

Pharmacophore

(An International Research Journal)

Available online at <http://www.pharmacophorejournal.com>

Review Article

PHYTOPHARMACOLOGICAL ASPECTS OF *SALACIA CHINENSIS*

U.A. Deokate*, S.S. Khadabadi

Govt. College of Pharmacy, Kathora Naka,
Amravati-444604 (M.S.), India

ABSTRACT

Salacia chinensis Linn. (Family Celastraceae) commonly known as Saptrangi and commonly used herb in Ayurvedic medicine. The present review is an attempt to compile information on pharmacological and phytochemical aspects of *Salacia chinensis* Linn. Its roots biologically active compounds such as triterpenes, phenolic compounds, glycosides and coloring agents which show various medicinal properties. The root extract shows various activities like, antioxidant, anticaries, antiulcer, antidiabetic, hypoglycemic, antiobesity and skin lightening agent. This review will help reader with detail understanding of Salacia root's properties.

Keywords: *Salacia chinensis*, Saptrangi, Root, Phytochemistry, Pharmacology.

INTRODUCTION

Salacia chinensis Linn. (Synonyms: *Salacia prinoidea*) Family: Celastraceae (Spike-thorn family) commonly called as Saptrangi, Dimal, Modhupal, Ingli, Cherukuranti, Nisul-bondi. This is a small erect or straggling tree or large, woody, climbing shrub found almost throughout India including Andaman & Nicobar Islands.¹

Three species of *Salacia* i.e. *S. chinensis*, *S. reticulata*, *S. oblonga* are used traditionally in Ayurveda, Unani systems as antidiabetic agent. Preclinical research and isolated clinical trials studying these effects have been promising. Fruits & Roots are the useful parts. Ripe fruits are eaten. Roots have been used as an antidiabetic drug. *Salacia chinensis* have been used in India and in other countries as a tonic, blood purifier and to treat amenorrhea and dysmenorrhea. Its root bark was used in gonorrhoea, rheumatism and skin diseases. Its aqueous extract showed significant hypoglycemic activity. Root bark boiled in oil or as decoction or as powder is used for the treatment of rheumatism, gonorrhoea, itches, and asthma, thirst and ear diseases.²⁻⁵ The root is dark yellow externally and light yellow internally. It has characteristic odor and bitter in taste. The TS of root (figure 1) shows wavy cork, cortex consisting of brown matter, uniseriate and few biseriate medullary rays and vascular bundle consists mainly of secondary xylem and phloem. Pith, Pericycle and endodermis is absent. Starch is present in the cortex region. The standardization parameters like ash value and extractive value have been also studied. The ash value results for this plant are Total ash 4.825%w/w; Water-soluble ash 2.75%w/w Acid-insoluble ash 3.5 %w/w. The extractive value results are found to be 3.275%w/w in Water-soluble extractive value and 1.8%w/w in Alcohol-soluble extractive value [6]. Table 1 is showing Fluorescence analysis of root powder.

The roots contains the phytoconstituents like alkaloides, glycosides, polyphenols, flavanoides, coumarins, proteins, carbohydrates, gums and mucilage, fixed oil and volatile oil. Triterpenoids like lupanes, hopanes, friedelanes are abundant in root and stem of plant. Salacinol from the stems of *S. chinensis* was found to alpha- glucosidase inhibitor. Mangiferin showed inhibitory effect on rat lense aldose reductase. Figure 2 and Table 2 has given is giving idea about chemical composition of salacia roots.

Recent studies have demonstrated that *Salacia* roots are very useful in type 2 diabetes and obesity-associated hyperglycemia, dyslipidemia and related cardiovascular complications and it may be due to the fact that it modulate multiple targets⁷ like peroxisome proliferator-activated receptor-alpha-mediated lipogenic gene transcription, angiotensin II/angiotensin II type 1 receptor, alpha-glucosidase, aldose reductase and pancreatic lipase. These activities are due to the constituents like mangiferin, salacinol, kotalanol and kotalagenin 16-acetate.⁸⁻¹⁰ Clinical trial studies have also been carried out to confirm the effects. Table 3 is explaining the all activities of salacia roots.

Table 1: Fluorescence analysis of root powder

Treatment	Day light	UV light (254nm)
Powder as such	Yellow color	Light green
Powder + 1N NaOH(Aq.)	Brown	Dark brown
Powder + 1N NaOH(Alc.)	Yellowish brown	Light yellow
Powder + 1N HCL	Green	Light green
Powder + Iodine	Dark brown	Brown
Powder + Ammonia	Yellow	Greenish yellow
Powder + 5% FeCl3	Dark yellow	Dark brown
Powder + 1N H2SO4	Black	No color

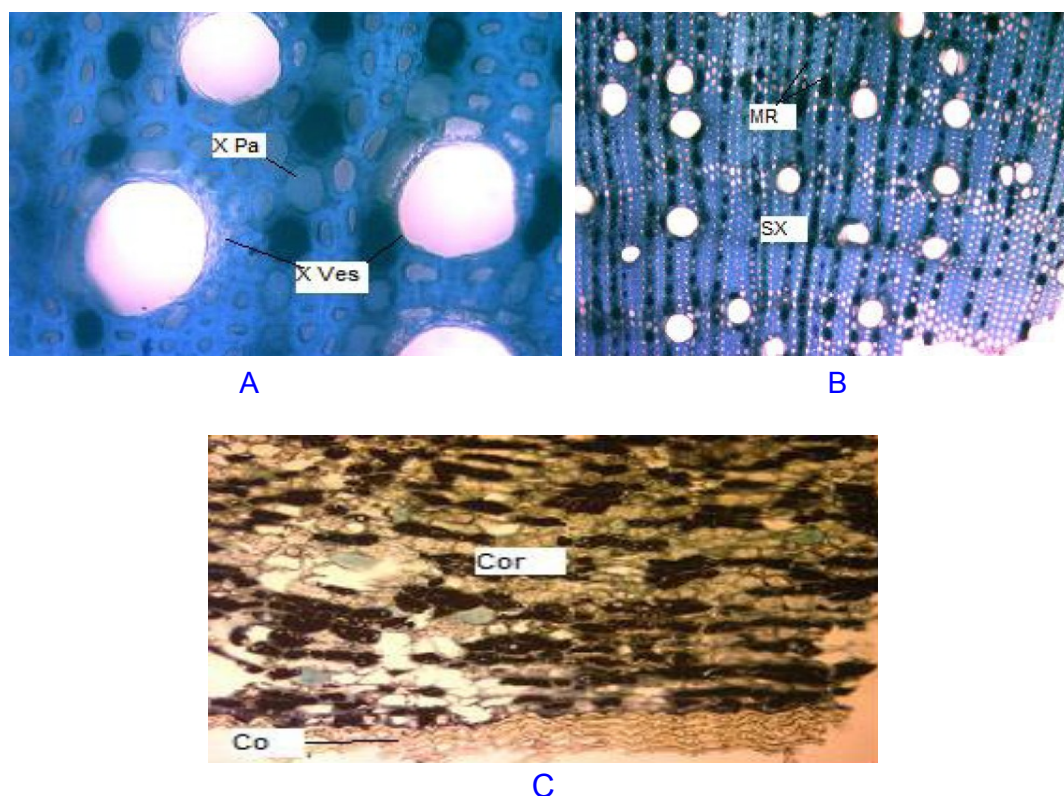


Figure 1: A, B & C: *Salacia* roots microscopy showing vascular bundles, medullary rays, and Annular Rings.

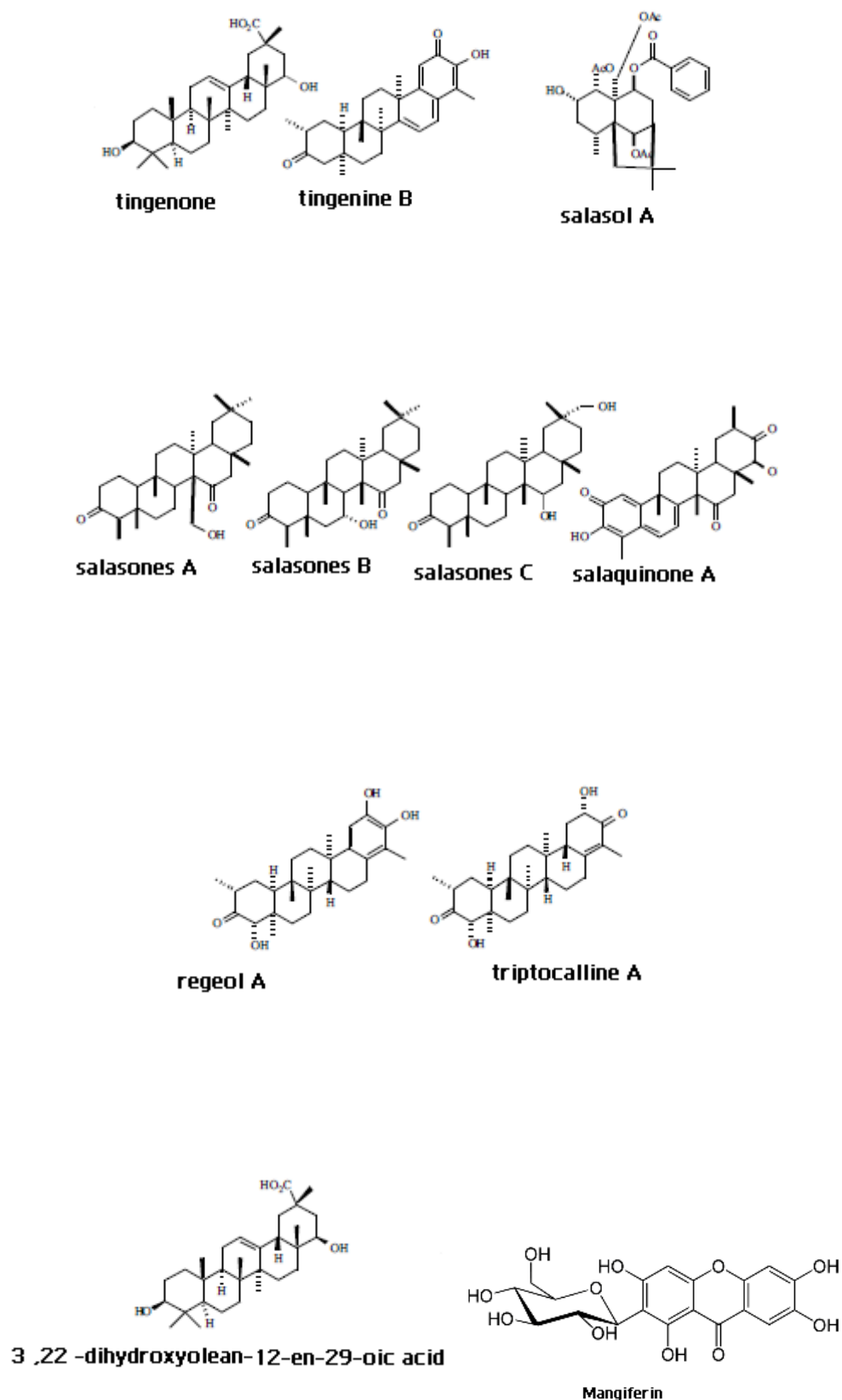


Figure 2: Chemical constituents of *S. chinensis*

Table 2: Phytochemistry of roots of *Salacia chinensis* Linn.

Constituents	Plant part
Triterpenes foliasalacins D1 (I), D2 (II), and D (III), Phenolic glycosides named foliachinenosides A1 (1), A2 (2), A3 (3), B1 (4), B2 (5), C (6), and D (7); Foliasalaciosides A1,A2, B1, B2,C and D; megastigmane glycosides named foliasalaciosides E1 (I)-I (1-7)	Leaves ¹¹
Megastigmane glycosides foliasalaciosides A1 (1) (I), A2 (2), B1 (3), B2 (4), C (5), and D (6)	Leaves ¹² (Methanolic extract)
Triterpenes like 28- hydroxy -3- oxo- 30- lupanoic acid, 29- nor-21 α - H- hopane- 3,22- dione, 21 α - H- hop- 22(29)- ene- 3 β , 30- diol, and betulin, 3 – oxo- lupane- 30 – al; Betuline, 29- nor- 21- α H- hopane- 3, 22- dione and 21- α H- hop- 22 (29)- ene- 3 β ,30- diol	Stems (n-hexane extract) ¹³
α -glucosidase inhibitor salacinol, Dimer(II), octaacetate, hexamethyl ether, Friedelane-type triterpenes, salasones D and E, norfriedelane-type triterpene, salaquinone B, polyacylated eudesmane-type sesquiterpine, salasol B; Two new friedelane-type triterpenes, salasones D and E, a new norfriedelane-type triterpene, salaquinone B, and a new polyacylated eudesmane-type sesquiterpine, salasol B,	Stems ¹⁴
Two new triterpenoids, named 7 α ,21 α -dihydroxyfriedelane-3-one (1) and 7 α ,29-dihydroxyfriedelane-3-one (2) and 21 α ,30-dihydroxyfriedelane-3-one	Stems ¹⁵ (ethyl acetate extract)
Friedelane-type triterpenes, salasones A, B, and C, norfriedelane-type triterpene, salaquinone A, acylated eudesmane-type sesquiterpine, salasol A	Stems (80% of methanolic extract) ¹⁶
3 β ,22 β -dihydroxyolean- 12- en- 29- oic acid, tingenone, tingenine B, regeol A, triptocalline A, and mangiferin	Stems (80% of methanolic extract) ¹⁷
1,3-diketofriedelane derivatives : six closely related triterpenes, P, Q, R, S, T and V	Root bark ¹⁸⁻¹⁹

Table 3: Pharmacology of roots of *Salacia chinensis* Linn.

S.No.	Activity	Model	Plant part	Conclusion
1	Antidiabetic activity ²⁰	Maltose or sucrose loaded rats	Methanolic extract of stems	Inhibitory effects on intestinal alpha-glucosidase, rat lens aldose reductase, formation of Amadori compounds and advanced glycation end-products, nitric oxide production from lipopolysaccharide-activated mouse peritoneal macrophage, and radical scavenging activities.
2	Antidiabetic activity ²¹	Streptozotocin (STZ) - induced diabetic rats.	<i>Salacia chinensis</i> and <i>coccinia indica</i> and <i>hipophae rhamnoides</i>	Found to be effective
3	Antihyperglycemic activity ²²	Streptozotocin (STZ) - induced diabetic rats.	Mangiferin purified from methanolic root ext. of <i>Salacia chinensis</i> (<i>S. chinensis</i>) mangiferin possess antidiabetic activity against STZ-induced diabetic rats.	Mangiferin is found responsible for antidiabetic activity.
4	Hypotensive activity ²³	Hypotensive activity in anesthetized female rats in estrus, and for vasodilator activities on isolated thoracic aortic rings in vitro.	Stem ethanolic extract	n-butanol extract from stems of <i>Salacia chinensis</i> possesses a hypotensive effect. The mechanism involved may be an indirect effect by stimulated release of nitric oxide from vascular endothelial cells and causes vasodilatation.
5	Hepatoprotective activity ²⁴	Wistar strain of albino rats of either sex against CCl ₄ induced	Root extract	Found to be effective
6	Anticaries activity ²⁵	Prevents glucan adhesion on tooth plane and inhibits glucosyltransferase activity, and can be used for	<i>Salacia extract</i>	<i>Salacia extract</i> inhibit sucrose-dependent biofilm formation by MS, similar to acarbose, and have potential as anti-plaque anti-caries agents

		preventing caries.		
7	Reproductive function activity [26]	Sprague–Dawley male & female rats	S. chinensis extract	no effects on the reproductive outcome such as estrous cycle of F0 females or any parameters for reproductive function or survival, growth, sensory reflex or function development of F1 pups even at a remarkably high dosage level, 2000 mg/kg/day,
8.	Anticancer activity [27]	Against the four cancer cell lines Hep-G2, LU, KB, and MCF-7.	Eight triterpenoids from this plant	The new compound showed good activity against all four tested cell lines.

CONCLUSION

Salacia chinensis is a traditional South and Southeast Asian herb medicine and has been reported to have an antidiabetic function via α -glucosidases inhibitory activity. The various active constituents have been found to affect multiple targets in diabetes, obesity and associated cardiovascular diseases through modulating PPAR- α -mediated lipogenic gene transcription and angiotensin II/angiotensin II type 1 receptor, inhibiting α -glucosidase, aldose reductase and pancreatic lipase. Although toxicological studies have suggested minimal adverse effects of this plant in rodents, a clinical trial is crucial to further confirm the safety of *Salacia* roots. In addition, mechanistic studies are necessary in order to know drug interaction of *Salacia* root with other therapeutic interventions.

REFERENCES

1. Mehra, Handa (1969), “*Researchers in Pharmacognosy in India*”, Bulletin of Punjab University, 20,275.
2. “*Encyclopedia of World Medicinal Plants*”, Vol.1, 1713, 2418
3. Almeida, MR & SM, Almeida (1994), “Identification of some plants from 'Hortus Malabaricus'”, *Journal of the Bombay Natural History Society*, 90 (3), 423-429.
4. Singh, A and Duggal, S (2010), “Salacia Species: Hypoglycemic Principles and Possible Role in Diabetis Management”, *Integrative Medicine*, 2(4).
5. Mehra, PN and Handa, SS (1967), “True Identity of Saptrangi Metab Abstract Casearia-Esculenta-D Casearia-Tomentosa-D Salacia-Chinensis-D Hypo Glycemic”, *Indian Journal of Pharmacy*, 29,341.

6. Dholwani, Kishor kumar; Saluja, Ajay K; Bhatt, Shibani and Chauhan, Namrata (2009), "Pharmacognostical Profiling of Roots of *Salacia prenoides* L", *Phcog. J.*, 1(3).
7. Yuhao, Lia; Tom Hsun-Wei, Huanga and Johji, Yamaharab (2008), "Salacia root, a unique Ayurvedic medicine, meets multiple targets in diabetes and obesity", *Life Sciences*, 82(21-22), 1045-1049.
8. Yoshikawa, M; Nishida, N; Shimoda, H; Takada, M; Kawahara, Y and Matsuda, H (2001), "Polyphenol constituents from *Salacia* species: Quantitative analysis of mangiferin with alpha-glucosidase and aldose reductase inhibitory activities", *Yakugaku Zasshi*, 121:371-378.
9. Matsuda, H; Morikawa, T; Toguchida, I; Yoshikawa, M (2002), "Structural requirements of flavonoids and related compounds for aldose reductase inhibitory activity", *Chemical and Pharmaceutical Bulletin (Tokyo)*, 50, 788-795.
10. Nadagouda, Smitha G; Karigar, Asif A; Joshi, VG and Sikarwar, Mukesh S (2010), "Validated HPTLC method for mangiferin in *Salacia chinensis*", *Journal of Pharmacy Research*, 1(3).
11. Yoshikawa Masayuki Zhang, Yi; Wang, Tao; Nakamura, Seikou and Matsuda, Hisashi (2008), "New triterpene constituents, foliasalacins A1-A4, B1-B3, and C, from the leaves of *Salacia chinensis*", *Chemical & Pharmaceutical Bulletin*, 56(7), 915-920.
12. Krishnan, V and Rangaswami, S (1967), "Chemical components of *Salacia chinensis* Linn. Stems and leaves", *Curr Sci.* 36, 596-597.
13. Krishnan, V and Rangswami, S (1967), "Proanthocyanidins of *Salacia chinensis* linn", *Tetrahedron Lett*, (26), 2441-2446.
14. BS, Joshi; VN, Kamat and N Viswanathan (1973), "Triterpenes of *Salacia prinoides*", *DC Tetrahedron*, 29(10), 1365-1374.
15. D, Rogers; DJ, Williamsa; BS, Joshib; VN, Kamatb and N, Viswanathanb (1974), "Structure of new triterpene ether from *Salacia prinoides* dc: x-ray investigation of the dibromo derivative", *Tetrahedron Letters*. 15(1), 63-66.
16. Tran Thi, Minha; Nguyen Thi Hoang, Anhb; Vu Dao, Thanga and Tran Van, Sungb (2008), "Study on Chemical Constituents of *Salacia chinensis* L. Collected in Vietnam Z", *Naturforsch.* (63b), 1411-1414.
17. T, Kishi A; Pongpiriyadacha, Y; Matsuda, H; Yoshikawa, MJ (2003), "Structures of new friedelane-type triterpenes and eudesmane-type sesquiterpene and aldose reductase inhibitors from *Salacia chinensis*", *Morikawa, Nat. Prod.* 66 (9), 1191-1196.
18. Yoshikawa, Masayuki and Morikawa, Toshio (2003), "Structures of New Friedelane- and Norfriedelane-Type Triterpenes and Polyacylated Eudesmane-Type Sesquiterpene from *Salacia chinensis* LINN. (*S. prinoides* DC, Hippocrateaceae) and Radical Scavenging Activities of Principal Constituents", *Chem Pharm Bull*, 51(9), 1051-1055.
19. Masayuki, Yoshikawa; Yi, Zhang; Seikou, Nakamura; Tao, Wang and Hisashi, Matsuda (2008), "The absolute stereostructures of three rare DB-friedobaccharane skeleton triterpenes from the leaves of *Salacia chinensis*", *Tetrahedron*. (64), 7347-7352.

20. Yoshikawa, M; Pongpiriyadacha, Y; Kishi, A; Kageura, T; Wang, T; Morikawa, T and Matsuda, H (2003), "Biological activities of *Salacia chinensis* originating in Thailand: the quality evaluation guided by alpha-glucosidase inhibitory activity", *Yakugaku Zasshi*. 123(10), 871-80.
21. Govind, Prasad Dubey; Aruna, Agarwal; Neers, Vyas; Victor, G and Rajamanickam (2010), "Herbal Formulation for the Prevention and Management of Diabetes Mellitus and Diabetic Micro-Vascular Complications".
22. Sellamuthu, PS; Muniappan, BP; Perumal, SM and Kandsamy, M (2009), "Antihyperglycemic effect of mengiferin in Streptozotocin induced Diabetic rats", *Journal of health sciences*.55 (2), 206-214.
23. Jansakul, C; Jusapalo, N and Mahattanadul, S (2005), "Hypotensive effect of n-butanol extract from stem of *Salacia chinensis* in rats", *Acta horticulturae*, 678, 107-114.
24. Asuti, Naveen (2010), "Hepatoprotective activity of ethanolic extract of root bark of *Salacia chinensis*", *Journal of Pharmacy Research*, 3(4), 833-834.
25. C, Vuong, and CI, Hoover (2010), "Inhibition of Sucrose-Dependent Biofilm Formation by α -Glucosidase Inhibitors".
26. Yang, Jihonga; Luo, Shaozhonga; Song, Jingfenga; Masakazu, Kobayashib; Junji, Akakib; Kousaku, Yamashitab; Makoto, Tamesadab and Tatsuo Umemurac, "Effects of *Salacia chinensis* extract on reproductive outcome in rats", *Food and Chemical Toxicology*.
27. Tran, Thi Minha; Nguyen Thi, Hoang; Anhb, Vu Dao Thanga, and Tran Van, Sungb (2010), "Study on Chemical Constituents and Cytotoxic Activities of *Salacia chinensis* Growing in Vietnam Z", *Naturforsch*. 65b, 1284-1288.