



**A REVIEW STUDY ON ETHNOPHARMACOLOGICAL & PHYTOCHEMICAL
COMPARISON BETWEEN *SYZYGIUM CUMINI* & *SYZYGIUM JAMBOS* OF GENUS
SYZYGIUM (FAMILY: MYRTACEAE).**

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Article Received on 27/03/2019

Article Revised on 17/04/2019

Article Accepted on 07/05/2019

ABSTRACT

This article aims to provide detailed information on two species of *Syzygium cumini* & *Syzygium jambos* from genus *Syzygium* of family Myrtaceae. An extensive search on electronic databases and conference papers was done to come across significant articles on different pharmacological activities with the traditional use of this genus. The presence of various phytochemical constituents have been reported that have significant prominence on the bioactive belongings. Hence, this current review is a detailed outline on the prospective medicinal values from previous studies. However, further exploration on the possible underlying mechanisms with the isolation of more respective active compounds remains under investigation.

KEYWORDS: *Syzygium cumini*, *Syzygium jambos*, pharmacological activities & phytochemical activities.

1. INTRODUCTION

Over 50% of all modern drugs are of natural product source and they play an significant role in drug development programs of the pharmaceutical industry.^[1] The use of herbal medicines worldwide has provided an excellent opportunity worldwide to look for therapeutic lead compounds from an ancient system of therapy, i.e. Ayurveda, which can be utilized for development of new drug. Epidemiological evidence suggests that dietary factors play an important role in human health and in the treatment of certain chronic diseases including cancer.^[2,3] The collective role of plants in the treatment of disease is emerged by their employment in all major systems of medicine irrespective of the underlying philosophical premise. Cultivation of medicinal plants with laboratory generated species is taken on the basis of chemical composition and ethnopharmacological investigation.^[4]

The genus *Syzygium* is one of the genera of the family of Myrtaceae that is widely distributed throughout tropical & subtropical regions. The whole plants possess some medicinal values according to ayurvedic, unani & sidha due to presence of volatile oil as well as other bioactive phytoconstituents. The fruit is edible & traditionally used for miscellany medicinal possessions. The present study has been performed to assess the phytochemical & pharmacological properties of two species of this genus. *Syzygium cumini* & *Syzygium jambos* are extensively known for their ethnopharmacological assets. Different

researchs were reviewed from previous studies to carry out this review study.

2. METHODOLOGY

The bibliographic research was performed in the following databases: PubMed, Google Scholar, Scopus, ScienceDirect, Classical text books of Ayurveda and other compilatory treatises where these databases were searched for relevant studies on about two species of this genus (*Syzygium cumini* & *Syzygium jambos*) in terms of phytopharmacological information. No limit was placed on the search time frame in order to retrieve all relevant papers. About 134 papers have been reviewed including journal articles and proceedings as well as the reference lists of articles for additional relevant studies.

3. *Syzygium cumini*

3.1 Plant Profile: *Syzygium cumini* (L.) Skeels (Myrtaceae) commonly known as Indian blackberry; commonly known as Black Plum in English, Jamun in Hindi, Jambu in Sanskrit and Jaman in Urdu.^[5] It is a large tree distributed throughout South & West asia also in Thailand, Philippines, Madagascar Africa, Caribbean and Tropical America. The tree is commonly grow in damp places and in evergreen forests & planted as an ornamental tree in gardens and roadsides 8. The berries are sweetish sour to taste. The ripe fruits are used for health drinks, making preserves, squashes, jellies and wine.^[6] This plant grows up to 30 meters and girth of 3.6 meters with a bole up to 15meters.^[7] The plant is

carminative, digestive, antihyperglycaemic, antihelminthic and antibacterial agent also used to cure diabetes, pharyngitis, spleenopathy, urethrorrhoea, ringworm infection, to strengthen teeth and gums,^[8,9] biliousness, dysentery, sore throat, bronchitis, thirst, asthma and ulcers.^[10] diabetes, constipation, leucorrhoea, fever, gastropathy and dermatopathy and to inhibit blood discharge in the faeces.^[11-12]

Taxonomical Classification

Kingdom- Plantae
Order- Myrtales
Family- Myrtaceae

Genus- *Syzygium*

Species- *cumini*

Synonyms

1. *Eugenia jambolana* Lam.
2. *Myrtus cumini* Linn.
3. *Syzygium jambolana* DC.
4. *Syzygium jambolanum* (Lam.) DC.
5. *Eugenia djouant* Perr.
6. *Calyptanthes jambolana* Willd.
7. *Eugenia cumini* (Linn.) Druce. and
8. *Eugenia caryophyllifolia* Lam.

3.2 Phytochemical Review

Plant Parts	Phytoconstituents present in <i>Syzygium cumini</i>	Reference
Stem Bark	Betulinic acid, β -sitosterol, friedeanol, epi-friedeanol, eugenin, β -sitosterol-D-glucoside, Kamepferol-3-O- glucoside	[13-14]
	Quercetin, myricetin, astragalol, and gallic acid.	[15]
Fruit	Malic acid, oxalic acid Gallic acid, tannins, cyanidine and diglycoside glucose, fructose, mannose, and galactose (principal sugar moieties).	[16]
	Ca, Mg, Na, K, Cu and vitamins such as thiamine, riboflavin, nicotinic acid,20	[17]
	anthocyanins, delphinidin, petunidin, malvidin-diglucosides.	[18,19, 20]
Seed	Glucoside jamboline, chlorophyll, fat, resin, gallic acid, ferulic acid guaicol, resorcinol, dimethyl ether, corilagin, protein, calcium.	[21-22]
	Phenolic such as ellagic acid, gallic acid, caffeic and ferulic acids and derivatives, guaicol, resorcinol dimethyl ether, corilagin.	[23]
	Monoterpenoids like β -pinene, terpinene, terpinolene, borbeneol, β -phellandrene, a-terpineol and eugenol,	[24]
	flavonoid such as rutin, quercetin. 11 and β -sitosterol	[25]
Leaf	Gallitanins, essential oil (terpenes, 1-limonene and dipentene), monoterpenoid terpinene, terpinolene, borbeneol, terpineol and eugenol, complicated mixture of polyphenol such as gallic acid, methylgallate, kaempferol, ellagic acid, ellagitannin, nilocitin, myricetin 3-O-D-glucuronopyranoside, 3-O- β -D-glucuronopyranoside and two flavanol glycosides such as nearsetin 2-O-(4'-O-acetyl)-a-L rhamnopyranoside, and myricetin 4''-O-acetyl''-2-O-gallate.	[26-28]
	Sitosterol, betulinic acid, crategolic (maslinic) acid, n-hepatcosane, nnonacosane, n-hentriacontane, n-octacosanol, n-triacontanol and ndotricontanol (by GLC), sugars – glucose, fructose, acidsoxalic, citric, glycolic acids and aminoacids – glycine, alanine, tyrosine and leucine.	[29]
	Quercetin	[30]
	Oleanolic acid, crategolic acid (maslinic acid) and flavonoids - isoquercitrin, quercetin	[31]
Flower	Kaempferol, quercetin, myricetin, isoquercetin (quercetin-3- glucoside), myricetin- 3 - L - arabinoside, quercetin-3-D-galactoside, dihydromyricetin, oleanolic acid, acetyl oleanolic acid, eugenol-triterpenoid A and eugenol-triterpenoid B.	[32, 33]
	Ellagic acids, isoquercetin, quercetin, kampferol and myricetin	[34]
Root	Myricetin 3-o-glucoside and myricetin 3-o-robinoside.	[35]
Essential oils	α -terpeneol, myrtenol, eucarvone, muurolol, α -myrtenal, 1, 8-cineole, geranyl acetone, α -cadinol and pinocarvone	[36]
	Terpenes, 1-limonene and dipentene.	[37]
	Llauric (2.8%), myristic (31.7%), palmitic (4.7%), stearic (6.5%), oleic (32.2%), linoleic (16.1%), malvalic (1.2%) and vernolic (3%) acids. 13 Novel compounds such as 5,6 dihydroxy-3-[(4-hydroxy-6-(hydroxymethyl)-3,5-di[3,4,5-trihydroxy-6-(hydroxymethyl)tetrahydro-2h 2pyranyl]oxy]2-methoxy-10,13 dimethylperhydrocyclopenta [a] phenanthren-17-yl (phenyl) methyl acetate, 14 3,15- dihydroxy ? 3 androstene [16, 17-C](6' methyl , 2'-1,3- dihydroxy-1-propene) 4H pyran and 3-hydroxy androstane [16,17-C](6' methyl, 2'-1-hydroxy –isopropene-1-yl) 4,5,6 H pyran. 15 androstane [16,17-C](6' methyl, 2'-1-hydroxy –isopropene-1-yl) 4,5,6 H pyran.	[38]
	The essential oils isolated from the freshly collected leaf (accounting for 82% of the oil)	[39]

	α -Pinene, camphene, β -Pinene, myrcene, limonene, cis-Ocimene, trans-Ocimene, γ -Terpinene, terpinolene, bornyl acetate, α -Copaene, β -Caryophyllene, α -Humulene, γ -Cadinene and δ -Cadinene	[40]
	Trans-ocimene, cis-ocimene, β -myrcene, α -terpineol, dihydrocarvyl acetate, geranyl butyrate, terpinyl valerate,	[41]
	α -terpineol, β -caryophyllene, α -humulene, β -selinene, calacorene, α -muurolol, α -santalol, cis-farnesol: lauric, myristic, palmitic, stearic, oleic, linoleic, malvalic, sterculic and vernolic acids	[42]
	Unsaponifiable matter of the seed fat was also chemically investigated.	[43]

3.3 Pharmacological Review

Pharmacological Activity of <i>Syzygium cumini</i>	Plant Part	Reference
Antihyperlipidaemic effect	Aqueous extract of pulp	[44]
	Seed kernel	[45]
	Fruit pulp	[46]
	Ethanoilc extract of kernels	[47]
	seeds	[48]
free radical-scavenging and antilipidperoxidative activity	Aqueous seed powder extract	[49]
	Extracts of fruit pulp, seed coat and kernel	[50-52]
Antioxidant effect	Aqueous seed powder extract)	[49, 53]
	Methanolic Leaf extract	[54]
	Methanolic extract of leaves, bark and seeds	[55-58]
Hepatoprotective effect	Ethanolc Pulp extract	[59]
	Alcoholic extract of the pulp	[60]
	Aqueous leaf extract	[61]
	Methanolic seed extract	[62]
Antiarthritic effect	Methanolic seed powder extract	[61]
Antiulcer effect	Ethanolc seed powder extract)	[63]
	Seed kernel extract	[64]
	Fruit extract	[65]
Antiallergic effect	Aqueous leaf extract	[66]
	Aqueous leaf extract	[67]
Antibacterial effect	Aqueous and acetone bark extract	[68]
	Stem, leaf and fruit extracts (Against <i>Roultella plantikola</i>)	[69]
	Seed extract (Against multidrug-resistant human bacterial pathogens)	[70]
	Ethyl acetate, petroleum ether and methanolic leaf extracts (Against <i>Salmonella typhimurium</i> , <i>Bacillus subtilis</i> , <i>Pseudomonas aeruginosa</i> , <i>Staphylococcus aureus</i> and <i>Enterobacter aerogenes</i>)	[71]
	Acetone, aqueous and ethanolic bark extracts (Against <i>Vibrio cholera</i>)	[72]
	Aqueous leaf extract (Against <i>Klebsiella sp.</i> , <i>Salmonella paratyphi A & B</i> , <i>Citrobacter sp.</i> , <i>Proteus mirabilis</i> , <i>Escherichia coli</i> , <i>Staphylococcus aureus</i> , <i>Shigella sonnei</i> , <i>Pseudomonas aeruginosa</i> , <i>Salmonella typhimurium</i> , <i>Shigella boydii</i> , <i>Streptococcus faecalis</i> , <i>Shigella flexneri</i> and <i>Salmonella typhi</i>)	[73]
Anti-inflammatory effect	Ethyl acetate and methanolic seed powder extract	[74]
	Ethyl-acetate and methanolic leaves and seeds extracts	[75,76]
Nephroprotective activity	FIIc, isolated from aqueous pulp Extract	[77]
Antidiarrhoeal effect	Ethanolc bark extract	[78]
	Ethanolc fruit extract	[79]
Central nervous system effect	Methanolic and ethyl acetate seed extracts	[80]
Neuropsychopharmacological	Ethyl acetate and methanolic seed powder extract	[80]

Hypothermic	Chloroform and methanolic fruit extracts	[81]
Antifertility effect	Flowers extract (Oleanolic acid)	[82]
Radioprotective effect	Hydroalcoholic seed powder extract	[83]
	Leaf extract	[84]
	Dichloromethane extract of leaf and Hydroalcoholic seed extract	[85,86]
	Leaf extract	[87]
Chemoprotective	Aqueous and ethanolic SC	[88-90]
Diuretic	Petroleum ether, chloroform, methanolic and aqueous bark extracts	[91]
Cardioprotective	the hydroalcoholic leaves extract	[92,93]
	aqueous suspension of seed extract	[94]
	Methanolic extract of fruit	[95]
Antiviral	Hot aqueous bark extract	[96]
	Aqueous leaves extract	[97, 98]
Ascaricidal	Ethanolic, hexane and ethyl acetate extracts of fruit & leaf	[99]
Antilishmanial	The essential oil (α -pinene)	[100]
Antinociceptive	Hydro-alcoholic leaf extract	[101]
	Methanolic fruit extract	[102]
Antifungal	Methanolic fruit extract (Against <i>Fusarium oxysporium</i> , <i>Rhizoctonia solani</i> and <i>Sclerotium rolfsii</i>)	102
	n-hexane, alcohol and aqueous extracts of different plant parts (barks of stem and roots, fruits and leaves) (Against <i>Ascochyta rabiei</i>)	[103]
Anti-diabetic	Seeds, fruit pulp whole fruit bark leaves and flowers	[104,105]
Anticlastogenic	Alcoholic seed extract	[106]
Anti cancer	Fruit extract (Ellagitannins)	[107,108,109]

4. *Syzygium jambos*

4.1 Plant Profile: *Syzygium jambos* Alston (syn. *Eugenia jambos* L.; *Jambosa jambos* Millsp.; *Jambosa vulgaris* DC.; *Caryophyllus jambos* Stokes) (Family: Myrtaceae) is an evergreen tree. It is native to Southeast Asia. It is a small tree with spreading branches, leaves, simple, opposite, lanceolate, narrowed into short petioles, secondary nerves joined by a prominent looping intramarginal vein. Flowers greenish white in short terminal racemose cymes, stamens many, yellowish white, fruits pale yellow to pinkish white, globose, seeds 1-2, grey in large cavity of the succulent pulp.^[110] Due to medicinal properties this plant has some traditional use such as to treat fever, diarrhea, dysentery, rheumatism, sore eyes, asthma, bronchitis, hoarseness.^[111] epilepsy, diabetes,^[112] herpes simplex type 1 and type 2, vesicular

somatitis virus.^[113,114] toothache, mouth sores, cough, wound dressing,^[115] respiratory disorders, eczema, malaria, and infectious diseases.^[116] Other study claimed anesthetic, diuretic febrifuge activity,^[117,118] of the plant.

Taxonomical Classification

Kingdom- Plantae
Order- Myrtales
Family- Myrtaceae
Genus- *Syzygium*
Species- *jambos*

Synonyms

1. *Eugenia jambos* *Myrtus cumini* Linn.
2. *Jambosa jambos* *Syzygium jambolanum* (Lam.) DC.

4.2 Phytochemical review

Plant Parts	Phytoconstituents present in <i>Syzygium jambos</i>	Reference
Leaves	Polyphenols, anthraquinones, tannins, and steroids	[118]
	Friedelin (1)	[119]
	Amyrin acetate	[120]
	Betulinic acid (2)	[121]
	Lu peol	[121]
Bark	Friedelolactone (3)	[122]
	Friedelanol (4)	[123]
	Polyphenols, anthraquinones, tannins, and steroids	[118]
	Triterpenes and saponins	[124]
	Triterpenoids such friedelin, β -amyrin acetate, betulinic acid, and lupeol	[125]

4.3 Pharmacological Review

Pharmacological Activity of <i>Syzygium jambos</i>	Plant Part	Reference
Antifungal activity (Against <i>T. mentagrophytes</i> and <i>T. soudanense</i>)	Ethyl acetate extract of the stem bark	[125]
Antidermatophytic activity	Crude extract and fractions	[126]
Antibacterial activity potential (against sensitive strains of <i>Staphylococcus aureus</i> , <i>Bacillus subtilis</i> , <i>Enterococcus gallinarum</i> , <i>Escherichia coli</i> , <i>Pseudomonas aeruginosa</i> , <i>Klebsiella pneumoniae</i> , <i>Proteus vulgaris</i> , <i>Enterococcus faecium</i> , <i>Salmonella typhi</i> , and <i>Vibrio cholera</i>)	Fruit extracts	[127]
	Bark extracts	[128]
Antibiotic-modulating activity (potentiate the activity of antibiotics on more than 70% of bacteria) of extracts at MIC/2 on more than 70% tested strains of <i>S. aureus</i>	Methanol extracts of bark and leaves	[129,130]
Antimicrobial activity (Minimum inhibitory concentrations for <i>faecalis</i> 797.5 µg/mL and <i>A. hydrophilia</i> 384.6 µg/mL <i>B. cereus</i> 182.6 µg/mL and <i>S. aureus</i> 346.5 µg/mL)	Acetone and aqueous extracts from the bark	[131]
Hepatoprotective agent	Methanolic extract of leaf	[132]
Analgesic effects	Hydro-alcoholic leaf extracts	[133]
Anti oxidant, anti inflammatory, anti diabetic, anticancer, anti ulcer, anti pyretic, cardio vascular diseases, anti hyperlipidimic and neurological disorders like alzheimer's, anti parkinsonism	Extract of fruit, leaf & bark.	[134]

CONCLUSION

Large scale literature study results revealed that the featured plants have potential pharmacological activities against various diseases performed *in vivo* & *in vitro*. The phytoconstituents which are present in the plants are mainly alkaloid, tannin, glycoside, carbohydrate, saponin, steroids and flavonoids which are responsible for the bioactivity. Further significant review is needed to find out the appropriate estimation and revival on pharmacological effect. Chemical investigations are needed to find out the future lead compound to develop drugs. Plant sources are being used for a long time to explore our medicinal sector because of their less side effects and more effectiveness.

CONFLICT OF INTEREST

We have no conflict of interest.

ACKNOWLEDGEMENT

We would like to give thanks to the Department of Pharmacy, Southeast University for technical support

REFERENCE

- Baker JT, Borriss RP, Carte B, Cordell GA, Soejarto DD. Natural product drug discovery and development: New perspective on international collaboration. *J Natl Prod.*, 1995; 58: 1325-57.
- Trichopoulos D, Willett WC. Nutrition and cancer. *Cancer Causes Cont*, 1996; 7: 3-4.
- Block G. The data support a role for antioxidants in reducing cancer risk. *Nutr Rev*, 1992; 50: 207-13.
- National Pharmacovigilance Protocol For Ayurveda, Siddha And Unani (Asu) Drugs; Gujarat Ayurved University, Jamnagar, Gujarat, 2008; 8-50.
- Deepti Katiyar *et al Adv. Appl. Sci. Res.*, 2016, 7(4): 1-12.
- Williamson, E. M., Jamun (*Syzygium cumini* (L.): A Review of Its Food and Medicinal Uses. Major Herbs of Ayurveda, Churchill Livingstone, China, 2002; 279-282.
- Jagetia GC and Baliga MS, *Syzygium cumini* (Jamun) reduces the radiation-induced DNA damage in the cultured human peripheral blood lymphocytes: a preliminary study. *Toxicol Lett.* 2002; 132: 19-25.
- Pepato MT, Folgado VBB, Kettelhut IC and Brunetti IL, Lack of antidiabetic effect of a *Eugenia jambolana* leaf decoction on rat streptozotocin diabetes. *Braz J Med Biol Res.*, 2001; 34: 389-395
- Balinga MS, Bhat HP, Balinga BRV, Wilson R, Palatty PL, *Food Res. Int.*, 2011; 1-14.
- Pari L and Saravanan G, Antidiabetic effect of Cogent db, a herbal drug in alloxan-induced diabetes mellitus. *CompBiochem PhysiolCToxicolPharmacol.* 2002; 131: 19-25.
- Mitra SK, Gopumadhavan S, Muralidhar TS, Anturlikar SD and Sujatha MB, Effect of D-400, a herbomineral preparation on lipid profile, glycated haemoglobin and glucose tolerance in streptozotocin induced diabetes in rats. *Indian J Exp Bio.*, 1995; 33: 798-800.
- Sengupta P, Das PB. Terpenoids and related compounds part IV triterpenoids the stem-bark of *Eugenia jambolana* Lam. *Indian Chem Soc*, 1965; 42: 255-258.

13. Bhargava KK, Dayal R, Seshadri TR. Chemical components of *Eugenia jambolana* stem bark. *Curr Sci*, 1974; 43: 645–646.
14. Panda DK, Ghosh D, Bhat B, Talwar SK, Jaggi M, Mukherjee R., *Meth. Find. Exper. Clin. Pharmacol.*, 2009; 31: 571–584.
15. Veigas JM, Narayan MS, Laxman PM, Neelwarne B. Chemical nature stability and bioefficacies of anthocyanins from fruit peel of *Syzygium cumini* Skeels. *Food Chem*, 2007; 105: 619–627.
16. Vijayanand P, Rao LJM, Narasimham P. Volatile flavour components of Jamun fruit (*Syzygium cumini*) Flavour Fragr J., 2001; 16: 47–49.
17. Duke James A, *Handbook of Medicinal Herbs*, 2nd edition, CRC Press, London, 2006; 422.
18. Brito, F. A., Lima, L. A., Ramos, M. F., Nakamura M.J., Cavalher-Machados S.C., Henriques M.G., Sampaino A.L., Pharmacological study of anti-allergic activity of *Syzygium cumini* (L) Skeels., *Brazilian journal of medical and biological research*, 2007; 40: 105-115.
19. Chaturvedi, A., Kumar M. M., Bhawani, G., Chaturvedi, H., Effect of ethanolic extract of *Eugenia jambolana* seeds on gastric ulceration and secretion in rats *Indian J. physiol Pharmacol*, 2007; 51(2): 131-140.
20. Daulatabad CMJD, Mirajkar AM, Hosamani KM, Mulla GMM. Epoxy and cyclopropenoid fatty acids in *Syzygium cumini* seed oil. *J Sci Food Agric*, 1988; 43: 91–94.
21. Gupta RD, Agrawal SK. Chemical examination of the unsaponifiable matter of the seed fat of *Syzygium cumini*. *Sci Cult*, 1970; 36: 298.
22. Waheed A, Miana GA, Ahmad SI, *Pak. J. Pharmacol*, 2007; 24(1): 13-17.
23. Timbola, A. K., Szpoganicz, B., Branco, A., Monache, F. D. and Pizzolatti, M. G., A new flavonol from leaves of *Eugenia jambolana*, *Fitoterapia*, 2002; 73: 174-176.
24. Rohit Kumar Bijauliya, Shashi Alok, Man Singh and Shanti Bhushan Mishra, Morphology, phytochemistry and pharmacology of *syzygium cumini* (linn.) - an overview, *IJPSR*, 2017; 8(6): 2360-2371.
25. Pandey M, Khan A. Hypoglycaemic effect of defatted seeds and water soluble fibre from the seeds of *Syzygium cumini* (Linn.) skeels in alloxan diabetic rats. *Indian journal of experimental biology*, 2002; 40(10): 1178- 1182.
26. Bhatia IS, Sharma SK, Bajaj KL. Esterase and galloyl carboxylase from *Eugenia jambolana* leaves. *Indian JExp Biol.*, 1974; 12: 550–552.
27. Kumar A, Naqvi AA, Kahol AP, Tandon S. Composition of leaf oil of *Syzygium cumini* L, from north India. *Indian Perfum*, 2004; 48: 439–441.
28. Gupta, G. S., Sharma, D. P., Triterpenoid and other constituents of *Eugenia jambolana* leaves, *Phytochemistry*, 1974; 13: 2013-2014.
29. Mahmoud, I. I., Marzouk, S. A., Moharram, F. A., El-Gindi, M. R., Hassan, A.M.K. Acylated flavonol glycosides from *Eugenia jambolana* leaves, *Phytochemistry*, 2001; 58: 1239-1244.
30. Nair, A. R. and Subramanian, S. S., Chemical examination of the flowers of *Eugenia jambolana*, *J. Sci. Industry, Res.*, 1974; 457-458.
31. Sagrawat H, Mann AS, Kharya MD. Pharmacological potential of *Eugenia jambolana*: a review. *Pharmacogn Mag*, 2006; 2: 96–104.
32. Mokkhasmit, M., Swasdimongkol, K., Ngarmwathana, W. and MPP Stanley, N.Kamalakkannan, V.P. Menon. J. *Ethanopharmacol*, 2003; 84(2-3): 205-209.
33. Duke James A, *Handbook of Medicinal Herbs*, 2nd edition, CRC Press, London, 2006; 422.
34. LewisYS, Dwarakanath CT, Johar DS. Acids and Sugars in *Eugenia jambolana*. *J. Sci. Idustur. Res.*, 1956; 15C: 280-81.
35. Herculano EDA, Costa CDF, Rodrigues AKBF, Junior JXA, Santana AEG, França PHB, et al., *Trop. J. Pharm. Res.*, 2014; 13(11): 1853-1861.
36. *The Wealth of India*, Vol-X, CSIR, New Delhi, 1982; 100-104.
37. Shankar, M. B., Parikh, J. R., Geetha, M., Mehta, R. S. & Saluja, A. K., Anti-diabetic activity of novel androstane derivatives from *Syzygium cumini* Linn, *Journal of Natural Remedies*, 2007; 712: 214-219.
38. Chandra D, Jackson EB, Ramana KV, Kelley R, Srivastava SK and Bhatnagar A, Nitric oxide prevents aldose reductase activation and sorbitol accumulation during diabetes. *Diabetes*, 2002; 51: 3095–3101.
39. *The Wealth of India*, Vol-X, CSIR, New Delhi, 1982; 100-104.
40. Vijayanand P, Rao LJM, Narasimham P. Volatile flavour components of Jamun fruit (*Syzygium cumini*) Flavour Fragr J., 2001; 16: 47–49.
41. Daulatabad CMJD, Mirajkar AM, Hosamani KM, Mulla GMM. Epoxy and cyclopropenoid fatty acids in *Syzygium cumini* seed oil. *J Sci Food Agric*, 1988; 43: 91–94.
42. Gupta RD, Agrawal SK. Chemical examination of the unsaponifiable matter of the seed fat of *Syzygium cumini*. *Sci Cult.*, 1970; 36: 298.
43. Rekha N, Balaji R and Deecaraman M, Effect of aqueous extract of *Syzygium cumini* pulp on anti-oxidant defense system in streptozotocin induced diabetic rats. *Iran J Pharmacol Therapeu*, 2008; 7: 137–145.
44. Ravi K, Rajasekaran S and Subramanian S, Anti-hyperlipidemic effect of *Eugenia jambolana* seed kernel on streptozotocin-induced diabetes in rats. *Food Chem Toxicol*, 2005; 43: 1433–1439.
45. Rabiea B, Manzar Z, Ahmad U, Shahnaz A, Azam Z, *J. Pak. Med. Assoc.*, 2011; 61(5): 433-437.
46. Kasiappan R, Subbaih R, Sorimuthu S, *Food Chem. Toxicol.*, 2005; 43: 1433–1439.
47. Sharma B, Balomajumder C, Roy P, *Food Chem. Toxicol.*, 2008; 46: 2376–2383.
48. AhmedF, Huded S and Malik FA, *In vitro* study on the radical scavenging and antilipidperoxidative

- effects of *Eugenia jambolana* aqueous extracts. *J Pharm Res.*, 2010; 3: 198–200.
49. Bhuiyan MS, Younus Mia M, Rashid MA. Antibacterial principles of the seeds of *Eugenia jambolana*. *Bangladesh J. Bot.*, 1996; 25(2): 239-41.
 50. Abalea V, Cillard J, Dubos MP, Sergent O, Cillard P and Morel I, Repair of iron induced DNA oxidation by the flavonoid myricetin in primary rat hepatocyte culture. *Free Radic Biol Med.*, 1999; 26: 1457–1466.
 51. Huang L, Yuan X, Aiken C, Chen CH. Bifunctional anti-human immunodeficiency virus type 1 small molecules with two novel mechanisms of action. *Antimicrobial agents and chemotherapy*, 2004; 48(2): 663-5.
 52. Prince PSM, Kamalakkannan N and Menon VP, *Syzygium cumini* seed extracts reduce tissue damage in diabetic rat brain. *J Ethnopharmacol*, 2003; 84: 205–209.
 53. Ruan ZP, Zhang LL and Lin YM, Evaluation of the antioxidant activity of *Syzygium cumini* leaves. *Molecule*, 2008; 13: 2545–2556
 54. Singh JP, Kaur A, Singh N, Nim L, Shevkani K, Kaur H, Arora DS, *LWT - Food Sci. Technol.*, 2016; 65: 1025-1030.
 55. Haroon R, Jelani S, Arshad FK, *Int. J. Res.-Granthaalayah*, 2015; 3(5): 13-26.
 56. Nair LK, Begum M, Geetha S, *JESTFT*, 2013; 7(1): 54-62.
 57. Rufino MSM, Alves RE, Fernandes FAN, Brito ES, *Food Res. Int.*, 2011; 44: 2072–2075.
 58. Das S and Sarma G, Study of the hepatoprotective activity of the ethanolic extract of the pulp of *Eugenia jambolana* (Jamun) in albino rats. *J Clin Diagn Res.*, 2009; 3: 1466–1474.
 59. Kirtikar KR and Basu B D. *Indian Medicinal Plants, Vol-II, Periodical Experts*, New Delhi, 1975; 1052-1053.
 60. Moresco RN, Sperotto RL, Bernardi AS, Cardoso RF, Gomes P, *Phytother. Res.*, 2007; 21: 793–795.
 61. Sisodia SS, Bhatnagar M, *Ind. J. Pharmacol.*, 2009; 41: 23–27.
 62. Ivan AR. *Medicinal plant of World: Chemical Constituents, Traditional Uses and Modern Medicinal Uses*, Human Press Totowa, New Jersey, 2006; 283-289.
 63. Chaturvedi A, Bhawani G, Agarwal PK, Goel S, Singh A, Goel RK, *Ind. J. of Physiol. & Pharmacol.*, 2009; 53: 16–24.
 64. Huang L, Yuan X, Aiken C, Chen CH. Bifunctional anti-human immunodeficiency virus type 1 small molecules with two novel mechanisms of action. *Antimicrobial agents and chemotherapy*, 2004; 48(2):663-5.
 65. Sood R, Swarup D, Bhatia D, Kulkarni DD, Dey S, Saini M, Dubey SC, *Ind. J. Exp. Biol.*, 2012; 50: 179-18.
 66. Kokate, C.K, AP Purohit and SB Gokhale. *Pharmacognosy*, 14th edition, NiraliPrakashan, Pune, 2008; 257-258.
 67. D. E. Djeussi, J. A. K. Noumedem, J. A. Seukep *et al.*, “Antibacterial activities of selected edible plants extracts against multidrug-resistant Gram-negative bacteria.” *BMC Complementary and Alternative Medicine*, 2013; 13 (1); 164.
 68. Pareek A, Meena RK, Yadav B, *Ind. J. Appl. Res.*, 2015; 5(9): 64-66.
 69. Bag A, Bhattacharyya SK, Pal NK, Chattopadhyay RR, *Microb. Res.*, 2012; 167: 352–357.
 70. M, Baravalia Y, Vaghasiya Y, Chanda S, *Ind. J. Pharm. Sci.*, 2009; 71: 406–412.
 71. Sharma A, Patel VK, Chaturvedi AN, *Ind. J. Pharmacol.*, 2009; 41: 129–133.
 72. Satish S, Raghavendra MP, Raveesha KA, *Adv. in Biol. Res.*, 2008; 2: 44–48.
 73. Kumar A, Ilavarasan R, Jayachandran T, Deecaraman M, Kumar RM, Aravindan P, *et al.*, Anti-inflammatory activity of *Syzygium cumini* seed. *Afr J Biotechnol*, 2008; 7: 941–943.
 74. Jain A, Sharma S, Goyal M, Dubey S, Jain S, Sahu J, Sharma A, Kaushik A, *Int. J. Phytomed.*, 2010; 2: 124-126.
 75. Kapoor LD, *Handbook of Ayurvedic Medicinal Plants*. CRC Press, London, 2001; 179–180.
 76. Tanwar RS, Sharma SB, Singh UR and Prabhu KM, Attenuation of renal dysfunction by anti-hyperglycemic compound isolated from fruit pulp of *Eugenia jambolana* in streptozotocin induced diabetic rats. *Indian J Biochem Biophys*, 2010; 47: 83–89.
 77. Mukherjee PK, Saha K, Murugesan T, Mandal SC, Pal M and Saha BP, Screening of anti-diarrhoeal profile of some plant extracts of a specific region of West Bengal, *India. J Ethnopharmacol*, 1998; 60: 85–89.
 78. Krishnakumar K, Augusti KT and Vijayammal PL, Hypolipidaemic effect of *Salacia oblonga* Wall. root bark in streptozotocin diabetic rats. *Med Sci Res.*, 2000; 28: 65–67.
 79. Kumar A, Padmanabhan N and Krishana MRV, Central nervous system activity of *Syzygium cumini* seeds. *Pak J Nutr*, 2007; 6: 698–700.
 80. Hirano T, Arimitsu J, Higa S, Naka T, Ogata A, Shima Y, *et al.*, Luteolin, a flavonoid, inhibits CD40 ligand expression by activated human basophils. *Int Arch Allergy Immunol*, 2006; 140: 150–156.
 81. Rajasekaran M, Bapna JS, Lakshmanan S, Ramchandran Nair AG, Veliath AJ and Panchanadam M, Antifertility effect in male rats of oleanolic acid, a triterpene from *Eugenia jambolana* flowers. *J Ethnopharmacol*, 1988; 24: 115–121.
 82. Satish S, Mohana DC, Ranhavendra MP, Raveesha KA. Antifungal activity of some plant extracts against important seed borne pathogens of

- Aspergillus sp. An International Journal of Agricultural Technology, 2007; 3(1): 109-19.
83. Jagetia GC, Shetty PC, Vidyasagar MS, *Ph. OL*, 2008; 1: 169-195.
 84. Jagetia GC, Baliga MS, *Die. Nahrung.*, 2003; 47: 181-185.
 85. Jagetia GC, Baliga MS, Venkatesh P, *J. Radia. Res.*, 2005; 46: 59-65.
 86. Jagetia GC, Balinga MS, *Toxicol. Let.*, 2002; 132(1): 19-25.
 87. Parmar J, Sharma P, Verma P, Goyal PK, *Asia. Pac. J. Can. Prev.*, 2011; 11: 261-265.
 88. Jain A, Sharma S, Goyal M, Dubey S, Jain S, Sahu J, Sharma A, Kaushik A, *Int. J. Phytomed.*, 2010; 2: 124-126.
 89. Goyal PK, Verma P, Sharma P, Parmar J, Agarwal A, *Asia. Pac. J. Can. Prev.*, 2010; 11: 753-758.
 90. Thapa BR and Walia A, Liver function tests and their interpretation. *Indian J Pediatr*, 2007; 74: 663-671.
 91. Ahmed F, Huded S and Malik FA, *In vitro* study on the radical scavenging and antilipidperoxidative effects of *Eugenia jambolana* aqueous extracts. *J Pharm Res*, 2010; 3: 198-200.
 92. Javanmardi J, Stushnoff C, Locke E and Vivanco JM, Antioxidant activity and total phenolic content of Iranian *Ocimum* accessions. *FoodChem*, 2003; 83: 547-550.
 93. Pizzale L, Bortolomeazzi R, Vichi S, Uberegger E and Conte LS, Antioxidant activity of sage (*Salvia officinalis* and *S fruticosa*) and oregano (*Origanum onites* and *O indercedens*) extracts related to their phenolic compound content. *J Sci Food Agric*, 2002; 82: 1645-1651.
 94. Ruan ZP, Zhang LL and Lin YM, Evaluation of the antioxidant activity of *Syzygium cumini* leaves. *Molecule*, 2008; 13: 2545-2556.
 95. Sood R, Swarup D, Bhatia D, Kulkarni DD, Dey S, Saini M, Dubey SC, *Ind. J. Exp. Biol.*, 2012; 50: 179-18.
 96. Schossler DRC, Mazzanti CM, Luz SCA, Filappi A, Prestes D, Silveira AF, *et al.*, *Syzygium cumini* and the regeneration of insulin positive cells from the pancreatic duct. *Braz J VetRes AnimSc.*, 2004; 41: 236-239.
 97. Bhanuprakash V, Hosamani M, Balamurugan V, Gandhale P, Naresh R, Swarup D, *Ind. J. Exp. Biol.*, 2008; 46: 120-127.
 98. Bhanuprakash V, Hosamani M, Balamurugan V, Singh RK, Swarup D, *Int. J. Trop. Med.*, 2007; 2: 3-9.
 99. Middleton E, Effect of plant flavonoids on immune and inflammatory cell function. *Adv ExpMed Biol*, 1998; 439: 175-182
 100. Li L, Zhang Y, Seeram NP, *Nat. Prod. Commun.*, 2009; 4: 217-219.
 101. Bhatia IS and Bajaj KL, Chemical constituents of the seeds and bark of *Syzygium cumini*. *Planta Med*, 1975; 28: 346-352
 102. Elangovan V, Shohami E, Gati I and Kohea R, Increased hepatic lipid soluble antioxidant capacity as compared to other organs of streptozotocin-induced diabetic rats: a cyclic voltammetry study. *Free Radic Res.*, 2000; 32: 125-134.
 103. Gumieniczek A, Hopkala H, Wojtowich Z and Nikolajuk J, Changes in nti-oxidant status of heart muscle tissue in experimental diabetes in rabbits. *Acta Biochim Po*, 2002; 49: 529-535.
 104. V. Kuete, S. Alibert-Franco, K. O. Eyong *et al.*, "Antibacterial activity of some natural products against bacteria expressing a multidrug-resistant phenotype," *International Journal of Antimicrobial Agents*, 2011; 37(2): 156-161.
 105. Al-Shamaony L, Al-Khazrajoi SM and Twaij HAA, Hypoglycaemic effect of *Artemisia herba alba*. II. Effect of a valuable extract on some blood parameters indiabetic animals. *JETHNOPHARMACO*, 1994; 43: 167-171.
 106. Gupta GS and Sharma DP, Triterpenoid and other constituents of *Eugenia jambolana* leaves. *Phytochemistry.*, 1974; 13: 2013-2014
 107. Daulatabad CMJ, Mirajkar AM, Hosamani KM and Mulla MMG, Epoxy and cyclopropenoid fatty acids in *Syzygium cumini* seed oil. *J Sci Food Agric*, 2006; 43: 91-94.
 108. Jasmine R, Daisy P, *Asia. J. Biochem.*, 2007; 2(4): 269-273.
 109. Sengupta P and Das PB, Terpenoids and related compounds. Part IV. Triterpenoids the stem-bark of *Eugenia jambolana* Lam. *Indian Chem So.*, 1965; 42: 255-258.
 110. Bhargava KK, Dayar R and Seshadri TR, Chemical components of *Eugenia jambolana* stem bark. *Curr Sci.*, 1974; 43: 645-646
 111. Warriar PK, Nambiar VP, Ramankutty C. Indian medicinal plants. A Compendium of 500 species, 1996; 5: 229-31.
 112. Morton J. Rose Apple. USA: Fruits of Warm Climates Florida Flair Books, 1987.
 113. Teixeira CC, Weinert LS, Barbosa DC, Ricken C, Esteves JF, Fuchs FD. *Syzygium cumini* (L.) Skeels in the treatment of type 2 diabetes. Results of a randomized, double-blind, doubledummy, controlled trial. *Diabetes Care.*, 2004; 27: 3019-20.
 114. Athikomkulchai S, Lipipun V, Leelawittayanont T, Khanboon A, Ruangrunsi N. Anti-herpes simplex virus activity of *Syzygium jambos*. *J Health Res.*, 2008; 22: 49-51.
 115. Abad MJ, Bermejo P, Villar A. Antiviral activity of medicinal plant extracts. *Phytother Res.*, 1997; 11: 198-202.
 116. Iwu MM. 1993. Handbook of African Medicinal Plants. CRC Press:Florida, London, Tokyo I.K. Voukeng, V. P. Beng, and V. Kuete, "Multidrug resistant bacteria are sensitive to *Euphorbia prostrata* and six others Cameroonian medicinal plants extracts," *BMC Research Notes*, 2017; 10: 321.

117. S. Mohanty and I. E. Cock, Bioactivity of *Syzygium jambos* methanolic extracts: Antibacterial activity and toxicity. *Pharmacognosy Res.*, 2010; 2(1): 4–9.
118. M. M. Cowan, “Plant products as antimicrobial agents,” *Clinical Microbiology Reviews*, 1999; 12(4): 564–582.
119. Candy HA, McGarry EJ, Pegel KH. Constituents of *Syzygium cordatum*. *Phytochemistry*, 1968; 7: 889–890.
120. Matsunaga S, Tanaka R, Akagi M. Triterpenoids from *Euphorbia maculata*. *Phytochemistry*, 1988; 27: 535–537
121. Sholichin M, Yamasaki K, Kasai R, Tanaka O. ¹³C Nuclear magnetic resonance of lupane-type triterpenes, lupeol, betulin and betulinic acid. *Chem Pharm Bull*, 1980; 28: 1006–1008.
122. Patra A, Chaudhuri KS. Assignment of carbon-13 nuclear magnetic resonance spectra of some friedelanones. *MagnReson Chem*, 1987; 25: 95–100.
123. Gunusekera PS, Sultanbawa MUS. Chemical investigation of Ceylonese plants. Part 22. Extractives of *Trichadenia zeylanica* Thw. (Flacourtiaceae); isolation and structures of six new triterpenoids containing the friedelane skeleton. *J C S Perkin*, 1997; 1: 483–490.
124. A. Paudel, H. Hamamoto, Y. Kobayashi, S. Yokoshima, T. Fukuyama, and K. Sekimizu, “Identification of novel deoxyribofuranosyl indole antimicrobial agents,” *The Journal of Antibiotics*, 2012; 65(2): 53–57.
125. J.-R. Kuate, S. Mouokeu, H. K. Wabo, and P. Tane, “Antidermatophytic triterpenoids from *Syzygium jambos* (L.) Alston (Myrtaceae),” *Phytotherapy Research*, 2007; 21(2): 149–152.
126. Zacchino SA, Lopez S, Pezzenati G *et al.* *In vitro* evaluation of antifungal properties of phenylpropanoids and related compounds acting against dermatophytes. *J Nat Prod*, 1999; 62: 1353–1357.
127. S. Murugan, D. P. Uma, P. N. Kannika, and K. R. Mani, “Antimicrobial activity of *Syzygium jambos* against selected human pathogens,” *International Journal of Pharmacy and Pharmaceutical Sciences*, 2011; 3: 44–47.
128. Katiyar D, Singh V, Ali M. Recent advances in pharmacological potential of *Syzygium cumini*: A review. *Adv. Applied Sci. Res.*, 2016; 7: 1-2.
129. Brice E. N. Wamba, Paul Nayim, Armelle T. Mbaveng, Igor K. Voukeng, Joachim K. Dzotam, Ornella J. T. Ngalani, and Victor Kuete, *Syzygium jambos* Displayed Antibacterial and Antibiotic-Modulating Activities against Resistant Phenotypes, *Evidence-Based Complementary and Alternative Medicine*, 2018; 12.
130. L. C. Braga, A. A. M. Leite, K. G. S. Xavier *et al.*, “Synergic interaction between pomegranate extract and antibiotics against *Staphylococcus aureus*,” *Canadian Journal of Microbiology*, 2005; 51(7): 541–547
131. Corine Djadjo Djipa, Michel Delmée, Joëlle Quetin-Leclercq, Antimicrobial activity of bark extracts of *Syzygium jambos* (L.) Alston (Myrtaceae). *Journal of Ethnopharmacology*, 2000; 71(1-2): 307–313.
132. N. Thamizh Selvam, V. Venkatakrishnan, R. Dhamodharan, S. Murugesan, and S. Damodar Kumar, Hepatoprotective activity of methanolic extract of *Syzygium jambos* (Linn.) leaf against paracetamol intoxicated Wistar albino rats. *Ayu*, 2013; 34(3): 305–308.
133. Avila-Peña D, Peña N, Quintero L, Suárez-Roca H. Antinociceptive activity of *Syzygium jambos* leaves extract on rats. *J Ethnopharmacol*, 2007; 112: 380–385
134. Ramirez RO, Roa Jr CC. The gastroprotective effect of tannins extracted from duhat (*Syzygium cumini* Skeels) bark on HCl/ethanol induced gastric mucosal injury in Sprague-Dawley rats. *Clinical hemorheology and microcirculation*, 2003; 29(3,4): 253-61.