

**A COMPREHENSIVE BIOLOGICAL, ETHNO- PHARMACOLOGICAL
and PHYTOCHEMICAL UPDATE REVIEW ON AYURVEDIC PLANT
OF *TERMINALIA CHEBULA* (HORTOKI) OF BANGLADESH**

¹A.K.M. Mohiuddin, ²Maidul Islam, ¹Shahin Mahmud, ³Md. Aminul Islam Apu,
³Joyanta Halder, ¹Md. Sadek Hosen Khoka, ¹Hasibul Haque Rakib, ¹Binita Shome and
¹Md. Shariful Islam *

¹Faculty of Life Science, Department of Biotechnology and Genetic Engineering, Mawlana Bhashani Science and Technology University, Santosh, Tangail-1902, Bangladesh.

²Department of Biochemistry and Biotechnology, Khwaja Yunus Ali University, Enayetpur, Sirajgonj-6751, Bangladesh.

³Department of Biotechnology and Genetic Engineering, Faculty of Applied Science and Technology, Islamic University, Kushtia-7003, Bangladesh.

Article Received on
07 March 2015,

Revised on 30 March 2015,
Accepted on 20 April 2015

***Correspondence for**

Author

Md. Shariful Islam

Faculty of Life Science,
Department of
Biotechnology and
Genetic Engineering,
Mawlana Bhashani
Science and Technology
University, Santosh,
Tangail-1902, Bangladesh

ABSTRACT

In the new era of Biotechnology, modern medicine system is so so advanced but still now, some of the common diseases are successfully and easily treated with the ayurvedic or herbal medicinal treatment; which is a major and important part of the modern treatment system. *Terminalia chebula* is termed the “king of medicine” in the ayurvedic land of science (medicinal harbal science), due to its huge quantity of pharmacological, biological and phytochemical rich constituents. HORTOKI is the Bengali term of this plant. The demand for herbal therapeutics is now increasing in modern biological science gradually worldwide. *Terminalia chebula* is one of the most commonly used plants in traditional systems of medicine in Bangladesh. It is a mild, safe and effective laxative in traditional medicine. It is reported to contain various pharmacological activities including antioxidant, antidiabetic, antibacterial, antiviral, antifungal, anticancerous,

antiulcer, antimutagenic, wound healing, immunomodulatory, cardioprotective effect, anti-aging, cytoprotective and hepatoprotective activities. It has been reported to contain various biochemical constituents including tannins, chebulinic acid, ellagic acid, gallic acid, punicalagin and flavonoids. Several pharmacological investigations for different biological activities of *Terminalia chebula* in various in vivo and in vitro test models have been carried

out based on the presence of biochemical ingredients and pharmacological findings. This update review gives a bird's eye view on the biological and ethno- pharmacological properties of various phytoconstituents and the biological uses of *Terminalia chebula* (HORTOKI) to enrich our knowledge about this plant.

KEYWORDS: *Terminalia chebula*, Pharmacological studies, Phytoconstituents, King of medicine, chebulinic acid, antimutagenic, antidiabetic.

1. INTRODUCTION

Terminalia chebula ((HORTOKI) is a moderate tree used in traditional ayurvedic medicinal uses. it is a popular traditional medicine not only used in Bangladesh but also in other countries of Asia and Africa. This is used in ayurvedic medicine due to the wide spectrum of pharmacological, phytochemical and biological activities associated with the biologically active chemical compounds present in this plant.^[15] It is used for the treatment of number of diseases like cancer, paralysis, cardio vascular diseases, ulcers, leprosy, arthritis and gout etc. It has been reported as antioxidant^[1], antidiabetic^[2], antiviral^[3], antiulcerogenic^[4], antinociceptive^[5], hepatoprotective^[6], antibacterial^[7], antimutagenic^[8], immunomodulatory^[9], cardioprotective^[10], anticancer^[11], antifungal^[12] radioprotective^[13] activities etc. In developing countries more than 80% peoples are dependent on medicinal plants which were estimated by world health organization (WHO).^[14] It is a well known fact that the demand for the ayurvedic drug treatment of various ailments is increasing day by day and plant drugs from the ayurvedic system are being explored more, not only in Bangladesh but also globally around the world. As a result, many research studies and findings are being undertaken and there is a need for an update and to put them together. That's why in this article an attempt has taken to recapitulate available pharmacological, phytochemical and biological studies for *Terminalia chebula*. This gives a wide knowledge about the ayurvedic plants and their importance in personal healthcare and hygiene products. It is considered a valuable source of unique natural products for development of drugs against various diseases and also for the development of industrial products.^[16] It is good to increase the appetite, as digestive aid liver stimulant, as stomachic, as gastrointestinal prokinetic agent and mild laxative. It stimulates the liver and perform by protecting expelling the waste excretory products from the intestines. It increases the frequency of stools, prevent aging, and provide immunity and body resistance against disease form.^[17] Active phytochemical constituents contain the triterpenes glucoside 1, arjungenin and the chebulosides 1&2. Other constituents contains tannins up to

30%, chebulic acid 3-5%, chebulinic acid 30%, tannic acid 20-40%, ellagic acid, 2,4-chebulyi- β -D-glucopyranose, gallic acid, ethyl gallate, punicalaginterflavin A, terchebin, anthraquinone, flavonoids like luteolin, rutins, and quercetin.^[18, 19,20] This phytochemical constituent's act as a various immunomodulatory functions in the human body with medicinal treatments.

2. TAXONOMY

Scientific name: *Terminalia chebula*

Bengali name: HORTOKI

Botanical description

Kingdom: Plantae

Division: Magnoliophyta

Class: Magnoliopsida

Order: Myrtales

Family: Combretaceae

Genus: *Terminalia*

Species: *chebula*



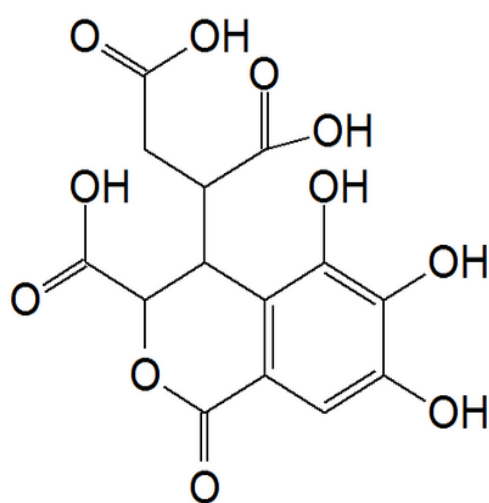
2.1 Botanical Description

The botanical description of *terminalia chebula* contains, it is a medium-sized, deciduous tree up to 25m tall, 60-80cm in diameter, crown rounded, spreading branches, dark brown branches with woody scales; leaves are thin- coriaceous, ovate, rounded at base, petiole up to 2cm long, 5-7cm long spikes flowers, 5 lobed calyx, absent corolla, 10 stamens, celled-1, ellipsoid drupe fruit, yellow to orange-brown when ripe.^[21-24]

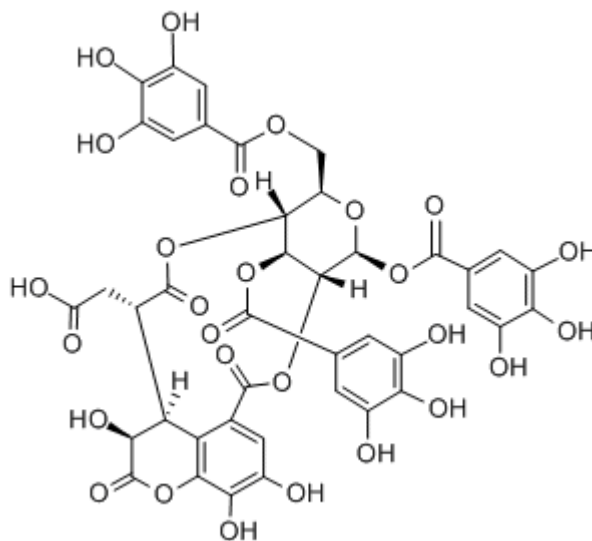
2.2 Phytochemistry of *Terminalia chebula*

Biologically active phytochemical constituents are includes chebulic acid^[25], chebulinic acid^[26], ellagic acid^[27], gallic acid^[27], chebulagic acid^[28], 1,6 di-*O*-galloyl- β -D-glucose, 3,4,6 tri-*O*-galloyl- β -D-glucose, 2,3,4,6 tetra-*O*-galloyl- β -D-glucose, 1,2,3,4,6 penta-*O* galloyl- β -D-glucose, ellagitannin contains punacalagin, casuarinin, corilagin, terchebulin, chebulanin, neochebulinic acid, chebulagic acid and phenolic compounds.^[30,31,32] High phenolic content, especially hydrolyzable tannins, anthraquinone, flavonol, carbohydrates, glucose and sorbitol^[29], with pharmacological studies from reverse phase chromatography there are some valuable active phytoconstituents has been reported including gallic acid, methyl gallate,

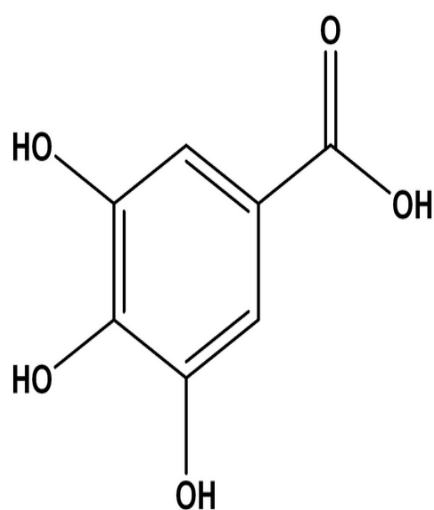
ethyl gallate, chebulagic acid, tetra-*O*-galloyl- β -D-glucose, ellagic acid, chebulinic acid and penta-*O* galloyl- β -D-glucose.^[33] It also contains nutrients such as vitamin C, protein, amino acids and minerals.^[34]



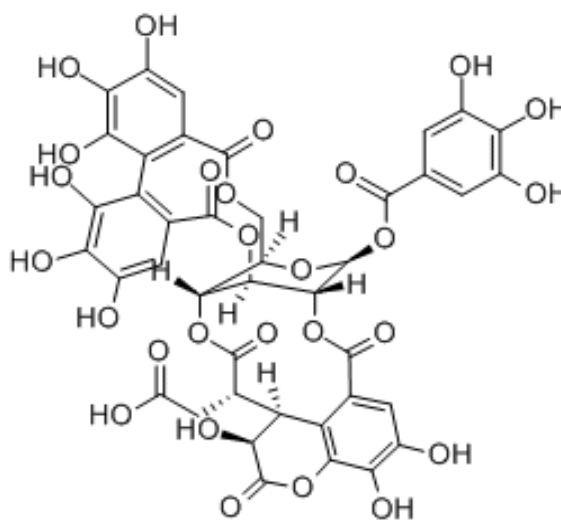
Chebulic acid



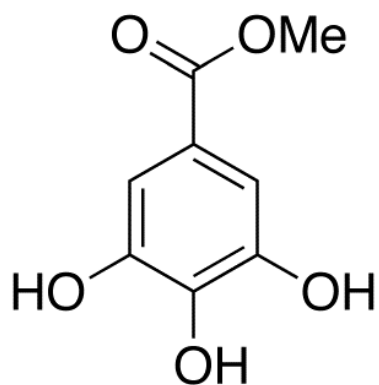
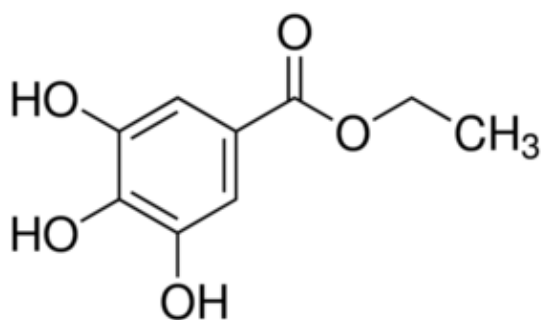
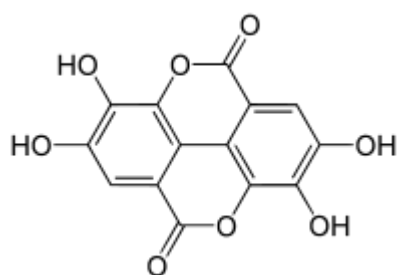
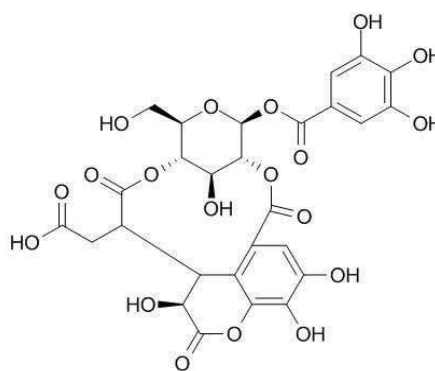
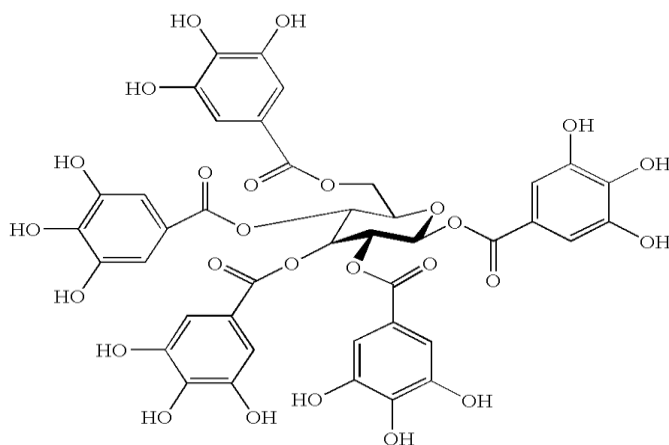
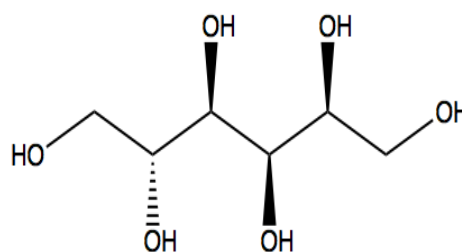
Chebulinic acid

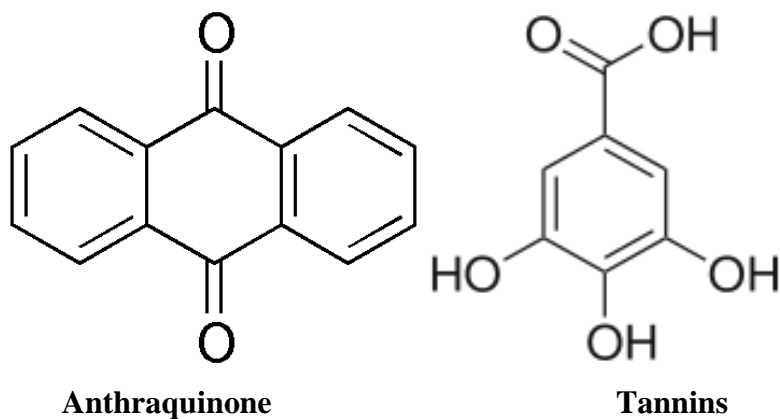


Gallic acid



Chebulagic acid

**Methyl gallate****Ethyl gallate****Ellagic acid****Chebunanin****Penta-O-galloyl-β-D-glucose****Sorbitol**

Figure1: phytochemical constituents of *Terminalia chebula*Table 1: pharmacological studies of *Terminalia chebula*

Serial	Pharmacological activity	Phytochemical compounds	Mode of Extraction	Organism	References
1	Anticancer	Chebolic acid	Methanol extract	<i>Salmonella typhi</i>	[7,25]
2	Antibacterial	Gallic acid	Ethanol extract, Ether, alcoholic, water extract	<i>Salmonella typhi</i> , <i>Staphylococcus aureus</i> , <i>Helicobacter pylori</i> , <i>Bacillus subtilis</i> etc.	[37,38,39,64]
3	Anticaries	chebulinic acid	Aqueous extract	<i>Streptococcus mutans</i>	[37,26, 40]
4	Anticonvulsant	ellagic acid	Ethanol, chloroform, Petroleum ether aqueous Extract	Rats	[37,27, 41]
5	Antidiabetic	chebulagic acid	Chloroform extract, Ethanol extract	Streptozotocin induced Diabetic rats, Adult albino male rats	[28,37,42,43]
6	Antifungal	1,6 di- <i>O</i> -galloyl- β -D-glucose	Aqueous, alcoholic, Ethyl acetate extract, 70% of methanol, Ethylacetate, hexane, Chloroform Extract	<i>Aspergillus niger</i> , <i>Aspergillus flavus</i> ,	[30,37,44,45]
7	Antimutagenic	penta- <i>O</i> galloyl- β -D-glucose	Acetone, aqueous chloroform extract Chloroform, aqueous Extract	<i>Salmonella typhimurium</i>	[33,37,46,47]
8	Antioxidant	Sorbitol	Water, methanol & 95% of ethanol extract	Fermented products Adult male albino rats	[29,37,48,49]
9	Antiulcer	Anthraquinone	Methanolic extract	Wistar albino male rats	[29,37,50]

10	Antiviral	Chebullanin	Acetone extract, Aqueous extract	Swine influenza A virus, Hepatitis B virus	[30,37,51,52]
11	Cardio protective	phenolic compounds	95% of ethanol extract	Adult albino male rats	[30,31,37,53]
12	Cytotoxic	chebulagic acid and	Acetone extract	Cancer cell lines	[31,32,37,54]
13	Immunodulatory	neochebulinic acid	Alcohol extract	Male wistar rats	[31,32,37,55]
14	Radioprotective	Punacalagin	Aqueous extract	Rats	[32,37,56]
15	Wound healing	Casurarinin	Hydroalcoholic extract, 90% of ethanol extract	Induced diabetic rats, Wistar albino rats	31,[37,57,58]
16	Dermal Wounds	Corilagin	Dry powder mixed With water	Rabbit	[30,32,59,60]
17	Anti-hyperglycemic Effect	Terchebulin	Water extract of dry fruits	Diabetic rats	[32,59,61]
18	Anticlastogenic Effect		Methanolic Extracts	Mouse bone marrow cells	[59,62]
19	Typhoid Fever		Aqueous extract		[59,63]
20	Antiplasmodial activity		Acetone seed extract		[64]
21	Inhibits free radical induced hemolysis		Aqueous extract		[65]
22	Xanthine/xanthine oxidase inhibition, 2,2-diphenyl-1- picrylhydrazyl (DPPH) radicals scavenging acitivity		Aqueous extract		[66]
23	Stronger antioxidant activity than alpha		Acetone extract		[67]
24	Gastrointestinal motility improving, Increase gastric emptying time, Protection against duodenal ulcer		Fruit extract		[68], [69]
25	Anti-oxidative and membrane stabilizing activities		(95% ethanolic extract		[70]
26	Reduces irradiation effects, Breaks Gamma radiation induced strand in		Aqueous extract)	Mice	[71], [72]

	plasmid PBR322 DNA				
27	Renoprotective activity, Reduction in blood glucose		Fruit, seed	Rats	[73], [74]
28	Hypocholesterolemi c activity, Induced atherosclerosis		Aqueous extract		[75], [76]
29	Cytoprotective activity, Development of duodenal ulcers, Inhibitory effect on cellular aging				[77], [78]
30	Free radical scavenging activity, Inhibited oxidative stress		Ethanol extract		[79]
31	Anti-microbial activity		Methanol extract		[80]
32	Radio Protecting Ability and Phytochemical analysis		Aqueous extract		[81]
33	Antinociceptive activity		Petroleum ether (PE), chloroform (CH), ethanol (ETH) and water Extracts		[82]
34	The Molluscicida L activity		Ethanol extract		[83]
35	Spasmogenic Activity		Aqueous extract		[84]
36	Hepatoprotective Activity		Leaf powder Mixed with 1% Gum accai Suapension		[85]
37	Inhibition of HIV 1 Integrase		Hot water Extract		[86]
38	Antidiabetic And renoprotective		Chloroform Extract		[87]

39	Biochemical Studies		Ethanol extract		[88]
40	Hepatocellular carcinoma		Aqueous extract		[89]
41	Anti lithiatic Activity		Aqueous extract		[90]
42	Anti-aging Activities		Methanol extract		[91]
43	Potent Sources of natural antioxidant.		Methanol extract		[92]
44	Using DPPH, deoxyribose, reducing power, chelating power		Hexane extracts		[93]
45	Exhibit antioxidant activity at different magnitude of potency		Warm water Extract		[94]
46	Improves glucose tolerance and brings down Fasting blood Glucose in		Water extract of Dry fruits	Rats	[95]
47	Against multi drug resistant diabetic foot ulcer isolates.		Methanol, Isopropanol, Chloroform, Diethyl ether and Hexane		[96]
48	Against gram-positive Bacteria than against gram-negative bacteria.		Aqueous extracts		[97]
49	Potential bactericidal Activity.		Methanol, Ethanol, ethyl Acetate water and Chloroform extract Of leaf		[98]
50	The antibacterial activity		Ethanol extract	<i>Salmonella typhi</i> , <i>Staphylococcus aureus</i> , <i>Bacillus subtilis</i> Etc.	[99]
51	Alkaloids from all plant Parts showed good antimicrobial activity		Alkaloids Extracted from Different parts (leaf, stem, stem Bark, and fruits)		[100]
52	Potential bactericidal and		Methanol and Aqueous extracts		[101]

	potent Antioxidant				
53	Significant response in wound Types		Ethanolic extract		[102]
54	Dry powder mixed With water		Significant improvement On Maturation	Rabbit	[103]
55	Powerful anti-bacterial and angiogenic Activity		Extracted with Warm water		[104]
56	Acceleration of the Healing process		Alcoholic extract Of the leaves		[105]
57	Performed against common pathogenic bacterial and fungal strains		Crude ethyl Acetate and ether Extract	<i>Staphylococcus aureus, proteus vulgaris</i> and <i>Escherichia coli</i>	[106]

2.3 pharmacological and biological uses

Terminalia chebula is called “The king of medicines” because of its high content of alkaloids, secondary metabolites, flavonoids and the astonishing power of healing with a wide range of biological and pharmacological uses.^[16] Important biological uses by this plant includes antibacterial, antifungal, antiviral, antimutagenic, adaptogenic, anti-anaphylactic, hypocholesterolemic, gastrointestinal motility improving, anti-ulcerogenic, hepatoprotective, cardioprotective, radioprotective, antidiabetic and retinoprotective, antispasmodic, wound healing, purgative, immunomodulatory and chemopreventive activities.^[16] Gallic acid acts as an anti-inflammatory response binding with receptors.^[35] It is used as a blood purifier.^[36] Several pharmacological investigations for different biological activities of *Terminalia chebula* in various in vivo and in vitro test models have been carried out based on the presence of chemical ingredients. A summary of the findings of some of these pharmacological studies is presented below in Table 1.

CONCLUSION

We are now living in a modern era as a result medical science are developing day by day inspite of this a large segment of the world population still now depends on the plants origin medicine. *Terminalia chebula* is one of the world most valuable ayurvedic plants having a wide variety of pharmacological and medical activities. From the ancient time, plants have been widely used as curative agents for variety of ailments. *Terminalia chebula* serves as a great source of a variety of biologically active phytoconstituents as for example chebulic acid, chebulinic acid, gallic acid, chebulagic acid and other related compounds that

compounds are result to antimicrobial, antitoxidant, antihyperglycemic, anticancer, and protective effects on various vital organs those are includes nervous, heart, kidney, liver. To treat a large number variety of health problems *Terminalia chebula* plant is generally used. By showing the biodiversity of both nutritional as well as medicinal components *Terminalia chebula* is known as the root of medicine. Day by day the investigation on medicinal plants are rising so fast as a result herbal products are becoming safe and effective to the people . We wish this work will help to create awareness about medicinal plant research and their future possibility.

ACKNOWLEDGEMENT

We cordially thankful to all teachers from the Department of Biotechnology and Genetic Engineering (BGE), Faculty of Life Science, Mawlana Bhashani Science and Technology University (MBSTU), Tangail-1902, Bangladesh; For their valuable suggestions and inspiration during our Review article proceedings.

REFERENCES

1. Suchalatha S and Devi CS. Antioxidant activity of ethanolic extract of *Terminalia chebula* fruit against isoproterenol – induced oxidative stress in rats. Indian Journal of Biochemistry and Biophysics., 2005; 42: 246-249.
2. Rao NK and Nammi S. Antidiabetic and renoprotective effects of the chloroform extract of *Terminalia chebula* Retz. Seeds in streptozotocin-induced diabetic rats, BMC Complement Altern Med., 2006; 6: 17.
3. Kim TG, Kang SY, Jung KK, Kang JH, Lee E, Han HM and Kim SH. Antiviral activities of extracts isolated from *Terminalia chebula* Retz., *Sanguisorba officinalis* L., *Rubus coreanus* Miq and *Rheum palmatum* L. against hepatitis B virus, Phytotherapy research., 2001; 15(8): 718-720.
4. Sharma P, Prakash T, Kotresha D, Ansari MA, Sahrm UR, Kumar B, Debnath J, Goli D. Antiulcerogenic activity of *Terminalia chebula* fruit in experimentally induced ulcer in rats. Pharm Biol., 2011; 49(3): 262-68.
5. Sarabjit Kaur & R K Jaggi, Antinociceptive activity of chronic administration of different extracts of *Terminalia bellerica* Roxb. and *Terminalia chebula* Retz. Fruits, Indian Journal of Experimental Biology, Vol. 48, September 2010; 925-930.

6. S.Vidya et al., Hepato-Protective Activity Of *Terminalia chebula* Leaves In Paracetamol Induced Hepato-Toxicity In Rats, International Journal of Advances in Pharmaceutical Research, 2011, 2(4), 127 – 132.
7. Kannan P, Ramadevi SR and Waheeta Hopper. Antibacterial activity of *Terminalia chebula* fruit extract, African Journal of Microbiology Research. 2009; 3(4): 180-184.
8. Grover IS and Bala S; Antimutagenic activity of *Terminalia chebula* (myroblan) in *Salmonella typhimurium*. Indian Journal of Experimental Biology, 1992; 30(4): 339-341.
9. Aher V, Wahi AK; Immunomodulatory activity of alcohol extracts of *Terminalia chebula* Retz combretaceae. Tropical Journal of Pharmaceutical Research, 2011; 10(5): 567-575.
10. Suchalatha S, Devi CS; Protective effect of *Terminalia chebula* against lysosomal enzyme alterations in isoproterenol induced cardiac damage in rats. Experimental clinical cardiology, 2005; 10(2): 91-95.
11. Saleem A, Husheem M, Harkonen P and Pihlaja K; Inhibition of cancer cell growth by crude extract and the phenolics of *Terminalia chebula* Retz fruit. Journal of Ethnopharmacology, 2002; 81(3): 327- 336.
12. Saheb S, More SM, Junne SB, Wadje SS; The Antifungal activity of five *Terminalia* species checked by paper disk method. International Journal of Pharma Research and Development, 2011; 3(2).
13. Jagetia GC, Baliga MS, Malagi KJ, Sethukumar KM; The evaluation of the radioprotective effect of Triphala (an ayurvedic rejuvenating drug) in the mice exposed to g-radiation. Phytomedicin, 2002; 9(2): 99–108.
14. Sharma M, Pandey Govind P; Ethanomedicinal Plants for Prevention and treatment of tumors. International Journal of Green pharmacy, 2009; 3(1): 2-5.
15. Prakash DV, SreeSatya N, Sumanjali Avanigadda S, Vangalapati M; Pharmacological review on *Terminalia chebula*. International Journal of Research in Pharmaceutical and Biomedical Sciences., 2012; 3(2): 679-683.
16. Chattopadhyay RR, Bhattacharyya SK (2007). Plant Review *Terminalia chebula*. Pharmacognos. Rev., 2007; 23: 145-150.
17. Vaibhav Aher and Arun Kumar Wahi. Immunomodulatory activity of alcohol extracts of *Terminalia chebula* Retz combretaceae, Tropical Journal of Pharmaceutical Research., 2011; 10(5): 567-575.
18. Kumar KJ. Effect of geographical variation on contents of tannic acid, gallic acid, chebulinic acid and ethyl gallate in *Terminalia chebula*. Natural Products., 2006; 2(3-4): 170-75.

19. Juang LJ, Sheu SJ, Lin TC. Determination of hydrolyzable tannins in the fruit of *Terminalia chebula* Retz. By high-performance liquid chromatography and capillary electrophoresis. *J Sep Sci.*, 2004; 27(9): 718–24.
20. Srivastava A, Chandra A, Singh M, Jamal F, Rastogi P, Rajendran SM, Bansode FW, Lakshmi V. Inhibition of hyaluronidase activity of human and rat spermatozoa *in vitro* and antispermatogenic activity in rats *in vivo* by *Terminalia chebula*, a flavonoid rich plant. *Reproductive Toxicol.*, 2010; 29: 214–24.
21. <http://ayurvista.blogspot.in/search?q=terminalia>
22. Boer, E., et al. 1995. *Terminalia* L. In Lemmens, R.H.M.J., Soerianegara, I. & Wong, W.C. (Eds.): *Plant Resources of South-East Asia*. No. 5(2): Timber tree: Minor commercial timber. Prosea Foundation, Bogor, Indonesia., 475-478., 483.
23. Fundter, J.M., et al. *Terminalia chebula* Retz. In Lemmens, R.H.M.J. & Wulijarni-Soetjipto, N. (Eds.): *Plant Resources of South-East Asia*. No. 3: Dye and tannin-producing plants. Prosea Foundation, Bogor, Indonesia., 1992; 122-125.
24. Prakash Chandra Gupta, Biological and Pharmacological Properties of *Terminalia chebula* Retz. Haritaki)- An Overview, *International Journal Of Pharmacy And Pharmaceutical Sciences.*, 2012; 4(3).
25. Lee HS, Koo YC, Suh HJ, Kim KY and Lee KW; Preventive effects of chebulic acid isolated from *Terminalia chebula* on advanced glycation endproduct-induced endothelial cell dysfunction. *Journal of Ethnopharmacol.*, 2010; 131(3): 567-574.
26. Quanbin Han, Jingzheng Song, Chunfeng Qiao, Lina Wong and Hongxi Xu; Preparative isolation of hydrolysable tannins chebulagic acid and chebulinic acid from *Terminalia chebula* by high-speed counter-current chromatography. *J. Sep. Sci.*, 2006; 29: 1653-1657.
27. Patel Madhavi G, Patel Vishal R, Patel Rakesh K; Development and Validation of Improved RP-HPLC method for Identification and Estimation of Ellagic and Gallic acid in Triphalachurna. *International Journal of ChemTech Research*, 2010; 2(3): 1486-1493.
28. Bharat Reddy D, Reddy TCM, Jyotsna G, Sharan S, , Priya N, Lakshmi pathi V , Reddanna P; Chebulagic acid a COX–LOX dual inhibitor isolated from the fruits of *Terminalia chebula* Retz induces apoptosis in COLO-205 cell line. *Journal of Ethnopharmacology*, 2009; 124(3): 506-512.
29. Creencia E, Eguchi T, Nishimura T, Kakinuma K; Isolation and structure elucidation of the biologically active components of *Terminalia chebula* Retz (Combretaceae). *KIMIKA*, 1966; 12: 1-10.
30. Juang LJ, Sheu SJ, Lin TC, *J. Sep. Sci.*, 2004; 27: 718-724.

31. Han Q, Song J, Qiao C, Wong L, Xu H. *J.Sep. Sci.*, 2006; 29: 1653-1657.
32. Chattopadhyay RR, Bhattacharyya SK. Plant Review *Terminalia chebula*. Pharmacognos. Rev., 2007; 23: 145-15.
33. Anil Mahajan, NandiniPai, Simultaneous isolation and identification of phytoconstituents from *Terminalia chebula* by preparative chromatography, *J. Chem. Pharm. Res.*, 2010; 2(5): 97-103
34. Mahesh R, Ramesh T, Nagulendran KR, Velavan S, Hazeena BV. Effect of *Terminalia chebula* on Monoamine Oxidase and Antioxidant enzyme activities in aged rat brain. Pharmacognos. Mag., 2007; 3: 12 16.
35. Feng-lin H, Li-ming Y, Shwu-fen C, Li-hsuan W, Chung-yi H, Pan-chun- L, Shwu-jiuan L. Biotransformation of Gallic Acid by *BeauveriaSulfurescens*. Appl. Microbial. Biotechnol., 2007; 74: 659-666.
36. Thomas J, Joy PP, Mathew G, Skaria S, Duethi BP, Joseph TS. Agronomic practices for aromatic and medicinal plant, Directorate of arecanut and spices Development India. Calicut, Kerala, India, 2000; 124-128.
37. Dr S. Aruna, Dr L.V. Nandakishore;Haritaki A Boon To Herbalism – A Review, Scholars Academic Journal of Biosciences (SAJB), 2014; 2(2): 132-136
38. Kannan P, Ramadevi SR, Hopper W; Antibacterial activity of *Terminalia chebula* fruit extract”, African Journal of Microbiology Research, 2009; 3(4): 180-184.
39. Malekzadeh F, Ehsanifar H, Shahamat M, Levin M, Colwell RR; Antibacterial activity of black myrobalan (*Terminalia chebula* Retz) against *Helicobacter pylori*. International Journal of Antimicrobial Agents, 2001; 18(1): 85–88.
40. Jagtap AG and Karkera SG; Potential of the aqueous extract of *Terminalia chebula* as an anticaries agent. Journal of Ethnopharmacology, 1999; 68(1-3): 299–306.
41. Hoga d eMaheshwar G, Deshpande S.V and PramodH.J. Anticonsultant activity of *Terminalia chebula* Retz in rats, Journal of Herbal Medicine and Toxicology. 2010; 4(2): 123- 126.
42. Gandhipuram P, Kumar S, Arulselvan P, Kumar DS, Subramanian SP; Anti Diabetic activity of fruits of *Terninalia chebula* on Streptozotocin induced Diabetic rats. Journal of Health Science, 2006; 52(3): 283-291.
43. Rao NK, Nammi S; Antidiabetic and renoprotective effects of the chloroform extract of *Terminalia chebula* Retz. Seeds in streptozotocin-induced diabetic rats. BMC Complement Altern Med, 2006; 6: 17.

44. Saheb S, More SM, Junne SB, Wadje SS; The Antifungal activity of five *Terminalia* species checked by paper disk method. International Journal of Pharma Research and Development, 2011; 3(2).
45. Vivek KB, Rahman A, Shukla S, Shukla S, YassirArafat SM, Hossain AM, Mehta A; In vitro kinetics and antifungal activity of various extracts of *Terminalia chebula* seeds against plant pathogenic fungi. Archives of Phytopathology and Plant Protection, 2010; 43(8): 801-809.
46. Grover IS and Bala S; Antimutagenic activity of *Terminalia chebula* (myroblan) in *Salmonella typhimurium*. Indian Journal of Experimental Biology, 1992; 30(4): 339-341.
47. Kaur S, Arora S, Kaur K, Kumar S; The in vitro antimutagenic activity of Triphala an Indian herbal drug. Food Chem Toxicol, 2002; 40(4): 527-34.
48. Suchalatha S, Devi CS; Antioxidant activity of ethanolic extract of *Terminalia chebula* fruit against isoproterenol – induced oxidative stress in rats. Indian Journal of Biochemistry and Biophysics, 2005; 42: 246-249.
49. Chia-Lin Chang and Che-San Lin; Development of antioxidant activity and pattern recognition extracts and its fermented products, Research Institute of Biotechnology, Hungkuang University., 2010.
50. Raju D, Ilango K, Chitra V, Ashish K; Evaluation of Anti-ulcer activity of methanolic extract of *Terminalia chebula* fruits in experimental rats. Journal of Pharmaceutical Science and Research, 2009; 1(3): 101-107.
51. Hongbo Ma, Yunpeng Diao, Danyu Zhao, Kun Li and Tingguo Kang; A new alternative to treat swine influenza A virus infection: extracts from *Terminalia chebula* Retz. African Journal of Microbiology Research, 2010; 4(6): 497- 499.
52. Kim TG, Kang SY, Jung KK, Kang JH, Lee E, Han HM, Kim SH; Antiviral activities of extracts isolated from *Terminalia chebula* Retz., *Sanguisorba officinalis* L., *Rubus coreanus* Miq and *Rheum palmatum* L. against hepatitis B virus. Phytotherapy research, 2001; 15(8): 718-720.
53. Suchalatha S, Devi CS; Protective effect of *Terminalia chebula* against lysosomal enzyme alterations in isoproterenol induced cardiac damage in rats. Experimental clinical cardiology, 2005; 10(2): 91-95.
54. Kaur S, Michael H, Arora S, Harkonen PL, Kumar S; The in vitro cytotoxic and apoptotic activity of Triphala--an Indian herbal drug. Journal of Ethnopharmacology, 2005; 97(1): 15–20.

55. Aher V, Wahi AK; Immunomodulatory activity of alcohol extracts of *Terminalia chebula* Retz combretaceae. Tropical Journal of Pharmaceutical Research, 2011; 10(5): 567-575.
56. Jagetia GC, Baliga MS, Malagi KJ, Sethukumar KM; The evaluation of the radioprotective effect of Triphala (an ayurvedic rejuvenating drug) in the mice exposed to g-radiation. Phytomedicin, 2002; 9(2): 99–108.
57. Singh MP, Sharma CS; Wound healing activity of *Terminalia chebula* in experimentally induced diabetic rats. International Journal of Pharm Tech Research, 2009; 1(4): 1267-1270.
58. Choudhary GP; Wound healing activity of ethanolic extract of *Terminalia chebula* retz. International Journal of Pharma and Bio Sciences, 2011; 2(1): 48-52.
59. R. Rathinamoorthy and G. Thilagavathi; *Terminalia chebula* - Review on Pharmacological and Biochemical Studies, International Journal of PharmTech Research., Jan-March 2014; 6(1): 97-116.
60. Shreedevi M.S and sampathkumar.B, Effect of siddha herbal paste on wound healing in induced dermal wounds in rat, International journal of Ayurvedic and Herbal Medicine, 2011; 1(3): 86-91.
61. Y. K. Murali, Ramesh Chandra and P.S. Murthy, Antihyperglycemic Effect of Water Extract of Dry Fruits of *Terminalia chebula* in Experimental Diabetes Mellitus, Indian Journal of Clinical Biochemistry, 2004; 19(2): 202-204.
62. Wasim Raja, Sonam Pandey and R.C. Agrawal, Studies on the Anticlastogenic Effect of *Terminalia chebula* extract on Cyclophosphamide-Induced Micronucleus Formation and Chromosome Aberrations in Swiss albino Mice, International Journal of Genetics, 2011; 1(2): 13-17.
63. Khan KH, Jain SK. Regular intake of *Terminalia chebula* can reduce the risk of getting typhoid fever. Advanced Biotech., 2009; 8: 10-15.
64. Prakash chandra gupta, biological and pharmacological properties of *terminalia chebula* retz. (haritaki)- an overview; international journal of pharmacy and pharmaceutical sciences., 2012; 4-3.
65. Mahesh R, Bhuvana S, Begum VM. Effect of *Terminalia chebula* aqueous extract on oxidative stress and antioxidant status in the liver and kidney of young and aged rats. Cell Biochem Funct., 2009; 27(6): 358–363.
66. Naik GH, Priyadarsini KI, Naik DB, Gangabhairathi R, Mohan H. Studies on the aqueous extract of *Terminalia chebula* as a potent antioxidant and a probable radioprotector. Phytomedicine., 2004; 11(6): 530–538.

67. Chen X, Sun F, Ma L, Wang J, Qin H, Du G. *In vitro* evaluation on the antioxidant capacity of triethylchebulate, an aglycone from *Terminalia chebula* Retz fruit. Indian J Pharmacol., 2011; 43(3): 320–3
68. Tamhane MD, Thorate SP, Rege NN, Dahanukar SA. Effect of oral administration of *Terminalia chebula* on gastric emptying: An experimental study. J Postgrad Med., 1997; 43(1): 12–13.
69. Sharma P, Prakash T, Kotresha D, Ansari MA, Sahrm UR, Kumar B, et al. et al. Antiulcerogenic activity of *Terminalia chebula* fruit in experimentally induced ulcer in rats. Pharm Biol., 2011; 49(3): 262–268.
70. Tasduq SA, Singh K, Satti NK, Gupta DK, Suri KA. *Terminalia chebula* (fruit) prevents liver toxicity caused by sub-chronic administration of rifampicin, isoniazid and pyrazinamide in combination. Hum Exp Toxicol., 2006; 25: 111-18.
71. Naik GH, Priyadarsini KI, Naik DB, Gangabthagirathi R, Mohan H. Studies on the aqueous extract of *Terminalia chebula* as a potent antioxidant and a probable radioprotector. Phytomedicine., 2004; 11(6): 530–538.
72. Gandhi NM, Nayar CKK. Radiation protection by *Terminalia chebula* some mechanistic aspects. Mol Cell Biochem., 2005; 277(1–2): 43–48
73. Kannan VR, Rajasekar GS, Rajesh P, Balasubramanian V, Ramesh N, Solomon EK, et al. et al. Anti-diabetic activity on ethanolic extracts of fruits of *Terminalia chebula* Retz. Alloxan induced diabetic rats. Am J Drug Discov Dev., 2012; 2: 135–142.
74. Senthilkumar GP, Subramanian SP. Biochemical studies on the effect of *Terminalia chebula* on the levels of glycoproteins in streptozotocin-induced experimental diabetes in rats. J Appl Biomed., 2008; 6: 105–115.
75. Maruthappan V, Shree KS. Hypolipidemic activity of Haritaki (*Terminalia chebula*) in atherogenic diet induced hyperlipidemic rats. J Adv Pharm Tech Res., 2010; 1: 229–235
76. Israni DA, Patel KV, Gandhi TR. Anti-hyperlipidemic activity of aqueous extract of *Terminalia chebula* and Gaumutra in high cholesterol diet fed rats. Int J Pharm Sci., 2010; 1(1): 48–59.
77. Lee HS, Koo YC, Suh HJ, Kim KY, Lee KW. Preventive effects of chebulic acid isolated from *Terminalia chebula* on advanced glycation endproduct-induced endothelial cell dysfunction. J Ethnopharmacol., 2010; 131(3): 567–574.
78. Na M, Bae M, Keng SS, Min BS, Yoo JK, Kamiryo Y, et al. et al. Cytoprotective effect on oxidative stress and inhibitory effect on cellular aging of *Terminalia chebula* fruit. Phytother Res., 2004; 18(9): 737–741.

79. Na MK, Bae KH, Kang SS, Min BS, Yoo JK, Kamiryo Y, Senoo YI, Yokoo S, Miwa N. Cytoprotective effect on oxidative stress and inhibitory effect on cellular aging of *Terminalia chebula* fruit. *Phytotherapy Res.*, 2004; 18(9): 737–41.
80. Rathinamoorthy. R, Udayakumar S and Thilagavathi G, Antimicrobial Efficacy of *Terminalia Chebula* Fruit Extract Treated Cotton Fabric For Healthcare Applications, *International Journal of Pharmaceutical Sciences and Nanotechnology*, 2012; 4: 1549-1556.
81. Naik GH, Priyadarsini KI, Naik DB, Gangabhagirathi R, Mohan H. Studies on the aqueous extract of *Terminalia chebula* as a potent antioxidant and a probable radioprotector. *Phytomedicine.*, 2004; 11: 530-38.
82. Sarabjit Kaur & R K Jaggi, Antinociceptive activity of chronic administration of different extracts of *Terminalia bellerica Roxb. and Terminalia chebula* Retz. *Fruits*, *Indian Journal of Experimental Biology*, Vol. 48, September 2010; 925-930.
83. Upadhyay A, Singh DK. Molluscicidal activity of *Sapindus mukorossi* and *Terminalia chebula* against the freshwater snail *Lymnaea acuminata*. *Chemosphere.*, 2011; 83(4): 468-74.
84. Seyyed Ali Mard et al, Spasmogenic Activity of the Seed of *Terminalia chebula* Retz in Rat Small intestine: In Vivo and In Vitro Studies, *Malaysian J Med Sci.*, 2011; 18(3): 18-26.
85. S.Vidya et al., Hepato-Protective Activity Of *Terminalia chebula* Leaves In Paracetamol Induced Hepato-Toxicity In Rats, *International Journal of Advances in Pharmaceutical Research*, 2011; 2(4): 127 – 132.
86. Ahn MJ, Kim CY, Lee JS, Kim TG, Kim SH, Lee CK, Lee BB, Shin CG, Huh H, Kim J. Inhibition of HIV-1 integrase by galloyl glucoses from *Terminalia chebula* and flavonol glycoside gallates from *Euphorbia pekinensis*. *Planta Med*, 2002; 68(5): 457-59.
87. Nalamolu Koteswara Rao and Srinivas Nammi, Antidiabetic and renoprotective effects of the chloroform extract of *Terminalia chebula* Retz. seeds in streptozotocin-induced diabetic Rats, *BMC Complementary and Alternative Medicine.*, 2006; 6: 17.
88. Gandhipuram Periasamy Senthilkumar, Sorimuthu Pillai Subramanian, Biochemical studies on the effect of *Terminalia chebula* on the levels of glycoproteins in streptozotocin-induced experimental diabetes in rats, *J. Appl. Biomed.*, 2008; 6: 105–115.
89. Srisesharam Srigopalram, Indira A Jayraaj, Effect of *Terminalia Chebula* Retz On Den Induced Hepatocellular Carcinogenesis In Experimental

90. Tayal, et al. Cytoprotective role of the aqueous extract of *Terminalia chebula* on renal epithelial cells, IBJU, 2012; 38(2): 204-214.
91. Manosroi A, Jantrawut P, Akihisa T, Manosroi W, Manosroi J. In vitro anti-aging activities of *Terminalia chebula* gall extract. Pharmaceutical Biology., 2010; 48: 469- 481.
92. Bibhabasu Hazra et al, RCesoeamrchp Artaicrleative study of the antioxidant and reactive oxygen species scavenging properties in the extracts of the fruits of *Terminalia chebula*, *Terminalia belerica* and *Embllica officinalis*, BMC Complementary and Alternative Medicine 2010; 10: 20.
93. Harpreet Walia, Subodh Kumar and Saroj Arora, Comparative antioxidant analysis of hexane extracts of *Terminalia chebula* Retz. prepared by maceration and sequential extraction method, Journal of Medicinal Plants Research, 2011; 5(13): 2608-2616.
94. Hua-Yew CHENG,et al, Antioxidant and Free Radical Scavenging Activities of *Terminalia chebula*, Biol. Pharm. Bull., 2003; 26(9): 1331—1335.
95. Y. K. Murali, Ramesh Chandra and P.S. Murthy, Antihyperglycemic Effect of Water Extract of Dry Fruits of *Terminalia Chebula* In Experimental Diabetes Mellitus, Indian Journal of Clinical Biochemistry, 2004; 19(2): 202-204.
96. Arumugam Suresh et al, Screening of Antibacterial Properties of Indian Medicinal Plants Against Multi Drug Resistant Diabetic Foot Ulcer Isolates, International Journal Of Phytopharmacology., 2012; 3(2): 139-146.
97. Maheshwar G. Hogade, Sunil Jalalpure, Sonali Kuthar, International Journal of Pharmacy and Pharmaceutical Science Research, 2011; 1(1): 26-29.
98. Golam Mostafa, M, Mahdia Rahman, M. Manjurul Karim, Antimicrobial Activity of *Terminalia Chebula*, Int. J. Med. Arom. Plants, 2011; 1(2): 175-179.
99. Kannan P, Ramadevi SR, Hopper W. Antibacterial activity of *Terminalia chebula* fruit extract. African J Microbiol Res, 2009; 3(4): 180-84.
100. Geeta singh, padma kumar Evaluation of antimicrobial activity of alkaloids of *Terminalia chebula* Retz. against some multidrug resistant Microorganisms, International Journal of Green Pharmacy, 2012; 57-62.
101. Dolly Singh, Therapeutical Effect of Extracts of *Terminalia chebula* In Inhibiting Human Pathogens and free radicals, International Journal of Bioscience, Biochemistry and Bioinformatics, 2012; 2(3).
102. Choudhary G.P., Wound healing activity of the ethanolic extract of *Terminalia chebula* retz., International Journal of Pharma and Bio Sciences, 2011; 2(1): 48-52

103. Fundter, J.M., et al. *Terminalia chebula* Retz. In Lemmens, R.H.M.J. & Wulijarni-Soetjipto, N. (Eds.): Plant Resources of South-East Asia. No. 3: Dye and tannin-producing plants. Prosea Foundation, Bogor, Indonesia. 1992; 122-125.
104. Kun Li et al, Tannin extracts from immature fruits of *Terminalia chebula* Fructus Retz. Promote cutaneous wound healing in rats BMC Complementary and Alternative Medicine., 2011; 11: 86.
105. Sugun L, Sing S, Sivakuma P, Sampat P, Chandrakasa G. Influence of *Terminalia chebula* on dermal wound healing in rats. Phytotherapy Res., 2002; 16(3): 227-31.
106. Archana Sharma, Suchitra Meena and Nachiketa Barman, Efficacy of ethyl acetate and ether extract of *Terminalia chebula* Retz against some human pathogenic strains, International Journal of PharmTech Research, 2011, 3(2): 724-727.