al., assessed the antioxidant and invitro antiinflammatory activities of alcoholic extract of Mimusopselengi leaves. The leave extract exhibited dose dependent free radical scavenging property in peroxynitrite, superoxide and hypochlorous acid models and the IC50value were found to be (205.53 \pm 2.30), (60.5 \pm 2.3), (202.4 \pm 5.3) µg/mL respectively.

• ontotoxic activity:

The cytotoxic effect of ethanolic extract of barks of M.elengi was investigated on meristimatic cells of root tips of Allium cepa. The experiment was carried out by using different concentrations (2.5, 5, 10 mg/ml) of standard cytotoxic drug cyclophosphamide and ethanolic extract. After 48 h and 96 h root length and mitotic index were calculated.

• Antibacterial activity:

The antibacterial activity of petroleum ether, chloroform, ethyl acetate and methanol extracts of the flowers of Mimusopselengi were screened againsvarious pathogenic Gram positive and Gram negative bacterial strains viz. Bacillus cereus, Enterobacterfaecali, Salmonella paratyphi, Staphylococcus aureus, Escherichia coli, Proteus vulgaris, Klebsiellapneumoniae, Pseudomonas aeruginosa and Serratiamarcescens by 'agar well diffusion' method.

• Hypotensive activity:

The methanolic extract of Mimusopselengi showed hypotensive activity in anaesthetized rats. On intravenous administration (i.v.) at a dose range of 2-16 mg/kg, it produced about a 7-38% fall in mean arterial blood pressure, in a dosEdependent manner. The effect was independent of adrenergic, muscarinic and histaminergic receptors.

• Wound Healing Activity:

In an study wound healing activity of extract of bark part of Mimusopselengi was evaluated. A methanolic extract was examined in the form of ointment in three types of wound models on mice: the excision, the incision and dead space wound model. The extract ointments showed considerable response in all the above said wound models as comparable to those of a standard drug Betadine ointment in terms of wound contracting ability, wound closure time, tensile strength and dry granuloma weight.

V. CONCLUSION

In spite of our great dependence on modern medicines and tremendous advances in synthetic drugs, a large portion of the world population still likes drugs of plants origin. M. elengi (Bakul) is one of the most important medicinal plants used in preparations of Ayurveda because of having a number of medicinal properties. It is the source of a variety of biologically active phytoconstituents which are responsible for antimicrobial, antioxidant, antihyperglycemic, anticancer and protective effects on various vital organs such as nerves, heart, kidney and liver.

REFERENCES

- [1] Ganu G, Garud A, Agarwal V, Talele S, JadhavS, Kshirsagar A. Anticonvulsant activity of a Mimusopselengi in experimental animals. J Pharm Res 2011; 4(9):2938-
- [2] Satishchandra, Sumithra M. Synergistic effect of Mimusopselengi and Moringa on high fat diet induced atheroma in rats. Int J AdvPharmaceut Res 2011; 2(6):293–300.
- [3] Prabhat, Ajaybhan, Navneet, Chauhan A. Evaluation of Antimicrobial Activity of Six Medicinal Plants against Dental Pathogens. Report Opinion 2010; 2(6):37. Ali MA, Mozid MA, YeasminMS, Khan AM, Sayeed MA. An Evaluation of
- [4] Antimicrobial Activities of Mimusopselengi Research Journal of Agriculture and
- [5] Biological Sciences 2008; 4(6): 871-874.
- [6] Gupta N, Jain U K. Investigation of Wound Healing Activity of Methanolic Extract of Stem Bark of MimusopselengiAfr J Tradit Complement Altern Med. 2011; 8(2): 98-103.
- [7] Ganu G, Jadhav S. In Vitro Antioxidant and In Vivo Antihyperglycemic Potential of Mimusopselengi in Alloxan-Induced Diabetes in Mice. Journal of Complementary and Integrative Medicine 2010; 7(1): 1553-3840.
- [8] Behbahanian DS, Malik A, Jahan N. Hypotensive effect of the methanolic extract of Mimusopselengi in normotensive rats. Phytomedicine 1999; 6(5):373.
- [9] Shah PJ, Gandhi MS, Shah MB, Goswami SS, Santani D. Study of Mimusopselengi bark in experimental gastric ulcers. J Ethnopharmacol 2003; 89(2-3):305.
- [10] Jana GK, Dhanamjayarao M, Vani M. Evaluation of anthelmintic potential of Mimusopselengi (Sapotaceae) leaf. J Pharm Res 2010; 3(10):2514-2515.

- [11] Alkilany AM, Lohse SE, Murphy CJ (2013) The gold standard: gold nanoparticle libraries to understand hydrogenation catalyst. AccChem Res 46:650–661.
- [12] Bag BG, Dash SS (2011) First self-assembly study of betulinic acid, a renewable nano-sized, 6-6-6-5 pentacyclicmonohydroxytriterpenic acid. Nanoscale 3:4564–4566
- [13] Zhang Y, Cui X, Shi F, Deng Y (2012) Nano-gold catalysis in fine chemical synthesis. ChemRevchemical synthesis. Chem Rev 112:2467–2505.