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Review Article

**FRANKINCENSE: A POTENTIAL ANTI-INFLAMMATORY
AGENT: AN OVERVIEW****Gh. Hassan Bhat¹, Mushtaq Ahmed Parray*², Ghulam Mohammad Jan³**¹Department of Botany, Govt. Degree College Dooru Anantnag J&K (India)²Department of Zoology, Govt. Degree College Shopian J&K (India)³Department of Chemistry, Govt. Degree College for Boys Anantnag J&K (India)**Abstract:**

Boswellia in the family *Burseraceae*, particularly *Boswellia sacra* (syn: *B. carteri*, *B. thurifera*, *B. bhaw-dajiana*), *B. frereana* and *B. serrata* (Indian frankincense.). Frankincense resin is edible and is used in traditional medicines in Africa and Asia for digestion and healthy skin. For internal consumption, it is recommended that frankincense be translucent, with no black or brown impurities. Exposure to 11-keto- β -boswellic acid (KBA), a lead boswellic acid in the novel solubilized frankincense extract Boswelan, is increased when taken with food.

Keywords: *Burseraceae*, resin, traditional medicines, translucent, boswellic acid**Corresponding author:****Mushtaq Ahmed Parray,**

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INTRODUCTION:

Since time immemorial, plants and their products have been the primary resource of food, shelter, clothing, flavors, and fragrances as also valuable ingredients for medicines for mankind. In this context, natural resins have played an important role. These have also been used as adhesives, as ingredients for cosmetic preparations, as fragrances in daily rituals and in religious ceremonies, as coating materials and also for their different curative powers [1–3]. In ancient times, Hindus, Babylonians, Persians, Romans, Chinese and Greeks as well as the people of old American civilizations used natural resins primarily for embalming and for its incense in cultural functions. They firmly believed that when these materials get in contact with fire, the smoke and the fragrance they produce not only soothe their souls but also please their gods. Burning of these natural resins had become an important component of their cultural life. They burned these resins during sacrificial ceremonies and in their daily rituals to prevent the influence of evil spirits on their souls or to honour the dead or living ones [4–6].

Boswellia serrata (Salai/Salai guggul) (Family: Burseraceae; Genus: *Boswellia*) is a moderate to large sized branching tree that grows in dry mountainous regions of India, Northern Africa and the Middle East [7,8]. The family of Burseraceae is represented in the plant kingdom with 17 genera and 600 species wide-spread in all tropical regions. There are about 25 known species belonging to Genus *Boswellia*, most of them occur in Arabia, northeastern coast of Africa and India. Since ancient times, three of these species have been considered as 'true Frankincense' producing trees [9,10].

Boswellia sacra Flueck, the first species, grows in South Arabia and is known amongst Arabians as 'maghrayt d' sheehaz' and the resin produced is known as 'luban dhakar'. *Boswellia carterii* Birdw, grows in Somalia and in the native language it is called 'moxor' and the resin produced is known as 'luban dhakar'. *Boswellia frereana* Birdw., is also a Somalian species and in the native language it is called 'jagcaar' and the resin produced is known as 'loban majdi' or 'maydi'. This is the most expensive brand of resin in the market [11]. Another resin producing species is *Boswellia serrata* Roxb., known as 'Indian olibanum', 'Indian frankincense', 'dhup' and 'salai' or 'salai guggul' is found in the middle and northern parts of Eastern India. It has been available as a high quality extract in India for nearly 25 years and marketed under the name Shallaki.

In India, the main commercial sources of *Boswellia serrata* are Andhra Pradesh, Gujarat, Madhya Pradesh, Jharkhand and Chhattisgarh. Regionally, it is also known by different names. It is tapped from

the incision made on the trunk of the tree, which is then stored in specially made bamboo basket. The semi-solid gum-resin is allowed to remain in the basket for about a month during which its fluid content locally known as 'ras' keeps flowing out. The residue, semi-solid to solid part, is the gum-resin which hardens slowly into amorphous, tear-shaped products with an aromatic scent. Then, it is broken into small pieces by wooden mallet or chopper and during this process all impurities including bark pieces etc. are removed manually. The gum-resin is then graded according to its flavour, colour, shape and size. Generally four grades i.e. Superfine, Grade I, Grade II and Grade III are available in the market. The fresh gum obtained from the tree is hot with pleasant flavour and slightly bitter in taste. It had been the 'frankincense' of ancient Egyptians, Greeks and Romans who used it as prized incense, fumigant as well as a multipurpose aromatic. It is generally used in making incense powder and sticks.

The oleo gum-resins contain 30–60% resin, 5–10% essential oils, which are soluble in the organic solvents, and the rest is made up of polysaccharides (~ 65% arabinose, galactose, xylose) which are soluble in water [12–14]. The resins have a fragrant aroma because of the presence of essential oils and these accounts for their commercial importance. The essential oil of gum-resin is one of the most commonly used oils in aromatherapy, paints and varnishes. Pure oleo gum-resin collected in the optimum season hardens slowly, retaining its golden colour and transparency. But the colour varies from golden brown to dark brown or dark greenish-brown depending on the locality, season, size of the tree and the wound-surface, collection process and storage. Darkening of colour of resin is also due to autoxidation, polymerization and enzymatic reactions. The resin is generally harvested all through the summer and autumn after the tree has been wounded in March or April. *Boswellia* tree can produce exudates in good quality only for three years. After this period, the quality of the collected resin decreases considerably. Therefore, the tree should be left to rest for some years after harvesting period.

HISTORICAL/TRADITIONAL APPLICATIONS:

Boswellia serrata is one of the ancient and most valued herbs in Ayurveda. "Gajabhakshya", a Sanskrit name sometimes used for *Boswellia*, suggests that elephants enjoy this herb as a part of their diet [15]. Three renowned ancient texts form the pillars of classical Ayurvedic Science, which has its roots in India: Charaka's Charaka Samhita (c.B.C. 700), the first fundamental medical text;

Susruta's *Susruta Samhita* (c.B.C. 600), which attempted to amass the entire medical knowledge, with special focus on surgery; and the two-volume tome comprising *Astanga Samgraha* and *Astanga Hridaya* (c.130-200 A.D.), written by Vagbhata the Elder and Vagbhata the Younger, which synthesized the works of Charaka and Susruta and summarized the eight parts of Ayurveda in prose and verse forms. The first two pillars of Ayurveda describe the antirheumatic (antiarthritis) activity of gugguls-the gum-resins of trees[16–20]. In addition to its beneficial use for arthritis, this gummy resin is also mentioned in traditional Ayurvedic and Unani texts as an effective remedy for diarrhoea, dysentery, ringworm, boils, fevers (antipyretic), skin and blood diseases, cardiovascular diseases, mouth sores, bad throat, bronchitis, asthma, cough, vaginal discharges, hair-loss, jaundice, hemorrhoids, syphilitic diseases, irregular menses and stimulation of liver. It is also diaphoretic, astringent, diuretic and acts both as internal and external stimulant. Modern medicine and pharmacology strongly point out to its use as an antiarthritic, antiinflammatory, antihyperlipidemic (controls blood lipids), antiatherosclerotic (anticoronary plaque), analgesic (pain-reliever) and hepatoprotective (protects the liver)[15,21–24]. Animal models and *in vitro* studies had shown that boswellic acids inhibit the synthesis of pro-inflammatory enzyme, which cause bronchoconstriction, chemo taxis, and increased vascular permeability [33–38]. Other anti-inflammatory drug like quercetin, also block this enzyme but in other fashion through its anti-oxidant activity whereas boswellic acids seem to be specific inhibitor of enzyme [39]. The enzyme causes inflammation by stimulating the p free radical damage, cell-adhesion, calcium displacement, and migration of inflammation-producing cells to the inflamed body area. But the boswellic acids have been shown to significantly reduce glycosaminoglycan degradation [40–43]. In other study it was found that the effect of *Boswellia* acid extract and ketoprofen occurs in deferent way [44].

In vitro studies it was recorded that boswellic acids were found to inhibit leukotriene synthesis via 5-LO, *Boswellia* acids has shown to be specific, non-redox inhibitors of leukotriene synthesis, either interacting directly with the enzyme or blocking it [45,46]. *Boswellia* acids was found to play an important role in chronic bronchitis cystic fibrosis, and acute respiratory distress syndrome [47,48]. It was found that 3-acetyl-11-keto- β -boswellic acid (AKBA) is the most potent inhibitor of 5-LO, an enzyme responsible for inflammation[49][50,51].[52,53]. The boswellic acid from *Boswellia serrata*, when tested on new model i.e. Papaya Latex Model, showed significant

activity of mean 35% inhibition of inflammation. Since the new model is reported to be sensitive to slowly acting remission-inducing drugs, its effectiveness on boswellic acid throws some light on its mechanism of action, which seems to be unlike aspirin and steroidal drugs[54]. Poeckel and Werz in 2006 have summarized the biological actions of boswellic acids on the cellular and molecular level and attempted to put the data into the perspectives of the beneficial effects manifested in animal studies and trials with human subjects related to inflammation and cancer[55]. Sharma *et al.*[56] have reported the effect of boswellic acids on bovine serum albumin (BSA)-induced arthritis in rabbits. Gayathri *et al.*[57] in 2007 have reported that pure compound from *Boswellia serrata* extract exhibits anti-inflammatory property in human peripheral blood mononuclear cells (PBMCs) and mouse macrophages through inhibition of tumor necrosis factor- α (TNF- α), interleukin-1 β (IL-1 β), NO and mitogen activated protein (MAP) kinases. Incensole acetate, a novel anti-inflammatory compound isolated from *Boswellia* resin inhibits nuclear factor-kappa B activation[58]. Boswellic acids are direct 5-LO inhibitors that efficiently suppress 5-LO product synthesis in common *in vitro* test models. However, the pharmacological relevance of such interference *in vivo* seems questionable [59]. Acetyl-11-keto- β -boswellic acid inhibits prostate tumor growth by suppressing vascular endothelial growth factor receptor 2-mediated angiogenesis [60]. Very recently, Pawaret *et al.* in 2011 have reported a simple, rapid, accurate, reproducible, selective and economic HPTLC method for routine quality control analysis as also quantitative determination of β -boswellic acid from *Boswellia serrata* Roxb. (exudate) and its formulations[62].

CONCLUSION:

Boswellic acids (triterpenoids), which represents a method of validating the authenticity of the essential oil. The chemistry of the essential oil is mainly monoterpenes and sesquiterpenes, with small amounts of diterpenoid components being the upper limit in terms of molecular weight. Analysis of frankincense from various *Boswellia* species with inhibitory activity on human drug metabolising cytochrome P450 enzymes. Boswellic acids was found to play an important role in chronic bronchitis cystic fibrosis, and acute respiratory distress syndrome [47,48]. It was found that 3-acetyl-11-keto- β -boswellic acid (AKBA) is the most potent inhibitor of 5-LO, an enzyme responsible for inflammation

REFERENCES:

- Vol. 1. New Delhi: Council of Scientific and Industrial Research; 1948. Anon (CSIR). The Wealth of India, Raw Materials.
- Felter H, Lloyd J. I and II. Cincinnati: Ohio Valley Company, US; 1898. Kings American Dispensatory.
- Howes FN. Age-old resins of the Mediterranean region and their uses. *Econ Bot.* 1950;4:307-16.
- Krishnamurthy T, Shiva MP. Salai guggul (from *Boswellia serrata* Roxb.): Its exploitation and utilization. *Indian Forest.* 1977;103:466-74.
- Verghese J. Olibanum in focus. *Perfumer Flavorist.* 1988;13:2-11.
- Holmes P. Frankincense oil. *Int J Arom.* 1999;9:56-61.
- Maupetit P. New constituents in olibanum resinoid and essential oils. *Perfumer Flavorist.* 1984;9:19-37.
- Leung AY, Foster S. 2nd ed. New York: John Wiley and Sons; 1996. Encyclopedia of common natural ingredients used in food, drugs and cosmetics; pp. 389-91.
- Wallis TE. 5th ed. London: J and A Churchill Limited; 1967. Textbook of Pharmacognosy; pp. 500-1.
- Evans WC. 14th ed. London: WB Saunders Company Ltd; 1996. Trease and Evans Pharmacognosy; p. 289.
- Chiavari G, Galletti GC, Piccaglia R, Mohamud MA. Differentiation between resins from *Boswellia carterii* and *Boswellia frereana* (Frankincense) of Somali region. *J Essential Oil Res.* 1991;3:185-6.
- Sharma RA, Verma KC. Studies on gum obtained from *Boswellia serrata* Roxb. *Indian Drugs.* 1980;17:225.
- Bhuchar VM, Agarwal AK, Sharma SK. Constituents of gum obtained from *Boswellia serrata* exudates. *Indian J Technol.* 1982;20:38.
- Gangwal ML, Vardhan DK. Carbohydrate contents of *Boswellia serrata*. *Asian J Chem.* 1995;7:677.
- Sharma S, Thawani V, Hingorani L. Pharmacokinetic study of 11-keto-beta-boswellic acid. *Phytomedicine.* 2004;11:255-60.
- Monograph *Boswellia serrata*. *Altern Med Rev.* 1998;3:306-7.
- Kirtikar KR, Basu BD. The antiinflammatory action of Indian medicinal plants. *Indian Med Plants.* 1935;1:521-9.
- Chatterjee GK, Pal SD. Antiinflammatory agents from Indian medicinal plants. *Indian Drugs.* 1984;21:431.
- Khare CP. 2004. Encyclopedia of India, Rational Western Therapy, Ayurvedic and other Traditional Usage.
- New Delhi: Ministry of Health and Family Welfare; 2007. Pharmacopoeia of India, Govt. of India; p. 2045.
- Dhiman AK. Delhi: Daya Publishing House; 2006. Ayurvedic Drug Plants; pp. 326-7.
- Hostanska K, Daum G, Saller R. Cytostatic and apoptosis-inducing activity of boswellic acids toward malignant cell lines *in vitro*. *Anticancer Res.* 2002;22:2853-62.
- Lemenih M, Teketay D. Frankincense and myrrh resources of Ethiopia: II. Medicinal and industrial uses. *Ethiopian J Sci.* 2003;26:161-72.
- Mathe C, Culioli G, Archier P. Characterization of archeological frankincense by gas chromatography mass spectrometry. *J Chromatogr.* 2004;1023:277-85.
- El-Khadem H, El-Shafei ZM, Elsekeify MA, Abdel Rahman MM. Derivatives of boswellic acids. *Planta Med.* 1972;22:157-9.
- Pardhy RS, Bhattacharyya SC. Tetracyclic triterpene acids from the resin of *Boswellia serrata* Roxb. *Indian J Chem.* 1978;16B:174-5.
- Pardhy RS, Bhattacharyya SC. β -Boswellic acid, acetyl- β -boswellic acid, acetyl-11-keto- β -boswellic acid and 11-keto- β -boswellic acid, four pentacyclic triterpene acids from the resin of *Boswellia serrata* Roxb. *Indian J Chem.* 1978;16B:176-8.
- Mahajan B, Taneja SC, Sethi VK, Dhar KL. Two triterpenoids from *Boswellia serrata* gum resin. *Phytochemistry.* 1995;39:453-5.
- Handa SS. Herbal raw material and traditional remedies. *Eastern Pharmacist.* 1995;3:24.
- Choudhary AC, Dikshit SK. Ayurvedic Pharmacopoeia. *Eastern Pharmacist.* 1999;10:52.
- Sane RT. Standardization, quality control, and GMP for herbal drug. *Indian Drugs.* 2002;39:184-90.
- Safayhi H, Mack T, Sabieraj J, Anazodo MI, Subramanian LR, Ammon HP. Boswellic acids: Novel, specific, non-redox inhibitors of 5-lipoxygenase. *J Pharmacol Exp Ther.* 1992;261:1143-6.
- Ammon HP, Mack T, Singh GB, Safayhi H. Inhibition of leukotriene B₄ formation in rat peritoneal neutrophils by an ethanolic extract of gum-resin exudates of *Boswellia serrata*. *Planta Med.* 1991;57:203-7.
- Wildfeuer A, Neu IS, Safayhi H, Metzger G, Wehrmann M, Vogel U, et al. Effects of boswellic acids extracted from a herbal medicine on the biosynthesis of leukotrienes and the course of experimental autoimmune encephalomyelitis. *Arzneim Forsch.* 1998;48:668-74.
- Ammon HP. Boswellic acids (components of frankincense) as the active principle in treatment of chronic inflammatory diseases. *Wien Med Wochenschr.* 2002;152:337-78.
- Ammon HP. Boswellic acids in chronic inflammatory diseases. *Planta Med.* 2006;72:1100-16.
- Schweizer S, von Brocke AF, Boden SE, Bayer E, Ammon HP, Safayhi H. Workup-dependent formation of 5-lipoxygenase inhibitory boswellic acids analogues. *J Nat Prod.* 2000;63:1058-61.
- Etzel R. Special extract of *boswellia serrata* (H15) in the treatment of rheumatoid arthritis. *Phytomedicine.* 1996;3:91-4.
- Ammon HP. Salai guggul-*Boswellia serrata* from a herbal medicine to a specific inhibitor of leukotriene biosynthesis. *Phytomedicine.* 1996;3:67-70.

40. Lee KH, Spencer MR. Studies on mechanism of action of salicylates V: Effect of salicylic acid on enzymes involved in mucopolysaccharide synthesis. *J Pharmacol Sci.* 1969;58:464–8.
41. Palmowski MJ, Brandt KD. Effect of salicylate on proteoglycan metabolism in normal canine articular cartilage *in vitro*. *Arthritis Rheum.* 1979;22:746–54.
42. Dekel S, Falconer J, Francis MJ. The effect of anti-inflammatory drugs on glycosaminoglycan sulphation in pig cartilage. *Prostaglandins Med.* 1980;4:133–40.
43. Brandt KD, Palmowski MJ. Effect of salicylates and other non-steroidal anti-inflammatory drugs on articular cartilage. *Am J Med.* 1984;77:65–9.
44. Reddy GK, Chandrakan G, Dhar SC. Studies on the metabolism of glycosaminoglycans under the influence of new herbal anti-inflammatory agents. *Biochem Pharm.* 1989;38:3527–34.
45. Ammon HP, Safayhi H, Mack T, Sabieraj J. Mechanism of anti-inflammatory actions of curcumin and boswellic acids. *J Ethnopharmacol.* 1993;38:113–9.
46. Safayhi H, Sailer ER, Ammon HP. Mechanism of 5-lipoxygenase inhibition by acetyl-11-keto-boswellic acid. *Mol Pharmacol.* 1995;47:1212–6.
47. Rall B, Ammon HP, Safayhi H. Boswellic acids and protease activities. *Phytomed.* 1996;3:75–6.
48. Safayhi H, Rall B, Sailer ER, Ammon HP. Inhibition by boswellic acids of human leucocyte elastase. *J Pharmacol Exp Ther.* 1997;281:460–3.
49. Singh GB, Atal CK. Pharmacology of an extract of salai guggul ex- *Boswellia serrata*. *Indian J Pharmacol.* 1984;16:51.
50. Kulkarni R, Patki P, Jog V, Gandage S, Patwardhan B. Treatment of osteoarthritis with a herbomineral formulation: A double blind, placebo-controlled, cross-over study. *J Ethnopharmacol.* 1991;33:91–5.
51. Chopra A, Lavin P, Patwardhan B, Chitre D. Randomized double blind trial of an Ayurvedic plant derived formulation for treatment of rheumatoid arthritis. *J Rheumatol.* 2000;27:1365–72.
52. Murray MT. Rocklin, CA: Prima Publishing; 1995. *The Healing Power of Herbs*; pp. 327–35.
53. Arora RB, Kapoor V, Basu N, Jain AP. Anti-inflammatory studies on *Curcuma longa* (turmeric) *Indian J Med Res.* 1971;50:1289–95.
54. Gupta OP, Sharma N, Chand D. A sensitive and relevant model for evaluating antiinflammatory activity – papaya latex-induced rat paw edema. *J Pharmacol Toxicol Methods.* 1992;28:15–9.
55. Poeckel D, Werz O. Boswellic acids: Biological actions and molecular targets. *Curr Med Chem.* 2006;13:3359–69.
56. Sharma ML, Bani S, Singh GB. Anti-arthritis activity of boswellic acids in bovine serum albumin (BSA)-induced arthritis. *Int Immunopharmacol.* 1989;11:647–52.
57. Gayathri B, Manjula N, Vinaykumar KS, Lakshmi BS, Balakrishnan A. Pure compound from *Boswellia serrata* extract exhibits antiinflammatory property in human PBMCs and mouse macrophages through inhibition of TNF alpha, IL-1beta, NO and MAP kinases. *Int Immunopharmacol.* 2007;7:473–82.
58. Moussaieff A, Shohami E, Kashman Y, Fride E, Schmitz ML, Renner F, et al. Incensole acetate, a novel anti-inflammatory compound isolated from *Boswellia* resin inhibits nuclear factor-kappa B activation. *Mol Pharmacol.* 2007;72:1657–64.
59. Siemoneit U, Pergola C, Jazzar B, Northoff H, Skarke C, Jauch J, et al. On the interference of boswellic acids with 5-lipoxygenase: Mechanistic studies *in vitro* and pharmacological relevance. *Eur J Pharmacol.* 2009;606:246–54.
60. Pang X, Yi Z, Zhang X, Sung B, Qu W, Lian X, et al. Acetyl-11-keto- β -boswellic acid inhibits prostate tumor growth by suppressing vascular endothelial growth factor receptor 2-mediated angiogenesis. *Cancer Res.* 2009;69:5893–900.
61. Anonymous. Indian herb hope for arthritis relief. *The Telegraph Calcutta.* 2008. Aug 4, [Last accessed on 2011 May 18]. p. 7.
62. Pawar RK, Sharma S, Singh KC, Sharma RK. Physico-chemical standardization and development of HPTLC method for the determination of β -boswellic acid from *Boswellia serrata* Roxb.(exudate) *Int J App Pharm.* 2011;3:8–13.
63. Basch E, Boon H, Davies T, Hashmi S, Hasskari J, Sollars D, et al. *Boswellia*: A evidence based systematic review by the natural standard research collaboration. *J Herb Phar.* 2004;4:63–83.
64. New York: Churchill Livingstone; 2002. *Dabur Research Foundation and Dabur Ayurved Limited. Major herbs of Ayurveda.*