

**Review Article** 

# A Descriptive Review on Traditional Herbal Drug-Terminalia Chebula

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# ABSTRACT

Terminalia chebula is an important medicinal plant in pharmaceutics and used in Unani System of Medicine (USM) from ancient time to combat various diseases and infections due to its potential medicinal use. This herb has been considered the valuable and cheap source of unique phyto-constituents (tannins, alkaloids, phytochemicals) which are used widely in the development of drugs with higher safety margins and lesser side effects against different types of diseases. It is commonly called Haritaki and belongs to the family Combretaceae. It is known as black myroban, Harad and Halaila in English, Hindi and Persian respectively. In Tibet, it is known as "king of the medicines". This herb has a unique mention in various traditional medicine systems due to its amazing power of healing. Unani physicians have been using this drug as a brain tonic, eye tonic, cardiotonic, stomach tonic, kidney tonic, blood purifier, astringent, purgative (bile), cholagogue, melanogogue, carminative, digestive anthelmintic, anti-inflammatory and antidysenteric agent. Hence this drug having a momentous position in Unani system of medicine and text.

The previous pharmacological studies showed that T. chebula possesses antibacterial, antifungal, antiviral, antidiabetic, antimutagenic, antioxidant, antiulcer and wound healing properties. It also prevents cardiac damage and is used for the treatment of kidney disease. T. chebula and its phytoconstituents have a therapeutic effect with no toxicity. This review was designed to lime light the T. chebula by describing its traditional, therapeutic and others uses.

**Keywords:** Anti-bacterial, Halaila, Medicinal Application, Phytoconstituents, Terminalia Chebula

# Introduction

Terminalia chebula (Halaila) is an important medicinal plant used in Unani System of Medicine (USM) from the dawn of civilization to combat various diseases and have been considered the precious and economical source of unique phytoconstituents which are utilized widely in the development of drugs against different diseases.<sup>1</sup> It is a flowering evergreen tree, belonging to the family Combretaceae. It has several common names such as Black myrobalan, ink tree or chebulic myrobalan and also known as "King of medicine" in Tibet. It is extensively exercised herb of USM not only used in India but also in



**Copyright (c) 2019 Journal of Advanced Research in BioChemistry and Pharmacology** https://www.adrpublications.in many other regions of Asia and Africa <sup>2</sup>. It is beneficial in cancer, paralysis, cardiovascular disease, leprosy, gout, epilepsy, ulcer, rheumatoid arthritis, piles, gastric disorders etc.<sup>2,3,4</sup> The pharmacological studies showed that T. chebula possesses antibacterial, antifungal, antiviral, antidiabetic, antimutagenic, antioxidant, antiulcer and wound healing properties etc. it is used widely in the formation of many extremely valuable Unani formulations to treat the many infectious diseases likes bleeding and ulceration of gums, conjunctivitis, diarrhea, dysentery, fungal infection of skin.<sup>2,5,6,7</sup> The fruit and bark are the main parts used for medicinal purpose in USM.<sup>2,6</sup> It strengthens the brain, eye, gums, prevent ageing and also increase body resistance against disease.<sup>2,5,6</sup>

#### **Botanical Description<sup>8</sup>**

#### Taxonomy

Kingdom: Plantae

Division: Mangnoliophyta

Class: Magnoliopsida

Order: Myrtales

Family: Combretaceae

Genus: Terminalia

Species: chebula

#### Habit and Habitat

The tree of T. chebula is a highly branched deciduous with 24-30 m height.<sup>8</sup> Leaves are 7 to 20 cm long, not clustered, distant, alternate, elliptic-oblong, acute, rounded. Flowers are short-stalked, hermaphrodites, yellowish white with a strong unpleasant odour and are found in terminal spikes. Drupes ellipsoidal, ovoid, yellow to orange-brown, sometimes tinged with red or black and hard when ripe, 3 to 5 cm long, ribbed on drying. Seeds are hard, pale yellow.<sup>9</sup> It is found throughout the greater parts of India from Ravi eastwards to West Bengal and Assam, ascending to an altitude of 1500 m in the Himalayas. It is also found in Bihar orissa, Madhya Pradesh, Maharashtra, Deccan and south India.<sup>10,11</sup> It grows in deciduous forests of Himachal Pradesh, Tamil Nadu, Kerala, Karnataka, Uttar Pradesh and hra Pradesh and West Bengal.<sup>8</sup>

#### Varieties

According to Basri Terminalia chebula is of 4 types; halelae-zard, halela-e-hindi (smaller and black coloured), halelae-siyah (large size) and hashaf waqaq also known as halaila chini.<sup>12</sup> However, some Unani Physicians described the following three varieties of halela-

Halail-e-siyah (choti har): The fruit falls off from tree before seed formation and turns black within a few days.

Halaila-e-zard: The fruit doesn't fall off from the tree but is semi-ripe, having a seed within it.

Halaila-e-kabuli (kabuli har): The fruit is fully developed, completely ripe and attain full growth.<sup>13</sup>

#### **Plant Profile**

Botanical Name: Terminalia chebula<sup>6,14,15,16,17</sup>

Common Name: Harada<sup>6</sup>

Family: Combretaceae<sup>6,17</sup>

#### **Vernacular Names**

Arabic: Ahleelalj, Ahilaj asfar<sup>5,18</sup>

Persian: Halela<sup>5,18,19</sup>

Hindi: Har, Hara, Harar<sup>5,6,19</sup>

English: Chebulic Myrobalan<sup>5, 16,17</sup>

#### Part Used

Bark<sup>14,20</sup>

Fruit<sup>2</sup>

#### Temperament

Cold in first degree and dry in second degree<sup>14,21</sup>

#### **Traditional Uses**

Unani physicians extensively described this herb as a medicinal plant to treat the different diseases since ancient time. It is widely used drug in Ayurveda, Siddha, Unani and Homeopathic system of medicine in India. it is a top listed herb in Unani Materica medica for treatment of asthma, hemorrhoids, sore throat, gastric disorders (vomiting, anorexia, flatulence), diarrhea, dysentery, spleenomegaly, epilepsy, leprosy, skin disorders, melancholia, gout and joints pain.<sup>6,9,22,23</sup>

Ibn Hubal, renowned Unani physician, in his famous book "Kitab Al-Mukhtarat Fi-Al-tib" mentioned that T. chebula acts as an excellent brain tonic, eye tonic, cardiotonic and blood purifier. Hence, the Unani physicians used this herb for the treatment of dementia, comjunctivitis, cataract, zoaf-e-basarat, palpitation.<sup>24</sup> It is used in Thai traditional medicine as a craminative, astringent and expectorant.<sup>25</sup> The "Triphala", a herbal preparation of "three fruits" from plants T. chebula, T. bellerica, Emblica officinalis, is used as a excellent laxative in chronic constipation, rejuvenator of the body, poor digestion and detoxifying agent of colon also<sup>26</sup>. Recent studies have shown that "Triphala" improve appetite and useful in treating cancer and detoxification<sup>27</sup>.

The fruits of halaila are used by Unani physicians both externally as well as internally for treatment purpose. Externally, the ointment (marham) of halaila (prepared from roghan gul, halaila powder and mom) was used by Unani physicians to cure the piles.<sup>5</sup> The gargle with its

decoction gives excellent results in stomatitis, bleeding and ulceration of gums and sore throat. The powder of Triphala can be used externally for hair wash and prevent hair falling and whitening. A fine powder of halaila is used as a tooth powder to strengthen the gums.<sup>5,21,22,24</sup> The paste of fruit with honey also beneficial in conjunctivitis due to its anti-inflammatory property.<sup>5</sup> The paste of fruits effectively decrease swelling, accelerate the healing and unsoiled the wounds. Halaila also prevents the collection of pus in skin disorders. The oil (roghan) of halaila is extremely helpful in healing of wound especially in burns.<sup>8</sup>

Internally, halaila is applied to cure a vast variety of diseases. The murabbah of halaila is used as an excellent brain tonic, cardiotonic, stomach tonic and in problem of constipation. According to Rhaze's (Rhazi), when halaila powder consumed regularly, it promotes memory, thinking and reasoning power, boosts the nervous system due to beneficial effects on the nerves of brain and also cure the ascites, spleenomegaly, leprosy, colitis and headache. Gastric disorders (anorexia, vomiting, indigestion, flatulence etc.), piles, enlargement of liver and liver, worms, colitis, epilepsy, diarrhea, dysentery can be treated well with halaila. All the verities of halaila are beneficial in chronic fever <sup>5,21,24</sup>. The decoction of haritaki or triphala is given along with honey in hepatitis. Haritaki powder with honey and ghee is also effective remedy for anemia. In obesity, its decoction with honey reduces the excessive body fats.8

#### **Phytochemical Properties**

Although several phytoconstituents like tannins, flavinoids, sterols, amino acids, fructose, resin, fixed oil etc. are present in T. chebula, however it is fairly rich in different tannins (about 32 percent tannin content). The tannin content of T. chebula mainly depends on its geographical distribution.<sup>28</sup> The principal components of tannin are chebulinic acid, chebulic acid, gallic acid, ellagic acid and corilagin.

The tannins of T. chebula (Halaila) are of hydrolysable type and there are approximately fourteen hydrolysable tannins such as chebulic acid, gallic acid, punicalagin, chebulanin, corilagin, neochenulinic acid, ellagic acid, chebulegic acid, chebulinic acid, 1,2,3,4,6 penta-O-galloyl-b-D-glucose, casuarinin, 3,4,6 tri-O-galloyl-D-glucose and terchebulin. All of these tannins are isolate from fruits of T. chebula.<sup>29</sup> Some important phytochemicals like anthraquinones, ethaedioic acid, sennoside, 4,2,4-chebulyl-d-glucopyranose, terpinenes and terpinenols have also been reported to be present. Triterpenoids and their glycosides have been isolated from stem bark of T. chebula<sup>30,31</sup>. Current research proved that T. chebula contains a lot of phenolics than any other plant.<sup>8</sup>

#### Precautions

The halaila should be carefully used by lean peoples, fast, in severe weakness, mental depression and in pregnancy.<sup>8</sup>

#### Pharmacological actions/ Pharmacological Studies

Anti-bacterial activity: Various extracts of T. chebula exhibits ant-bacterial property against different bacterial species.<sup>32</sup> Kannan et al have investigated two imported antibacterial compounds, Gallic acid and ethyl ester against methicillinresistant Staphylococcus, have been isolated from ethyl alcohol extract of the fruit of T. chebula. The T. chebula is well effective against Helicobacter pyroli, bacterium blame to gastritis, ulcer and sometimes stomach cancer.<sup>33</sup> The ether, alcoholic and aqueous extracts of T. chebula were tested against H. pylori, but the aqueous extract of the plant, at a concentration of 1-2.5 mg/ml, inhibited the urease activity of H. pylori.<sup>34</sup> Numerous biologically active components were isolated from butanol fraction of fruit extract of T. chebula and tested against 6 intestinal bacteria. Ethanedioic acid proved strong and moderate inhibitory action against Clostridium perfringens and Escherichia coli, respectively, with no undesirable effects on the growth of the four tested lactic acid-producing bacteria. Ellagic acid exerted an effective inhibitory effect against C. perfringens and E. coli, but little or no inhibition was observed for behenic acid,  $\beta$ -caryophyllene, eugenol, isoquercitrin, oleic acid,  $\alpha$ -phellandrene,  $\beta$ -sitosterol, stearic acid,  $\alpha$ - terpinene, terpinen-4-ol, terpinolene or triacontanoic acid.<sup>35</sup> The ethanolic extract of T. chebula fruit was found effective against both gram-positive and gram-negative bacteria such as Salmonella typhi SSFP 4S, Staphylococcus epidermidis MTCC 3615, Staphylococcus aureus ATCC 25923, Bacillus subtilis MTCC 441 and Pseudomonas aeruginosa ATCC 27853 suggesting its broad-spectrum antimicrobial activity.<sup>33</sup> Anti-fungal activity: Aqueous extract of T. chebula has been reported to exhibit antifungal activity against different varieties of dermatophytes such as Epidermophyton, Floccosum, Microsporum gypseum, Tricophyton rubrum and yeasts e.g. Candida albicans.<sup>36,37</sup> Aqueous, alcoholic and ethyl acetate extracts of leaves of T. chebula were also tested against five pathogenic fungi (Aspergillus flavus, A. niger, Alternaria brassicicola, A. alternate and Helminthosporium tetramera) using paper disc method and were found effective compared to that of the reference standard Carbendazim.38

Antiamoebic and immunomodulatory activities: In a study, T. chebula was evaluated in an experimental amoebic liver abscess in golden hamsters and in immunomodulation studies. The formulation had a maximum cure rate of 73% at 800 mg/ kg body weight in hepatic amoebiasis. In immunomodulation studies, humoral immunity was improved where T-cell counts remained unchanged in the animals, but the cell-mediated immune response was stimulated.<sup>39</sup>

#### Antiplasmodial activity

T. chebula, water extract revealed antiplasmodial activity

in vitro because it inhibits the uptake of [3H] hypoxanthine into the Plasmodium falciparum K1 multidrug-resistant strain and in vivo.<sup>40</sup>

#### **Molluscicidal Activity**

The ethanolic extract of T. chebula fruit powder exhibits molluscicidal activity against vector snail Lymnaea acuminate. Liquid chromatography analyses demonstrated that the active molluscicidal component in T. chebula was tannic acid. Therefore, T. chebula could be an effective source of molluscicides against the snail L. acuminates.<sup>41</sup>

#### **Anti-helminthes Activity**

The ovicidal and larvicidal activities of ethyl acetate, acetone and methanol extracts of dried leaves and seeds of T. chebula were examined in vitro on Haemonchus contortus based on egg hatch and larval development assays at 50, 25, 12.5, 6.25 and 3.13 mg/ml. The extracts of leaves and seeds of T. chebula showed complete inhibition at 50 mg/ml.<sup>42</sup>

Anti-viral activity: The extract of the fruit of T. chebula showed inhibitory effects on HIV-1 reverse transcriptase.<sup>43</sup> In a study, Hot water extract of T. chebula showed antiherpes simplex virus (HSV) activity in vivo and anticytomegalovirus (CMV) activity both in vitro and in vivo.<sup>44</sup> A study proved that T. chebula fruits contain 4 human HIV-type-1 integrase inhibitors, for example gallic acid and three galloyl glucoses and suggested that galloyl moiety had a key role for inhibition of the 3'-processing of HIV-1 integrase by these compounds.<sup>45</sup> T. chebula can also be used in sexually transmitted diseases (STDs) and AIDS.<sup>46</sup> Recently, acetone extract of T. chebula has emerged as a new substitute to treat pandemic swine influenza- An infection due to its low price, easy preparation and potential effect.<sup>47</sup>

#### Antimutagenic and Anti-carcinogenic Activities

Ponnusankar et al have performed by the effect of 70% methanolic fruit extract of T. chebula was examined on the growth of numerous malignant cell lines. One of the fractionated compounds from ethanolic fruit extract of T. chebula, chebulagic acid, showed potent dual inhibition against COX and 5-LOX. It also showed anti-proliferative activity against HCT-15, COLO-205, MDA-MB-231, DU-145 and K562 cell lines. A recent study has shown the ability of triphala to inhibit cytochrome P450.<sup>48</sup>

#### **Anti-oxidant Activity**

Chen et al have done the study on the T. chebula and found that it is an excellent anti-oxidant. In a study, 6 extracts and 4 pure compounds of T. chebula showed anti-lipid peroxidation, anti-superoxide radical formation and free radical scavenging properties at different concentration. The outcome established that tri-ethyl-chebulate was a powerful antioxidant as well as free-radical scavenger, which might donate to the anti-oxidative ability of T. chebula.<sup>49</sup>

#### Anti-diabetic Activity

The blood sugar level in normal and alloxan diabetic rats reduces significantly within 4 h by Oral administration of 75% methanolic extract of T. chebula (100 mg/kg body weight) and constant every-day administration of the medicine produced a sustained result.<sup>50</sup> The chloroform extract of T. chebula seeds (100, 200 and 300 mg/kg body weight) produced a dose-dependent decline in blood sugar level of diabetic rats equally in short term and long term study (300 mg/kg body weight for 8 weeks). Additional, notable renoprotective action was also detected in T. chebula treated rats.<sup>51</sup>

#### Anti-anaphylactic Activity

Shin et al have examined on the T. chebula along with numerous new medicinal plants help to resist against many stressors in different ways. T. chebula diminish the serum histamine levels and showing a strong anti-anaphylactic property when it administered following anaphylactic shock.<sup>52</sup>

https://pdfs.semanticscholar. org/582d/1ce689d0092c76d2e08c0dffa479ed32fab4.pdf

#### **Anti-nociceptive Activity**

The petroleum ether, chloroform, ethanol and water extracts of T. chebula fruits were evaluated for their analgesic activity using the tail immersion model in mice. The ethanolic extract of the plant exhibited analgesic response at 200- 400 and 800mg/kg body weight in acute pain and in chronic pain studied for 15 days with a maximum analgesic response on 14th day.<sup>53</sup>

#### Anti-ulcerogenic Activity

Sharma et al. have examined on the animals pre-treated at 200 and 500 mg/kg body weight with hydro alcoholic extract of T. chebula showed a reduction in lesion index, total affected area and percentage of lesion in comparison with control groups in the aspirin, ethanol and cold restraint stress-induced ulcer models.<sup>54</sup>

#### **Anti-arthritic Activity**

The hydroalcoholic extract of T. chebula produced a significant inhibition of joint swelling as compared to control in both formaldehyde-induced and CFA-induced arthritis. T. chebula treatment also reduced serum TNF- $\alpha$  level and synovial expression of TNF-R1, IL-6 and IL-1 $\beta$ . The authors believed that T. chebula could be used as a disease-modifying agent in the treatment of rheumatoid arthritis.<sup>55</sup>

#### Wound Healing Activity

Choudhary et al have done on the alcoholic extract of the leaves of T. chebula caused much faster healing of rat dermal wounds in-vivo due to improved rates of contraction and a decreased period of epithelialization for the topical administration. Biochemical studies revealed an increase in total protein, DNA and collagen contents in the granulation tissues of treated wounds.  $^{\rm 56}$ 

# **Cyto-protective Activity**

Manosroi et al. have performed on the different concentrations of gallic acid and chebulagic acid, isolated from fruit extract of T. chebula, blocked cyto-toxic T lymphocyte (CTL)-mediated cyto-toxicity. Granule exocytosis in response to anti-CD3 stimulation was also blocked by the above phyto-chemicals at the equivalent concentrations.<sup>57</sup>

# **Radio-protective Activity**

Gandhi et al. have estimates on the aqueous extract of the fruit of T. chebula (50  $\mu$ g) was able to neutralize 1, 1-diphenyl-2picrylhydrazyl, a stable free radical by 92.9% and protected the plasmid DNA pBR322 from undergoing the radiation-induced strand breaks.<sup>58</sup>

# **Cardio-protective Activity**

Cardioprotective effect of ethanolic extract of T. chebula fruits (500 mg/kg body weight) was investigated in isoproterenol-induced myocardial damage in rats. It was reported that the pretreatment with T. chebula extract had a cardioprotective effect due to the lysosomal membrane stabilization preventing myocardial necrosis and inhibition of alterations in the heart mitochondrial ultrastructure and function in the experimental rats.<sup>59</sup>

#### **Hepato-protective Activity**

The 95% ethanolic extract of T. chebula fruit showed hepatoprotective activity against anti-tuberculosis (anti-TB) drug-induced toxicity which could be attributed to its prominent anti-oxidative and membrane stabilizing activities.<sup>60</sup>

#### Anti-spermatogenic Activity

Gupta et al. have performed on the oral administration (300 mg/kg body weight for 28 days) of the bark of T. chebula extracted in acetone, methanol, 50% ethanol and in aqueous solvents caused histological alterations in seminiferous tubules in testes of treated mice.<sup>61</sup>

#### **Chemoproventive Activity**

In an investigation, T. chebula extract treatment prevened nickel chloride induced renal oxidative stress, toxicity and cell proliferation response in male Wistar rats. The authors suggested that T. chebula extract could be used as a therapeutic agent for cancer prevention as it blocked or suppressed the events associated with chemical carcinogenesis.<sup>62</sup>

#### Hypolipidemic and Hypocholesterolemic Activity

T. chebula extract administration showed hypolipidaemic activity against experimentally induced atherosclerosis and

hypocholesterolemic activity against cholesterol-induced hypercholesterolemia and atherosclerosis.<sup>63</sup> Triphala formulation was found to have hypolipidaemic effects on the experimentally induced hypercholesteremic rats.<sup>64</sup>

# Therapeutic Uses

- As Halela cause purgation of bile (safra), it is used in bilious and melancholic disorders.<sup>5,19</sup>
- It is beneficial in piles, epilepsy, diarrhea, dysentery, bleeding and ulceration of gums, leprosy and melancholia <sup>6,22,23</sup>.
- It strengthens the brain, eye, gums.<sup>5,6,7</sup>
- It is useful in Zof-e-meda and various diseases of eyes like Zof-e-Basarat (dim-vision), Dhalk and Conjunctivitis.<sup>5,6,18,19</sup>
- It is useful in gastric disorders such as anorexia, vomiting, indigestion, flatulence and constipation.<sup>3</sup>
- It is beneficial in Gout and Joints Pain (Wajaul Mafasil).<sup>3,20</sup>

# Dose

7gm-10.5gm<sup>21</sup>, 7gm-17gm<sup>20</sup>, 5gm-7gm<sup>18</sup>

# Substitute (Badal)

Post-e-Anar (Epicarp of pomegranate) and Mazu <sup>20</sup>.

Halela Kabuli<sup>6</sup>.

# Toxicology

# Harmful (Muzir)

It is harmful to intestine, anus, liver.<sup>6</sup>

#### **Corrective (Musleh)**

Unnab (Zizyphus vulgaris).6

Sapistan (Cordia latifolia Roxb).6

Almond oil (Prunus amygdalus Batsch).<sup>261</sup>

# Popular Unani Preparations (Murakkabat).<sup>5,6</sup>

Habb-e-muqil, Itrifal Saghir, Itrifal Kabir, Itrifal Kishneezi, Itrifal Zamani, Safoof-e-halela.

# Conclusion

In spite of the overwhelming influences and our addiction on modern medicines and synthetic drugs, a large section of the world population still likes drugs of plants origin for treatment of the 2,50,000 higher plant species on earth, more than 80,000 are medicinal. However, only 7000-7500 species are used for their medicinal values by traditional communities. T. chebula (haritaki) is an essential herb used by renowned Unani Physicians since ancient time to cure various disorders because of having a number of medicinal property. The main action of T. chebula are stomachic, blood purifier, diuretic, antidysentric, purgative (safra), sedative, astringent as well as it is a good eye tonic, cardio-tonic and brain tonic. Hence, it is widely used in Unani Medicine to treat gout, joints pain, conjunctivitis, low vision, piles, epilepsy, melancholia and leprosy, to strengthen the brain, eye, gums and stimulates digestive capacity. It is a good source of a variety of biologically active phytoconstituents such as chebulic acid, chebulinic acid, chebulagic acid, gallic acid, corilagin ellagic acid and other related compounds which are responsible for antimicrobial, antioxidant, antihyperglycemic, anticancer and protective effects on different human vital organs such as nerves, heart, kidney and liver. Traditionally, Terminalia chebula is used to treat a vast variety of health problems. Consequently, there is an urgent need to investigate the biological activity of its phytoconstituents for development of a new more effective, cheap, reliable herbal drug with better efficacy and higher safety margin.

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# References

- 1. Singh P, Malhotra H. Terminalia Chebula-A review pharmacognistic and phytochemical studies. *International Journal of Recent Scientific Research* 2017; 8(11): 21496-21507.
- 2. Dinesh MD, soorya TM, Vismaya MR et al. Terminalia chebula A Traditional Herbal Drug-A Short Review. *International Journal of Pharmaceutical Science Invention* 2017; 6(2): 39-40.
- Sharma PC, Velnu MB, Dennis TJ. Database on Medicinal Plants used in Ayurvedic- 7-Volume Set. CCRAS 2002; 1,3,5: 225-243.
- 4. Najmul GH. Advia K. Part I-IV. New Delhi: Idara Kitab us shifa Daryaganj, YNM: 308-312, 861-862,1352-1354.
- 5. Hakeem K, Mufradat M. Delhi: Ejaz Publishing House Daryaganj, YNM: 102-103, 363-364, 590-591.
- 6. Qasmi IA. Mufridat KU. I<sup>st</sup> ed. Aligarh: International Printing Press 2001; 129-130, 154-155, 236-238.
- 7. Nadkarni AK. Indian Materia Medica. Bombay: Popular Prakashan 1982; 1: 278-279.
- 8. Gupta PC et al. Biological and Pharmacological Properties of Terminalia Chebula Retz. (Haritaki)-An Overview. *Int J Pharm Sci* 2012; 4 (3): 62-68.
- 9. Kirtikar KR, Basu BD. Indian Medicinal plants. Periodical Expert Book Agency 1984; 2: 1429-1431.
- 10. Sharma PC, Velnu MB, Dennis TJ. Database on Medicinal Plants used in Ayurvedic. CCRAS 2002; 1,3,5: 225-243.
- 11. Standardisation of Single Drugs of Unani Medicine. Part I, IV. CCRUM Dept. of Ayush 2006: 86-90, 202-206.

- 12. Ahmed ZAB. Mufradat AJU, Aghzia AAW. Urdu translation by CCRUM, New Delhi; 2,3,5: 51-53, 96-98, 170-175, 349-352.
- 13. kabeeruddin, Akseer HMA. Part II Gulberg: Al-Shifa: 1448-1450.
- Chopra RN et al. Glossary of Indian Medicinal Plants. 1<sup>st</sup> ed. 4<sup>th</sup> Reprint. New Delhi: National Institute of Science and Communication 1996: 67, 73, 75, 242.
- Kirtikar KR, Basu BD. Indian Medicinal Plants. 2<sup>nd</sup> ed. Dehradun: International Book Distributor, 1981; 2: 1020-1023.
- Chatterjee A, Prakashi SC. The Treatise of Indian Medicinal Plants. New Delhi: National Institute of Science Communication CSIR 1997; 3: 69-71,203-204.
- 17. Rastogi RP, Mehrotra BN. Compendium of Indian Medicinal Plants. New Delhi: National Institute of Science Communication and Information Resources, Reprint 2004; 1: 119,105,407,653-654.
- Hussain AHM, Mufradat K. 3<sup>rd</sup> ed. Lahore: Sheikh Ghulam Ali and Sons, Kashmeeri Bazaar 1960: 433, 304-305,503-504,526-527.
- 19. Ahmad TN. Advia KU. New Delhi: Idara Kitab-us-Shifa Kocha Chelan Daryaganj, YNM: 105-107,460,759-761.
- 20. Najmul GH, Advia K. Part I-IV. New Delhi: Idara Kitab us shifa Daryaganj, YNM: 308-312, 861-862,1352-1354.
- Abdullah Ibn BZ. Aljamiul Mufradat Al Advia Wa Al Aghzia (Urdu Translation). New Delhi: CCRUM Ministry of Health and Family Welfare, 2003; 4: 354,-356, 436-439.
- 22. AK Nadkarni. Indian Materia Medica. Bombay: Popular Prakashan 1982; 1: 278-279.
- 23. Harichand MH. Taj-ul-Aqaqueer-Hindustan ki Jadi Butiyan. Nirala Jogi Publications, YNM; 1: 378-383,528-532,814-816.
- Hubal Ibn B. Kitab Al Mukhtarat Fit Tib (Urdu Translation by CCRUM). New Delhi: Ministry of Health and Family Welfare 2007; 2: 118,209,239-240.
- 25. Panunto W, Jaijoy K, Lerdvuthisopon N et al. Acute and chronic toxicity studies of water extract from dried fruits of Terminalia chebula Retz. In rats. *Int J Applied Research in Natural Products* 2011; 3(4): 36-43.
- 26. Mohd A, Khan U, Iqbal A et al. Comprehensive review on therapeutic strategies of gouty arthritis. *Pak J Pharm Sci* 2014; 27(5): 1575-1582.
- 27. Marc HC, Alan SJ, Josef SS et al. Rheumatology 3<sup>rd</sup> ed. Spain Mosby: An imprint of Elsevier Ltd 2003; 2: 1893-1936.
- 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.
- 29. Juang LJ, Sheu SJ, Lin TC. Determination of hydrolysable tannins in the fruit of Terminalia chebula Retz. by

high performance liquid chromatography and capillary electrophoresis. *J Sep Sci* 2004; 27(9): 718-24.

- Rastogi RP, Mehrotra BN. Compendium of Indian Medicinal Plants. New Delhi: National Institute of Science Communication and Information Resources, Reprint 2004; 27(9): 718-724.
- 31. Rastogi RP, Mehrotra BN. Compendium of Indian Medicinal Plants. New Delhi: National Institute of Science Communication and Information Resources, Reprint 2008; 4: 215-719.
- 32. Ahmad I, Mehmood Z, Mohammad F. Screening of some Indian medicinal plants for their antimicrobial properties. *J Ethnopharmacol* 1998; 62: 183-93.
- 33. Kannan P, Ramadevi SR, Hopper W. Antibacterial activity of Terminalia chebulafruit extract. *African Journal of Microbiol Res* 2009; 3(4): 180-84.
- 34. Malekzadeh F, Ehsanifar H, Shahamat M et al. Antibacterial activity of black myrobalan (Terminalia chebula Retz) against Helicobacter pylori. *J Antimicrobial Agents* 2001; 18: 85-88.
- 35. Kim HG, Cho JH, Jeong EY et al. Growth-inhibiting activity of active component isolated from Terminalia chebula fruits against intestinal bacteria. *J Food Prot* 2006; 69(9): 2205-2209.
- 36. Dutta BK, Rahman I, Das TK. Antifungal activity of Indian plant extracts. Mycoses 1998; 41(11-12): 535-36.
- 37. Mehmood Z, Ahmad I, Mohammad F et al. Indian medicinal plants: A potential source for anticandidal drugs. *Pharmaceutical Biology* 1999; 37(3): 237-42.
- 38. Shinde SL, More SM, Junne SB et al. The antifungal activity of five Terminalia species checked by paper disk method. *Int J Pharma Res and Develope* 2011; 3(2).
- 39. Sohni YR, Bhatt RM. Activity of a crude extract formulation in experimental hepatic amoebiasis and in immunomodulation studies. *J Ethnopharmacol* 1996; 54(2-3): 119-24.
- Pinmai K, Hiriote W, Soonthornchareonnon N et al. In vitro and in vivo antiplasmodial activity and cytotoxicity of water extracts of Phyllanthus emblica, Terminalia chebula and Terminalia bellerica. J Med Assoc Thai 2010; 93(7): 120-26.
- 41. Upadhyay A, Singh DK. Molluscicidal activity of Sapindus mukorossi and Terminalia chebula against the freshwater snail Lymnaea acuminate. *Chemosphere* 2011; 83(4): 468-74.
- 42. Kamaraj C, Rahuman AA. Efficacy of anthelmintic properties of medicinal plant extracts against Haemonchus contortus. *Res Vet Sci* 2011; 91(3): 400-404.
- 43. Mekkawy SE, Meselhy MR, Kusumoto IT et al. Inhibitory Effects of Egyptian Folk Medicines oh Human Immunodeficiency Virus (HIV) Reverse Transcriptase. *Chemical & Pharmaceutical Bulletin* 1995; 43(4): 641-

648.

- 44. Yukawa TA, Kurokawa M, Sato H et al. Prophylactic treatment of cytomegalovirus infection with traditional herbs. *Antiviral Res* 1996; 32(2): 63-70.
- 45. Ahn MJ, Kim CY, Lee JS et al. 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.
- 46. Vermani K, Garg S. Herbal medicines for sexually transmitted diseases and AIDS. *J Ethnopharmacol* 2002; 80(1): 49-66.
- 47. Ma H, Diao Y, Zhao D et al. A new alternative to treat swine influenza A virus infection: extracts from Terminalia chebula Retz. *African J Microbiol* 2010; 4(6): 497-99.
- 48. Ponnusankar S, Pandit S, Babu R et al. Cytochrome P450 inhibitory potential of Triphala-A Rasayana from Ayurveda. *Journal of Ethnopharmacol* 2011; 133(1): 5.
- 49. Chen X, Sun F, Ma L et al. In vitro evaluation on the antioxidant capacity of tri-ethylchebulate, an aglycone from Terminalia chebula Retz fruit. *Indian Journal of Pharmacol* 2011; 43(3): 6.
- 50. Sabu MC, Kuttan R. Anti-diabetic activity of medicinal plants and its relationship with their antioxidant property. *J Ethnopharmacol* 2002; 81: 155-60.
- 51. Rao NK, Nammi S. 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.
- Shin TY, Jeong HJ, Kim DK et al. Inhibitory action of water soluble fraction of Terminalia chebulaon systemic and local anaphylaxis. *Journal of Ethnopharmacol* 2001; 74: 6.
- Kaur S, Jaggi RK. Antinociceptive activity of chronic administration of different extracts of Terminalia bellerica Roxb and Terminalia chebula Retz Fruits. *Indian J Exp Biol.* 2010; 48(9): 925-30.
- 54. Sharma P, Prakash T, Kotresha D et al. Anti-ulcerogenic activity of Terminalia chebula fruit in experimentally induced ulcer in rats. *Pharm Biol* 2011; 49(3): 7.
- Nair V, Singh S, Gupta YK. Anti-arthritic and disease modifying activity of Terminalia chebula Retz. in experimental models. *J Pharm Pharmacol* 2010; 62(12): 1801-1806.
- 56. Choudhary GP. Wound healing activity of ethanolic extract of Terminalia chebula Retz. *International Journal of Pharma and Bio Sci* 2011; 2(1): 7.
- 57. Manosroi A, Jantrawut P, Akihisa T et al. In vitro antiaging activities of Terminalia chebula gall extract. *Pharm Biol* 2010; 48(4): 469-481.
- 58. Gandhi NM, Nair CKK. Radiation protection by Terminalia chebula: Some mechanistic aspects. *Mol*

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Cell Biochem 2005; 277(2): 43-48.

- 59. Suchalatha S, Shyamala Devi CS. Protective effect of Terminalia chebula against experimental myocardial injury induced by isoproterenol. *Indian J Exp Biol* 2004; 42(2): 174-78.
- 60. Tasduq SA, Singh K, Satti NK et al. Terminalia chebula (fruit) prevents liver toxicity caused by sub-chronic administration of rifampicin, isoniazid and pyrazinamide in combination. *Hum Exp Toxicol* 2006; 25(3): 111-118.
- 61. Gupta PC, Singh SK. Effect of different extracts of bark of Terminalia chebula on reproductive organs in male mice. ICRH & 20<sup>th</sup> Annual Meeting of the Indian Society for the Study of Reproduction & Fertility 2010: 8.
- 62. Prasad L, Khan TH, Jahangir T et al. Chemomodulatory effects of Terminalia chebula against nickel chloride induced oxidative stress and tumor promotion response in male Wistar rats. *J Trace Elements in Med and Biol* 2006; 20(4): 233-239.
- 63. Shaila HP, Udupa SL, Udupa AL. Hypolipidemic activity of three indigenous drugs in experimentally induced atherosclerosis. *Int J Cardiol* 1998; 67(2): 119-124.
- 64. Saravanan S, Shrikumar R, Manikandan S et al. Hypolipidemic effect of Triphala in experimentally induced hypercholesterolemic rats. *Yakugaku Zasshi* 2007; 127(2): 385-388.