

PHYTOCHEMISTRY AND PHARMACOLOGICAL ACTIVITIES OF *BERSAMA ENGLERIANA* GUERKE- AN OVERVIEW

Lather Amit^{1,2*}, Gupta Vikas¹, Tyagi Vaibhav², Kumar Vikash², Garg Siddhartha²

¹University Centre of Excellence in Research, BFUHS, Faridkot, India

²School of Pharmaceutical Sciences, Shobhit University, Meerut, India

*Amit Lather, School of Pharmaceutical Sciences, Shobhit University, Meerut, India

Email: amitlather244@yahoo.com

Article Received on: 14/11/10 Revised on: 30/11/10 Approved for publication: 09/12/10

ABSTRACT

Bersama engleriana Guerke (Melianthaceae) is the synonym of the *Bersama abyssinica*. Bark, leaf and root decoctions are widely taken as a purgative to treat a range of stomach disorders, such as abdominal pain, colic, diarrhea, cholera, intestinal worms, amoebiasis and dysentery. This plant mainly contains mangiferin, swinniol, Δ^4 -stigmaster-3 β -ol, 4-methylstigmaster-5, 23-dien-3 β -ol, sterols, flavones, phenols, triterpenes, anthraquinones, Hellebrigenin 3-acetate, hellebrigenin 3, 5-diacetate and bersaldegenin 1, 3, 5-orthoacetate. *Bersama engleriana* shows hypoglycemic, antimicrobial, antioxidant, antitumor, anti HIV activities. This review explains various traditional uses as well as phytochemical and pharmacological reports on *Bersama engleriana* which help the researcher to explore more potential of this plant.

KEYWORDS: *Bersama*, Purgative, Decoctions, Amoebiasis, Dysentery.

INTRODUCTION

During the last two decades, traditional systems of medicine and medicinal plant research have become subject of worldwide interest and consequence. In many developing nations of the world, large numbers of people still rely heavily on traditional healers and medicinal plants to meet their daily primary healthcare needs. Because of their perceived efficiency, negligible side effects in clinical experience and moderately low cost, herbal drugs are prescribed widely even when their biologically active compounds are unknown¹. In Upaveda, Ayurveda composed around 2500 BC, deals with medicine, healthcare and treatment of disease from indigenous drugs. From Veda it is learnt that Indo-Aryans used the 'Soma' (A Plant product) as a medicinal agent, which exhibits an amazing stimulating effect². Guerke belongs to Melianthaceae family. This plant is mainly found in Cameroon³. *Bersama abyssinica* is a poisonous and have been implicated in killing human and livestock. *Bersama abyssinica* (*Bersama engleriana*) is distributed from Guinea Bissau through the coastal countries of West Africa except Benin, east to Eritrea and Ethiopia and south to Angola, Zambia, Zimbabwe and Mozambique. Evergreen shrub to small tree up to 12(-25) m tall; bark grey, brown or mottled, scaly. *Bersama abyssinica* grows in lowland bush savanna, gallery forests and montane forests, from sea-level up to 2700 m altitude. *Bersama abyssinica* can be propagated by seed, cuttings, wildlings or root suckers⁴. Fruits are red woody capsules⁵. In Babungu (Cameroon) *Bersama abyssinica* is known as Fuaveti and Timber is used for construction purposes⁶.

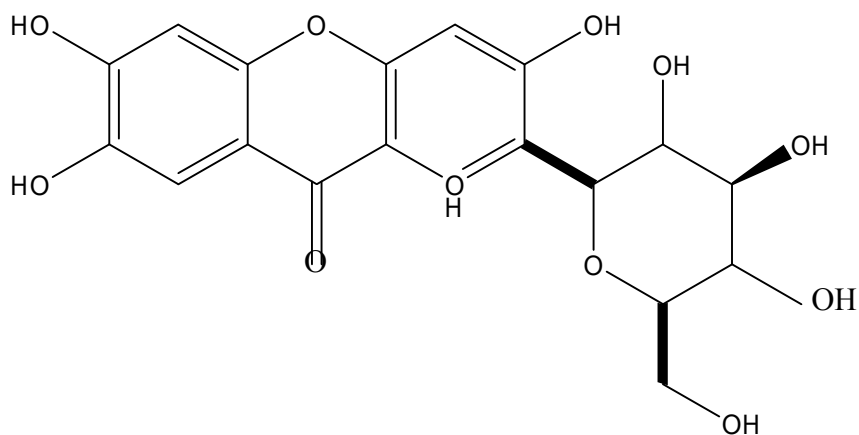
Traditional Uses

Bersama abyssinica is traditionally used to treat rheumatism⁷. *Bersama abyssinica* stem bark extract has been used as an aphrodisiac^{8,9}. Its leaves are also used to treat snake bite¹⁰. Bark, leaf and root decoctions are widely taken as a purgative to treat a range of stomach disorders, such as abdominal pain, colic, diarrhea, cholera, intestinal worms, amoebiasis and dysentery. Rabies, syphilis and gonorrhea are

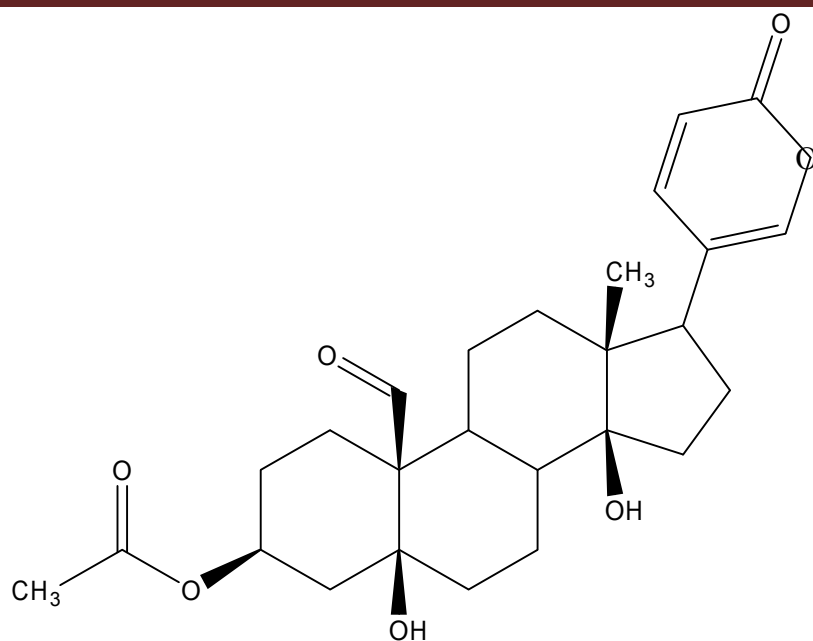
also treated with these decoctions. A stem bark decoction is drunk to cure cancer and rheumatism. As an aphrodisiac, powdered bark is added to beer or leaves are chewed. Stem bark and leaves are used to treat diabetes mellitus. Leaf decoctions are also taken to treat feverish pains, loss of appetite, debility, jaundice and leprosy. Extracts of growing shoots are used for external treatment of burns, ulcers and to clean wounds⁴. In Kenya boiled root and leaves of the plant is traditionally used in the treatment of general sexually transmitted diseases (STD)^{11, 12}.

Chemical Constituents

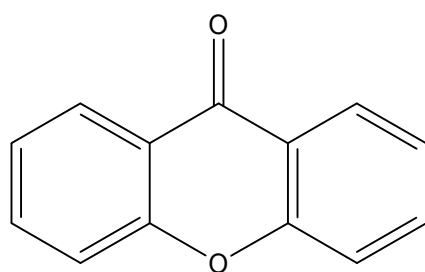
Fractionation of the CH₂Cl₂-MeOH (1:1) extract of the stem bark of *B. engleriana* afforded a xanthone C-glucoside (mangiferin) and isolation of three terpenoids swinniol, Δ^4 -stigmaster-3b-ol and 4-methylstigmaster-5, 23-dien-3-b-ol³. *Bersama engleriana* contains sugar moieties Glc (1-2) GlcA at C-3, Glc at C-28 and an aglycone moiety¹³⁻¹⁵. Phytochemical analysis revealed the presence of sterols, flavones and alkaloids in *B. engleriana* leaves¹⁶. The phytochemical tests indicated the presence of flavonoids, phenols; triterpenes and anthraquinones in methanol extract¹⁷. Hellebrigenin 3-acetate, hellebrigenin 3, 5-diacetate and bersaldegenin 1, 3, 5-orthoacetate were also isolated from *Bersama engleriana* plant¹⁸⁻²⁰. Five 3-O-glucuronide triterpene saponins were isolated from the stem bark of *Bersama engleriana* Gurke along with two known saponins, polyscias saponin C and aralia saponin 15, and one major C-glycoside xanthone, mangiferin²¹. Stem bark also contains a mixture of cardenolides including abyssinol A, B, C, bersaldegenin, hellebrigenin, and bufadienolide-O-acetate, as well as saponins, mangiferine, and gallic acid derivatives^{8, 9}. Stem bark of the plant contains oleanolic acid saponins^{22, 23}.



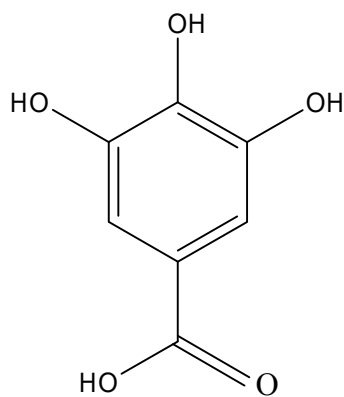
Mangiferin



Hellebrigenin 3-acetate



Xanthone



Gallic acid

Pharmacological Activities

Hypoglycemic Activity

The hypoglycemic properties of the aqueous and methanol extracts of the leaves of *Bersama engleriana* were evaluated in normoglycaemic rats in order to scientifically validate its traditional therapeutic use. With a dose of 300 mg/kg b.w. only the aqueous extract appeared to be significantly effective while at high dose, 600 mg/kg b.w., the aqueous and methanol extracts of *Bersama engleriana* reduced the blood glucose level. These results confirm the hypoglycemic properties of some extracts of the leaves of *Bersama engleriana* with the aqueous extract appearing more active²⁴. Bioactive molecules present in these extracts of *Bersama engleriana* may probably possess an insulin-like effect or stimulate the pancreatic β cells to produce insulin which in turn lowers the blood glucose level. Similar observations have been reported by some others scientists also²⁵⁻²⁷.

Antimalarial Activity

Methanol extract from *Bersama engleriana* were highly active on schizonts. Phytochemical analysis revealed the presence of sterols, flavones and alkaloids in *B. engleriana* leaves. One fraction and one pure product from *B. engleriana* leaves were highly active on the field parasites. *B. engleriana* was the least toxic with lower cytotoxicity than chloroquine. No acute toxicity was observed in 6N C57 BL mice for the pure products and highly active extracts^{16, 28, 29}.

Antitumor, antioxidant and antimicrobial activities

Methanolic extracts from the roots, stem bark, leaves and wood of *Bersama engleriana* showed antitumor, antioxidant and antimicrobial activities. The crown gall tumor and DPPH radical scavenging assays were used to detect respectively the antitumor and oxidant activities. Agar diffusion and liquid dilution were used for antimicrobial tests and the phytochemical assays were conducted according to Harbone methods. The single-dose oral toxicity test was performed in accordance with the OPPTS 870.1100 and OECD 401 guidelines. Pronounced tumor reducing activity was observed with the extracts from the roots and leaves. The DPPH scavenging activity showed that the extract from the leaves was the most active. The results of antimicrobial activity showed that all tested extracts were active against all tested microbial species, including Gram-positive and negative bacteria, the two *Candida* species and mycobacterium. Under the conditions of the studied toxicity, all extracts were found to be non-toxic¹⁷⁻²⁰. Hellebrigenin is the chemical constituent responsible for anticancer activity of the plant³⁰. Ethanol extract, water extract and methanolic extract of Stem bark and root bark *Bersama abyssinica* also show antibacterial activity against various bacterial strains at different concentration e.g. *Bacillus cereus*, *Neisseria gonorrhoea*, *Streptococcus pneumonia* etc³¹.

Anti HIV activity

Root bark extract of *Bersama abyssinica* displayed antiviral activity at concentrations that were nontoxic to MT-4 cells. *Bersama abyssinica* displayed anti-HIV activity against both HIV-1 and HIV-2 strains^{32, 33}.

Excopula Ejaculatory activity

Fictive ejaculation of *Bersama engleriana* is studied in spinal male rats. Anesthetized rats treated intravenously with aqueous and methanolic extracts from the dried leaves of *B. engleriana* in the absence and presence of dopamine or oxytocin. The inhibitory effect of *B. engleriana* extracts on the expression of fictive ejaculation in spinal male rat is mediated through dopaminergic and oxytocinergic pathways. This prolonged ejaculatory latency caused by *B. engleriana* could support its potential use in patients with rapid ejaculation³⁴.

CONCLUSION

Major thrust by whole of the pharmaceutical industry is paying attention towards design and development of new novel and indigenous plant based drugs through investigation of leads from traditional system of medicine³⁵. In recent years, ethno-botanical and traditional uses of natural compounds, especially of plant origin received much attention as they are well tested for their efficacy and generally believed to be safe for human use. The health care systems are going to become more and more expensive therefore, we have to develop technologies to essentially introduce and integrate herbal medicine system in our health care. The present review shows the pharmacological properties of various bioactive compounds present in *Bersama engleriana*.

REFERENCES

1. Valiathan MS. Healing Plants. Current Science 1998; 75:1122-1126.
2. Satyaavati GV, Raina MK, Sharma M. Medical plants of India. New Delhi: Indian Council of Medical Research. 1976. p. 76-77.
3. Djemgou PC, Hussien TA, Hegazy MEF, Ngandeu F, Neguim G, Tane P et al. C-Glucoside xanthone from the stem bark extract of *Bersama engleriana*. Phcog Res. 2010; 2:229-232.
4. Bosch CH. *Bersama abyssinica* Fresen. In: Schmelzer GH, Gurib-Fakim A (Editors). Plant Resources of Tropical Africa 11(1). Medicinal plants 1. PROTA Foundation. Wageningen, Netherlands: Backhuys Publishers, 2008. p.116.
5. Timberlake J, Golding J, Clarke P. Niassa Botanical Expedition June 2003. Occasional Publications in Biodiversity No. 12. Biodiversity Foundation for Africa. Fomona, Bulawayo, Zimbabwe, 2003. p.1-32.
6. Simbo DJ. An ethnobotanical survey of medicinal plants in Babungu, Northwest Region, Cameroon. Simbo Journal of Ethnobiology and Ethnomedicine 2010; 6(8):1-7.

7. Yineger H, Yewhalaw D, Teketay D. Ethnomedicinal plant knowledge and practice of the Oromo ethnic group in southwestern Ethiopia. *Journal of Ethnobiology and Ethnomedicine* 2008; 4(11):1-10.
8. Iwu MM. *Handbook of African Medicinal Plants*. CRC Press 1993.
9. Dweck AC. Impotence – evaluating traditional remedies. *Personal Care* 2007:5.
10. Tadesse M, Hunde, Getachew Y. Survey of Medicinal Plants Used To Treat Human Diseases In Seka Chekorsa, Jimma Zone, Ethiopia. *Ethiop J Health Sci* 2005; 15(2):89-106.
11. Njoroge GN, Bussmann RW. Ethanotherapeutic management of Sexually Transmitted Diseases (STDs) and reproductive health conditions in central province of Kenya. *Indian Journal of Traditional Knowledge* 2009; 8(2):255-261.
12. Jiofack T, Fokunang C, Guedje N, Kemeuze V, Fongnzossie E, Nkongmeneck BA. Ethnobotanical uses of medicinal plants of two ethnoecological regions of Cameroon. *International Journal of Medicine and Medical Sciences* 2010; 2(3):60-79.
13. Gauthier C, Legault J, Pichette A. Recent Progress in the Synthesis of Naturally Occurring Triterpenoid Saponins. *Mini-Reviews in Organic Chemistry* 2009; 6:321-344.
14. Tapondjou AL, Miyamoto T, Lacaille-Dubois MA. Glucuronide triterpene saponins from *Bersama engleriana*. *Phytochemistry* 2006; 67:2126-2132.
15. Jiagang D. Study on Mango Leaf and Mangiferin. 1st International symposium on screening functional components of agricultural residues and the study on Mangiferin. Baise, Guangxi, China 2009. p.63.
16. African Network for Drugs and Diagnostics Innovation (ANDI), Creating a sustainable platform for R&D innovation in Africa, Part 1: Health Product R&D Landscape in Africa, Part 2: Collection of Meeting Abstracts. WHO Founding meeting in Abuja, Nigeria, 6–8 October 2008. p.70.
17. Victor K, Armelle TM, Maurice T, Veronique PB, Francois-Xavier ETOA, Augustin EN et al. Antitumor, antioxidant and antimicrobial activities of *Bersama engleriana* (Melianthaceae). *Journal of Ethnopharmacology* 2008; 115(3):494-501.
18. Kupchan SM, Moniot JL, Sigel CW, Hemingway RJ. Tumor inhibitors. LXV, Bersenogenin, berscillogenin, and 3-epiberscillogenin, three new cytotoxic bufadienolides from *Bersama abyssinica*. *J. Org. Chem.* 1971; 36(18):2611–2616.
19. Kupchan M, Hemingway RJ, Hemingway JC. *Tetrahedron Letters* 149 (1968); 1. *Org. Chem.* 1969; 34:3894.
20. Kupchan SM, Ognyanov I. *Tetrahedron Letters* 1969. p.1709.
21. Tapondjou AL, Miyamoto T, Lacaille-Dubois MA. Glucuronide triterpene saponins from *Bersama engleriana*. *Phytochemistry*. 2006; 67(19):2126–2132.
22. Fai YM. An update of review on the presence of Oleanolic acid in Natural Products at Aug 2010. *Natura Proda Medica* 2010; (3):13-73.
23. Fai YM, Tao CC. A review of presence of Oleanolic acid in Natural Products. *Sample Review for Natura Proda Medical* 2009, p.1-271.
24. Njike GN, Watcho P, Nguetefack TB, Kamanyi A. Hypoglycaemic activity of the leaves extracts of *Bersama engleriana* in rats. *African Journal of Traditional. Complementary and Alternative medicines* 2005; 2(3):215-221.
25. Fuentes O, Arancibia-Avila P, Alarcon J. Hypoglycemic activity of *Bauhinia candicans* in diabetic induced rabbits. *Fitoterapia* 2004; 75(6):527-532.
26. Hemalatha S, Wahi AK, Singh PN, Chansouria JP. Hypoglycemic activity of *Withania coagulans* Dunal in streptozotocin-induced diabetic rats. *J Ethnopharmacol* 2004; 93:261-264.
27. Sepici A, Gurbuz I, Cevik C, Yesilada E. Hypoglycaemic effects of myrtle oil in normal and alloxan-diabetic rabbits. *J Ethnopharmacol* 2004; 93:311-318.
28. Ngemenya M, Titanji V, Akam T, Yong J, Tane P, Fanso-Free S et al. Antiplasmodial activity and toxicity of extracts and products from selected medicinal plants used in Cameroon. *Abstract/Acta Tropica* 2005; 96S:S1–S506.
29. Zofou D, Ngemenya MN, Tane P, Titanji VPK. Malaria drug discovery from Cameroonian medicinal plants: Screening efforts at the Biotechnology Unit, University of Buea, In Cameroon 3rd Stakeholders

- Meeting of the African Network for Drugs and Diagnostics Innovation. United Nations Office in Nairobi (UNON), Kenya 2010. p.137.
30. Patel B, Das S, Prakash R, Yasir M. Natural Bioactive Compound with Anticancer Potential. International Journal of Advances in Pharmaceutical Sciences 2010; 1:32-41.
 31. Geyid A, Abebe D, Debella A, Makonnen Z, Aberra F, Teka F et al. Screening of some medicinal plants of Ethiopia for their anti-microbial properties and chemical profiles. Journal of Ethnopharmacology 2005; 97:421–427.
 32. Moshi MJ. Current and future prospects of integrating traditional and alternative medicine in the management of diseases in Tanzania. Tanzania Health Research Bulletin 2005; 7:159-167.
 33. Asres K, Bucar F, Kartinig T, Witvrouw M, Pannecoupue C, De Clercq E. Antiviral activity against human immunodeficiency virus type 1 (HIV-1) and type 2 (HIV-2) of ethno botanically selected Ethiopian medicinal plants. Phytotherapy Research 2001; 15:62-69.
 34. Watcho P, Carro-Juarez M. Evaluation of the excopula ejaculatory potentials of *Bersama engleriana* in spinal male rats. Asian Journal of Andrology 2009; 11:533–539.
 35. Patwardhan B, Ashok DBV, Chorghade M. Ayurveda and natural products drug discovery. Current Science 2004; 80(6):789-799.