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Medicinal plant: Garcinia spp.

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Abstract— Garcinia is a tropical fruit tree with promising pharmacological properties. This review presents an overview of the bioactive compounds derivative from Garcinia fruits and their biological activities for promoting human health as food and medicine.

Keywords—Medicinal plant, Garcinia spp., Therapeutic properties, phytochemical properties.

I. INTRODUCTION

Plants are considered as the great reservoir of structurally diverse bioactive molecules such as phenolics, terpenoids, carotenoids, anthocyanins and flavonoids which are having therapeutic values and are useful in the treatment of various ailments. Now-a-days these bioactive molecules are widely used in the food, pharmaceutical and cosmetics industries (Hosakatte *et al.*, 2018).

Garcinia is a polygamous tropical tree or shrub under Clusiaceae family. It consists of 250 species, out of which about 30 species are indigenous to India. *Garcinia pedunculata* (Amlavethasa), *G. cowa*, and *G.Morella* (Indian gamboge) are grown in North-Eastern parts of India and Andaman Islands (Negi *et al.*,2008; Sharma and Devi,2015; Murthy *et al.*, 2020). *Garcinia* are rich source of nutrients, minerals, vitamins, and dietary fibers. It has the folklore claims such as rejuvenator, cardio tonic, asthma, obesity and arthritis. The mature fruit is eaten cooked or raw and also for pickle preparation.

Garcinia pedunculata is an evergreen tree. The tree is endemic to the south eastern regions of Asia such as parts of Myanmar and North-Eastern parts of India. The tree has a fluted trunk with short spreading branches. Leaves are lanceolate with prominent mid ribs. Male flowers are light green in sparsely flowered panicles, the female flowers are solitary. The fruit is round with a diameter ranging between 8cm and 12cm. It has a juicy interior with edible arils. The mature *G. pedunculata* fruit is greenish yellow and is consumed as a vegetable.

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II. MEDICINAL PROPERTIES OF GARCINIA

The fruits of *Garcinia* have been used since ancient times in traditional medicinal practices. These species provide a rich natural source of bioactive compounds with relevant therapeutic properties and anti-inflammatory effects, for the treatment of skin disorders, wounds, ulcers, dysentery, pain, infections, fever, cough, bronchitis, asthma, rheumatoid arthritis, obesity and having antioxidant, antiaflatoxigenic anti-inflammatory, leishmanicidal, and antiprotozoal activities (Joseph *et al.*,2005; Ali *et al.*,2017 ;Espirito *et al.*,2020).

III. PHYTOCHEMICAL PROPERTIES OF SOME SPECIES OF GARCINIA

Garcinia are rich sources of fiber, total phenols, and natural antioxidants with high amount of ascorbic acids .Extracts of the pericarp, epicarp, and seeds of Garcinia have demonstrated the phytochemicals such as pedunculol, garcinol, cambogin 3 and hydroxyl citric acid. Bennet and Lee (1989) and Rao et al.(1974;1980) have isolated the bioactive compounds namely benzoquinones, triterpenes and anthocyanins. Garcinol, being rich in derivates of poly-isoprenylated benzophenones, polyphenols, bioflavonoids (kolaviron, volkensiflavone, fukugetin) and xanthones (Sarma et al., 2016). Xanthones are the major class of phenolic compounds in Garcinia species, followed by benzophenones and biflavonoids. Xanthones have demonstrated effects against human cervical cancer, lung cancer cells, and hepatocellular carcinomas (Vo et al., 2015). These compounds have been associated with biological activities such as free-radical scavenging,

antiulcer effects, cytotoxicity, inhibition of nitric oxide synthase, chemoprevention of cancer, induction of apoptosis, anti-HIV, and trypanocidal effects (Hung et al.,2015; Fu et al.,2017). These compounds exhibiting a wide range of pharmacological activities such as antimicrobial, antioxidant, antitumour-promoting, cytotoxic, etc. (Jayaprakasha et al., 2006; Mundugaru et al., 2014; Adegoke et al., 1998; Asano et al., 1996; Bakana et al., 1987; Iinuma et al., 1996; Mackeen et al., 2000; Fu et al., 2014; Minami et al., 1994; 1995; 1996; Islam et al.,2015 ;Paul et al.,2017).Crude extracts as well as partially purified compounds from different parts of some species of Garcinia plants have shown potential antibacterial activities against Bacillus cereus, Bacillus coagulans, Bacillus subtilis, Staphylococcus aureus and Escherichia coli.

Some prominent *Garcinia* **species** are known to have good medicinal value and fruit extract has traditionally very well known for treatments of various diseases (Deore *et al.*, 2011).

G. pedunculata was estimated to contain moisture 88.20%, protein 0.50%, β carotene 45.00mg/100 g, vitamins 0.03, riboflavin 0.02, (thiamine and ascorbic acid142.83mg/100 g, resp.), minerals (sodium 1.80, potassium106.00, calcium 18.00, magnesium 23.00, iron 0.08, zinc0.15, copper 0.12, and phosphorus 17.00mg/100 g), phenolics(19.45mg gallic acid/100 g), and flavonoids (18.33mg rutin/g). The dried fruit rinds and pericarp of G. pedunculata have been reported to contain some benzophenones, pedenculol, hydroxy citric acid, garcinol, and cambogin, some of which are strong antioxidants (Sahu et al., 1989; Mudoi et al., 2012; Ravi et al., 2014; Sarma et al., 2015; Mundugaru et al., 2019). This fruit extract is reported to possess a variety of pharmacological benefits including antimicrobial, anti-inflammatory, hepatoprotective, and cardioprotective properties (Kagyung et al., 2010; Mundugaru et al., 2014, 2016; Ali et al., 2017). The aqueous extract of Garcinia pedunculata, exhibited significant neuroprotection against AlCl₃ induced neurotoxicity (Mundugaru et al., 2016, 2017).

In traditional system of medicine the leaves of G. lancifolia are used as stomachic and diuretic. The acidic fruits are used to prepare juice, pickle and curries. G. lancifolia is used as stomachic, diuretic and its fruit is used to cure dysentery and diarrhoea. The bark of G. lanceifolia has also been reported to contain prominent antibacterial and anthelmintic potential (Chowdhury and Handique, 2012; Bora *et al.*, 2014a; 2014b). The phytochemical analysis of different extracts of G. lancifolia leaf, stem and fruit revealed the presence of tannins, saponins, flavonoids, terpenoids, alkaloids and cardiac glycosides. The high phenolic content was observed in the methanol extract of leaf followed by methanol extract of stem and dichloromethane extract of leaf.

Antimicrobial and free radical scavenging xanthones from the latex of *G. cowa* (Mahabusarakam *et al.*, 2005; Na Pattalung *et al.*,1994; Auranwiwat *et al.*,2014), and antimalarial xanthones (Likhitwitayawuid *et al.*,1998 ;) from the stem bark of *G. cowa* have been reported.

A polyisoprenylated benzophenone known as garcinol isolated from stem bark of *G. huillensis* has been shown to possess chemotherapeutical activity against Gram-positive and Gram negative cocci.

Alpha-mangostin, rubraxanthone and xanthochymol isolated from *G. mangostana*, *G. diocia* and *G. subelliptica*, respectively, showed strong antibacterial activity (Iinuma *et al.*, 1996).

Crude extracts of leaves, fruits, root, stem and trunk bark of *G. atroviridis* exhibited antibacterial (Mackeen *et al.*, 2000).

IV. INDUSTRIAL RELEVANCE

This traditional medicines are assuming greater important because of its effective, safer, locally available and no side effects and more reliable medicine than synthetically produced drugs. *Garcinia* extracts can be utilized as nutraceuticals and as food biopreservatives which could be developed into value added products or medicine (Acuna *et al.*, 2012; Biswas *et al.*, 2017). To produce potentially more active and safer drugs the plant-derived compounds should be isolated which could improve the economy of pharmaceutical industries.

V. CONCLUSION

Though the fruits of *G. cowa* and *G. pedunculata* are underutilized, recent year the interests in research activities in the fields of chemistry and pharmacology has arisen in exploiting on the fruit species. The advanced technology for isolation of the bioactive compounds from plants is very important as it could help in structural modifications of the synthetic products from the fruits. Based on the mechanism and mode of action of these plants it is confirmed the curative and therapeutic effectiveness of the plant. Hence, much research effort on crop improvement and physiologically active components is needed.

REFERENCES

- Acuna UM, Dastmalchi K,Basile MJ, Kennelly EJ.Quantitative high performance liquid chromatography photo-diode array (HPLC–PDA) analysis of benzophenones and biflavonoids in eight *Garcinia* species. J. Food. Compos. Anal. 2012; 25: 215-220
- [2] Adegoke GO, Kumar MV, Sambaiah K, Lokesh BR. Inhibitory effect of *Garcinia kola* on the lipid peroxidation in rat liver homogenate. Indian J. Exp. Biol. 1998; 36: 907-910.
- [3] Ali MY, Paul S, Tanvir EM, Hossen MS, Rumpa NN, Saha M, Bhoumik NC, Islam MA, Hossain MS, Alam N, Gan SH, Khalil MI. Antihyperglycemic, antidiabetic, and antioxidant effects of *Garcinia pedunculata* in rats. 2017;2979760. doi: 10.1155 /2017/ 2979760. Epub 2017 Oct 19.
- [4] Aravind APA, Lekshmi N, Menon, Rameshkumar KB. Structural diversity of secondary metabolites in *Garcinia* species. pp.1-18. In: (Ed.KB.Rameshkumar). Diversity of *Garcinia* species in the Western Ghats: Phytochemical Perspective Jawaharlal Nehru Tropical Botanic Garden and Research Institute, Thiruvananthapuram, 2016.
- [5] Asano J, Chiba K, Tada M, Yoshii T. Cytotoxic xanthones from *Garcinia hanburyi*. Phytochemistry. 1996; 41(3):815-20. doi: 10.1016/0031-9422(95)00682-6.
- [6] Auranwiwat C, Trisuwan K, Saiai A, Pyne SG, Ritthiwigrom T.. Antibacterial tetraoxygenated xanthones from the immature fruits of *Garcinia cowa*. Fitoterapia, 2014; 98: 179-183.
- Bakana P, Claeys M, J Totte J, Pieters LA, Van Hoof L, Tamba-Vemba, Van den Berghe DA, Vlietinck AJ. Structure and chemotherapeutical activity of a polyisoprenylated benzophenone from the stem bark of *Garcinia huillensis*. J Ethnopharmacol. 1987; 21(1):75-84. doi: 10.1016/0378-8741(87) 90096-1.
- [8] Bennet GJ,Lee HH. Xanthones from guttiferae. Phytochemistry. 1989; 28(4): 967- 998
- [9] Biswas SC,Hazarika P,Dutta NB, Sarmah H. Few novel value added products prepared from fruits of *Garcinia pedunculata*. Research Journal of Chemical Sciences. 2017; 7(3): 23-29.
- [10] Bora NS, Kakoti BB, Gogoi B.Investigation of *in-vitro* anthelmintic activity of *Garcinia lanceifolia* bark in *Pheretima posthuma* (Indian adult earthworm). *Pharmanest.* 2014a; 5(3): 2007-2010.
- [11] Bora NS, Kakoti BB, Gogoi B. 2014b. Study on Antibacterial Activity of the Bark of *Garcinia lanceifolia* Roxb. International Scholarly Research Notices. Article ID 784579, 3 pages http://dx.doi.org/10.1155/2014/784579.
- [12] ChowdhuryT, Handique PJ. Evaluation of antibacterial activity and phytochemical activity of *Garcinia lancifolia* roxb. *International Journal of Pharmaceutical Sciences&Research.* 2012; 3(6): 1663-1667.
- [13] Deore AB, Sapakal VD, Naikwade NS. Antioxidant and hepatoprotective activity of *Garcinia indica* fruit rind. Pharmacie Globale-Inter. J. Compr. Pharm. 2011; 2(6):8.
- [14] Espirito Santo BL, Santana LF, Kato Junior WH, de Araujo F, Bogo D, Freitas KC, Guimaraes RA, Hiane PA, Pott A, Oliveira Filiu WF, Asato MA, Figueiredo PO

ISSN: 2456-1878 (Int. J. Environ. Agric. Biotech.) https://dx.doi.org/10.22161/ijeab.65.5 , Bastos PR. Medicinal potential of *Garcinia* species and their compounds. Molecules. 2020;25: 4513; doi:10.3390/ molecules 25194513

- [15] Fu WW, Tan HS, Xu HX. Research progress of chemistry and anti-cancer activities of natural products from Chinese *Garcinia* plants. Yao Xue Xue Bao . 2014; 49(2):166-74.
- [16] Hosakatte N, Dandin V, Dalawai D, Park SY, Paek K. Bioactive compounds from *Garcinia* fruits of high economic value for food and health. Phytochem. Spr. Nature, 2018; 1: 1-28.
- [17] Hung WL, Liu CM, Lai CS, Ho CT, Pan MH. Inhibitory effect of garcinol against 12-O-tetradecanoylphorbol13acetate-induced skin inflammation and tumorigenesis in mice. Journal of Functional Foods. 2015; 18: 432-444.
- [18] Iinuma M, Tosa H, Tanaka T, Kanamaru S, Asai F, Kobayashi Y, Miyauchi K, Shimano R. Antibacterial activity of some Garcinia benzophenone derivatives against methicillin-resistant *Staphylococcus aureus*. Biol Pharm Bull. 1996; 19(2):311-4. doi: 10.1248/bpb.19.311.
- [19] IslamMZ, Hoque MM, Asif-Ul-Alam SM, Monalisa K. Chemical composition, antioxidant capacities and storage stability of *Citrus macroptera* and *Garcinia pedunculata* fruits. Emirates Journal of Food and Agriculture. 2015; 27(3): 275-282.
- [20] Jayaprakasha GK, Negi PS, Jena BS..Anti-oxidative and antimutagenic activities of the extracts from the rinds of *Garcinia pedunculata*. Innov Food Sci Emerg Technol. 2006;7: 246-50.
- [21] Joseph GS, Jayaprakasha GK, Selvi AT, Jena BS, Sakariah KK. Antiaflatoxigenic and antioxidant activities of *Garcinia* extracts. Int J Food Microbiol. 2005;101(2):153-60. doi: 10.1016/j.ijfoodmicro.2004.11.001.
- [22] Kagyung R, Gajurel PR, Rethy P. Singh B.. Ethnomedicinal plants used for gastrointestinal diseases by Adi tribes of Dehang-Debang Biosphere Reserve in Arunachal Pradesh. Indian J Tradit Know. 2010; 9: 496-501
- [23] Likhitwitayawuid K, Padungcharoen T, Krungkrai J..Antimalarial xanthones from *Garcinia cowa*. Planta Medica1998; 64:70-72.
- [24] Mackeen MM, Ali AM, Lajis NH, Kawazu K, Hassan Z, Amran M, Habsah M, Mooi LY, Mohamed SM.. Antimicrobial, antioxidant, antitumour-promoting and cytotoxic activities of different plant part extracts of *Garcinia atroviridis* griff. ex T. anders. J Ethnopharmacol. 2000;72(3): 395-402. doi: 10.1016/s0378-8741(00)00245-2.
- [25] Mahabusarakam W, Chairerk P, Taylor WC. Xanthones from *Garcinia cowa* Roxb. Latex. Phytochemistry. 2005; 66(10):1148-53. doi: 10.1016/j. phytochem. 2005.02.025.
- [26] Minami H, Kinoshita M, Fukuyama Y, Kodama M, Yoshizawa T, Sugiura M, Nakagawa K , Tago H. Antioxidant xanthones from *Garcinia subelliptica*. Phytochemistry, 1994; 36(2):501-506
- [27] Minami H, Kuwayama A, Yoshizawa T, Fukuyama Y. Novel prenylated xanthones with antioxidant property from

the wood of *Garcinia subelliptica Chem. Pharm. Bull.* 1996; 44:2103-2106

- [28] Minami H, Takahashi E, Fukuyama Y, Kodama M, Yoshizawa T, Nakagawa K. Novel xanthones with superoxide scavenging activity from Garcinia subelliptica. Chem Pharmacol Bull 1995;43:347–349.
- [29] Mudoi T, Deka DC, Devi R. *In vitro* antioxidant activity of *Garcinia pedunculata*, an indigenous fruit of North Eastern (NE) region of India. International Journal of Pharm Tech Research. 2012; 4(1): 334-342.
- [30] Mundugaru R, Joy F, Shrinidhi R, Das L, Sudhakara, Ravishankar B.. Anti-inflammatory activity of aqueous extract of fruits of *Garcinia pedunculata* in experimental animals. Am J Pharma Tech Res. 2014; 4: 3.
- [31] Mundugaru R, Udaykumar P, Senthilkumar S, Bhat S. Cardioprotective activity of fruit of *garcinia pedunculata* on isoprenaline-induced myocardial infarction in rat. Bangladesh Journal of Pharmacology.2016; 11(1): 231-235.
- [32] Mundugaru, R., Varadharajan, MC., and Basavaiah R. 2014. Hepatoprotective activity of fruit extract of *Garcinia pedunculata*. Bangladesh Journal of Pharmacology. 9(4): 483-487.
- [33] Mundugaru R, Narayana SKK, Ballal SR, Thomas J, Rajakrishnan R..Neuroprotective activity of *Garcinia pedunculata* roxb ex buch ham fruit extract against aluminium chloride induced neurotoxicity in mice. Indian J. Pharm. Educ. Res. 2016; 50: 435-441.
- [34] Mundugaru R, Sivanesan SK, Udaykumar P, Joy F, Narayana SKK, Rajakrishnan L, Al Farhan AH, Jacob T, Rajagopal R, Hisham SM.Quality standardization and nephroprotective effect of *Garcinia pedunculata* Roxb. fruit extract. Indian J. Pharm. Educ. 2017; 51: 713-721
- [35] Mundugaru R, Udaykumar P, Kumar S, Narayana SKK, Jacob T, AlFarhan AH, Rajakrishnan L. Protective effect of *Garcinia pedunculata* fruit rind in acetic acid induced ulcerative colitis. Farmacia. 2019; 67: 160-166.
- [36] Murthy HN, Dalawai D, Dewir YH, Ibrahim A. Phytochemicals and Biological Activities of *Garcinia morella* (Gaertn.) Desr.: A Review. Molecules. 2020; 25(23): 5690. doi: 10.3390/molecules25235690.
- [37] Na Pattalung P, Thongtheeraparp W, Wiriyachitra P, Taylor WC. Xanthone of *Garcinia cowa*. Planta Medica ,1994; 60(4):365-368.
- [38] Negi PS, Jayaprakasha GK, Jena BS.Antibacterial activity of the extracts from the fruit rinds of *Garcinia cowa* and *Garcinia pedunculata* against food borne pathogens and spoilage bacteria. LWT-Food Sci. Technol. 2008; 41: 1857-1861.
- [39] Paul S, Ali MY, Rumpa NE, Tanvir EM,Hossen MS,Saha M, Bhoumik NC,Gan SH, Khalil MI. Assessment of toxicity and beneficiary effects of *Garcinia pedunculata* on the hematological, biochemical, and histological homeostasis in rats. Evid. Based Complementary Altern. Med. 2017; Article ID 4686104 | https://doi.org/10.1155/2017/4686104
- [40] Policegoudra RS, Saikia S, Das J, Chattopadhyay P, Singh L., Veer V. Phenolic content, antioxidant activity,

antibacterial activity and phytochemical composition of *Garcinia lancifolia*. Indian Journal of Pharmaceutical Sciences. 2012; 74(3): 268-271.

- [41] Rao AVR, Venkataswamy G, Yemul SS. Xanthochymol and isoxanthochymol two novel polyisoprenylated benzophenones from *Garcinia xanthochymus*. Indian J Chem.1980; 19B: 627-33.
- [42] Rao AVR, Sarma MR, Venkataraman K, Yemul SS. A benzophenone and xanthone with unusual hydroxylation patterns from the heartwood of *Garcinia pedunculata*. Phytochemistry. 1974; 13: 1241-1244.
- [43] Ravi M, Febin J, Shrinidhi R, Lipika D, Sudhakara B, Ravishankar B. Anti-inflammatory activity of aqueous extract of fruits of *Garcinia pedunculata* in experimental animals. Am. J. Pharma. Tech. Res. 2014; 4: 3-6
- [44] Sahu A, Das B, Chatterjee A. Polyisoprenylated benzophenones from *Garcinia pedunculata*. Phytochemistry. 1989; 28(4):1233-1235.
- [45] Sarma R, Das M, Mudoi T, Sharma KK, Kotoky JA, Devi R. Evaluation of antioxidant and antifungal activities of polyphenol-rich extracts of dried pulp of *Garcinia pedunculata* Roxb. and *Garcinia morella* Gaertn. (Clusiaceae) Tropical Journal of Pharmaceutical Research, 2016, 15(1): 133-140. doi.org/10.4314/tjpr.v15i1.19.
- [46] Sarma R, Kumari S, Elancheran R, Deori M, Devi R. Polyphenol rich extract of *Garcinia pedunculata* fruit attenuates the hyperlipidemia induced by high fat diet. Front Pharm. 2016; 7:294
- [47] Sharma R , Devi R. Ethnopharmacological survey of *Garcinia pedunculata* Roxb. Fruit extract in six different districts of Assam, India. International Journal of Pharmaceutical Science.2015; 4(1); 20-28.
- [48] Vo HT, Ngo NT, Bui TQ, Pham HD, Nguyen LD. Geranylated tetra oxygenated xanthones from the pericarp of *Garcinia pedunculata*. Phytochem. Lett. 2015; 13: 119-122.