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## Pharmacological activities of Baheda (*Terminalia bellerica*): A review

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### Abstract

*Terminalia bellerica* Roxb (combretaceae) is found widely throughout the Indian subcontinent, Sri Lanka, South- East Asia, Bangladesh as a medicinal plant. Plant and plant parts are used in the traditional system of medicines like Ayurveda, Siddha, Unani & Chinese medicine. The plant is constituted of Glucoside, Tannins, Gallic acid, Ethyl Gallate, Chebulinic acid which serves as an antioxidant, antimicrobial, antidiarrheal, anticancer, antihypertensive, hepatoprotective & antipyretic agent. This review sites the information on pharmacological activities of *Terminalia bellerica* which may serve as a source for further research studies.

**Keywords:** *Terminalia bellerica*, Phytoconstituents, Traditional use.

### 1. Introduction

Plants have been used as medicines from the ancient times. Throughout the world medicinal plants are widely and successfully used. A plant with active medicinal constituents are used to treat diseases in the traditional systems like Ayurveda, Siddha and Unani. In Asia, the use of medicinal plants are well established and are well documented. The plants those are recognized internationally mostly comes from this region. Plants, plant parts and plant products those are used for the preparation of medicines serves wee to uplift the economical status of the country and they are the natural wealth of a country. Medicinal plants has got significant role in saving the lives of rural area people. In India, 45,000 plant species have been identified and out of which 15-20 thousands plants are found to have good medicinal value. Study says about 6000 traditional plants are used in Indian traditional and herbal medicines. In this paper, we selected the plant *Terminalia bellerica* belonging to family Combretaceae to study its pharmacological effect.

### 2. Synonyms

Assam - Bhomora, Bhomra, Bhaira;  
Eng – Beleric Myrobalan;  
Guj - Bahedam, Baheda;  
Hindi - Bahera;  
Kan - Shanti, Shantikayi, Tare, Tarekayi;  
Mal - Tanni, Tannikai;  
Mar - Baheda;  
Ori - Baheda, Bhara;  
Sansk - Vibhita, Aksa, Aksaka, Bibhitaki;  
Tam - Thanakkai, Tanri, tanrikkai, Tani;  
Tel - Tannikkaya, Vibhitakami, Tani

### 3. Plant description

*Terminalia bellerica* is a large deciduous tree to 50 m tall and a diameter of 3 m with a rounded crown. The frequently buttressed bole at the base is branchless up to 20 m. The bark is bluish or ashy-grey covered with numerous fine longitudinal cracks, the inner bark yellowish. Leaves large, glabrous, alternate, broadly elliptic to obovate-elliptical, 4-24 cm x 2-11 cm, base rounded to cuneate, rufous-sericeous but soon glabrescent, with 6-9 pairs of secondary veins. Secondary and tertiary venation prominent on both surfaces, clustered

towards the ends of branchlets. Petiole 2.5-9 cm long. Young leaves copper-red, soon becoming parrot green, then dark green. Flowers solitary, small, 3-15 cm long, greenish white, simple, axillary spikes; calyx tube densely sericeous or tomentulose; flowers appear along with new leaves and have a strong honey-like smell. Fruit sub-globular to broadly

ellipsoid, 2-4 x 1.8-2.2 cm, densely velutinous or sericeous, light-yellow, obscurely 5-angled and minutely brown tomentosa. The generic name 'Terminalia' comes from Latin word 'terminus' or 'terminalis' (ending), and refers to the habit of the leaves being crowded or borne on the tips of the shoots.



Fig 1: Fruits



Fig 2: Plant branches



Fig 3: Plant as a whole

#### 4. Traditional uses

Fruits are laxative, astringent, anthelmintic and antipyretic; useful in hepatitis, bronchitis, asthma, dyspepsia, piles, diarrhoea, coughs, hoarseness of voice, eye diseases and scorpion-sting; used as a hair tonic. Decoction of the green fruit is used for cough. Pulp of the fruit is useful in dysenteric-diarrhoea, dropsy, piles and leprosy. Half ripe fruit is used as purgative. Kernel of the fruit is narcotic. Fruits are used in menstrual disorder in Khagrachari. Seed oil is used in rheumatism. Gum of the bark is demulcent and purgative. The triterpenoid present in the fruits possess significant antimicrobial activity. Kernel oil has purgative action and its prolonged use was well tolerated in mice (Ghani, 2003) [1].

#### 5. Phytoconstituents

Glucoside (bellericanin) [12], Gallo-tannic acid, Coloring matter, resins and a greenish yellow oil [9]. Ellagic acid, gallic acid, lignans (termilignan and thannilignan), 7-hydroxy 3'4' (methylenedioxy) flavone and anolignanB [11]. Tannins, ellagic acid, ethyl gallate, galloyl glucose and chebulagic acid, phyllembin,  $\beta$ -sitosterol, mannitol, glucose, fructose and rhamnase [12, 13].

#### 6. Pharmacological effects

**6.1 Analgesic activity:** ArifUllah Khan *et al.*, (2010) describes the antisecretory and analgesic activities of the crude extract of *Terminalia bellerica*. *T. bellerica* extract at the dose range of 300 - 1000 mg/kg inhibited the castor oil-induced intestinal fluid secretion in mice. The extract also dose-dependently (50 - 100 mg/kg) where it reduced the numbers of acetic acid-mediated in mice. These results indicate that *TB* exhibit antisecretory and antinociceptive effects, hence justifying its medicinal use in diarrhea and pain [4].

**6.2 Anti diarrhoeal activity:** The Anti diarrhoeal activity was performed using Castor oil induced diarrhoea, PGE2 induced entero pooling and gastrointestinal motility test (Bimlesh Kumar *et al.*, 2010). Aqueous and ethanolic extract of fruit pulp of *TB* at the doses of 334 mg/kg, 200 mg/kg, 143 mg/kg were used. Comparison of percentage protection in these models revealed that the extracts have more prominent anti-secretory effect than the reduction in gastrointestinal motility [14].

**6.3 Antihypertensive Effect:** Arif Ullah Khan *et al.*, (2008) was screened the effect of *TB* in hypertension. After administration of *TB*, they observed that fall in the arterial BP of rats under anaesthesia. In isolated guinea-pig atria, inhibition of force and rate of atrial contractions noted. In rabbit thoracic aorta, relaxation was observed after the induction of contractions which was induced by phenylephrine [4].

**6.4 Anti salmonella activity:** Madani A *et al.*, (2008) were studied the effect of *T. bellerica* against *Salmonella typhi* and *Salmonella typhimurium*. *In vitro* cellular toxicity also performed by them. In this study, Petroleum ether, chloroform, acetone, alcohol and aqueous extract of *TB* fruit taken for screening. When compared with other extracts bothalcoholic and aqueous extracts of *TB* showed significant anti-salmonella activity. There was no cytotoxicity was observed in *in vitro* cellular toxicity study [5].

**6.5 Anti-Spasmodic and Bronchodilatory Properties:** Anwarul Hassan Gilani *et al.*, (2008) were postulated that the crude extract of *TB* fruits elicited relaxation of spontaneous contractions in both isolated rabbit jejunum and guinea-pig ileum. Protective effect of *TB* against castor oil-induced diarrhea and carbachol-mediated bronchoconstriction also observed in rodents. In guinea-pig trachea, *TB* relaxed the CCh-induced contractions [6].

**6.6 Anti-microbial activity:** Elizabeth K M *et al.*, (2005) were conducted the antimicrobial activity of *TB* against 9 human microbial pathogens. The Aqueous extract of dry fruit at 4 mg concentration showed highest zone of inhibition against *S.aureus*. These pathogens were highly sensitive to the methanol extract also except *E. coli* (enteropathogen) and *P. aeruginosa*. Finally they concluded that *TB* dry fruit possesses potential broad spectrum antimicrobial activity [8].

**6.7 Antimicrobial and Toxicity Studies:** Badrul Alam *et al.*, (2011) postulated that the crude methanolic extract of the fruits of *Terminalia bellerica* Roxb along with its various organic fractions elicited both *in vitro* and *in vivo* antioxidant activity as well as antibacterial activity. Total antioxidant activity, scavenging free radical, authentic peroxy nitrite and reducing power assessment were performed. Finally they concluded that the EtOAc fraction elicited strong activity in all the model systems with moderate toxicity [3].

**6.8 Antioxidant activity:** Ramesh Kumar *et al.*, (2011) postulated that the crude aqueous extract of the fruits of *Terminalia belerica* Roxb have antioxidant properties since these contains enzymatic and non – enzymatic antioxidants, these can be very effective against microbes causing various diseases. *In vitro* assessment of the antioxidant activity of ethanolic fractions of both these plants to scavenge 2, 2-Diphenyl-1-picrylhydrazyl (DPPH) and highly reactive hydroxyl radicals showed that the semi pure compounds present in the fractions are useful potential source of antioxidants and can be used in the therapy of diseases like cancer, coronary heart disease, ageing and any other disease related to oxidative stress. These fractions being non-toxic showed significant antioxidant activity at scavenging free radicals. They also significantly scavenge hydroxyl radical which is known to cause cellular damage<sup>[9]</sup>.

**6.9 Wound healing activity:** Saha *et al.* (2011) postulated that the paste of *Terminalia belerica* Roxb have proper efficacy on wound healing. Herbal paste preparation showed significant ( $P < 0.05$ ) improvement on maturation, wound contraction and epithelialization. Therefore it may be concluded that the paste obtained from *Terminalia belerica* offers a distinctive advantage in wound healing<sup>[10]</sup>.

**6.10 Immunological activity:** Aurasorn Saraphan choti withthaya *et al.* postulated that *T. bellerica* extract affected T cell proliferation mainly through the same mechanism as PHA. The extract with LPS and PWM also affected B cell proliferation through T cell-independent and T cell-dependent mechanisms respectively. The results indicated that the extract affected cellular mediated immunity (CMI) rather than humoral mediated immunity (HMI)<sup>[11]</sup>.

**6.11 Acute and Sub-acute Toxicities:** Thanabhorn S. *et al.*, (2009) were conducted acute and sub-acute toxicity studies as per the OECD guideline. Single oral administration of the ethanolic extract of *T. belerica* at a dose of 5,000 mg/kg did not produce any toxicity. In sub-acute toxicity, repeated administration of 1,000 mg/kg of *T. belerica* over 14 days did not cause changes in terms of general behaviors, mortality, weight gain, hematological or clinical blood chemistry parameters. The results of histological examinations showed normal appearance of the internal organs when compared to those of the control group<sup>[16]</sup>.

**6.12 Immune response *In vitro*:** *In vitro* Phagocytic activity and lymphocyte proliferation assay were carried out in methanolic extract of on the mouse immune system (Aurasorn Saraphanchotiwiththaya *et al.*, 2008). In both assay, stimulation of macrophage phagocytosis and maximal activation of phytohemagglutinin were observed. Finally, the authors concluded that the methanolic extract of *T. belerica* affected the mouse immune system, specifically both the cellular and humoral immune response *in vitro*<sup>[11]</sup>.

**6.13 Hepatoprotective activity:** Sangeetha Shukla *et al.*, (2006) were evaluated the protective effect of TB fruit extract and its active principle, Gallic acid against CCl<sub>4</sub> intoxication. Treatment with extract (200, 400 and 800 mg/kg, p.o.) and gallic acid (50, 100 and 200 mg/kg, p.o.) showed dose-dependent recovery in biochemical parameters such as SGOT, SGPT and lipid peroxidase, glutathione but the effect was more pronounced with gallic acid<sup>[7]</sup>.

**6.14 Antibiofilm Activity:** The ethanolic extract of a plant *Terminalia bellerica* (common name = Baheda) was tested for its antimicrobial activity against the oral plaque forming bacteria *Streptococcus mutans*. It was found to significantly inhibit biofilm formation. It was found that the extract from *Terminalia bellerica* showed strong activity against *Streptococcus mutans*. The extract also prevents the formation of biofilm by the bacteria. The study suggests possible benefits of this herbal preparation which inhibit the biofilm formation by streptococci, a oral pathogens<sup>[15]</sup>.

**6.15 Anticancer Activity:** *P. emblica* and *T. bellerica* extracts demonstrated growth inhibitory activity, with a certain degree of selectivity against the two cancer cell lines tested. Synergistic effects (CI < 1) for *P. emblica*/doxorubicin or cisplatin at different dose levels were demonstrated in A549 and HepG2 cells. The *T. bellerica*/cisplatin or doxorubicin also showed synergistic effects in A549 and HepG2 cells. In some instances, the combinations resulted in antagonistic effects. The dose reduction level was different and specific to each combination and cell line<sup>[17]</sup>.

**6.16  $\beta$ -lactamase inhibitor activity:** The  $\beta$ -lactamase inhibitor activity of 68 extracts from Indian herbs and spices was surveyed. Most promising results of the  $\beta$ -lactamase inhibitor activity *in vivo* and *in vitro* were achieved from the herbal extracts of Baheda (*Terminalia bellerica*), Ginger (*Zingiber officinale*), Brahmi (*Bacopa monnieri*), Garlic (*Allium sativum*), Gurmar (*Gymnema sylvestre*), Satavar (*Asparagus racemosus*) and Pomegranate (*Punica granatum*) peels and seeds against *Staphylococcus aureus* as the test organism<sup>[18]</sup>.

**6.17 Antiulcer Activity:** The anti-ulcer activity of ethanolic extract of *Terminalia belerica* (Combretaceae) fruits ETB was investigated in pylorus ligation and ethanol induced ulcer models in wistar rats. In both models the common parameter determined was ulcer index. ETB at doses of 250,500 mg/kg orally produced significant inhibition of the gastric lesions induced by Pylorus ligation induced ulcer & Ethanol induced gastric ulcer. The extract (250 mg/kg & 500 mg/kg) showed significant ( $P < 0.05$ ) reduction in free acidity and ulcer index as compared to control<sup>[19]</sup>.

**6.18 Antithrombotic and Thrombolytic activity:** An *in vitro* model was used to check the clot lysis and antithrombotic effect of *Terminalia belerica* fruits along with Streptokinase as a positive control. From this study it was found that after addition of Streptokinase clot formation is delayed upto more than 90 min whereas after addition of test solution it was found that as the concentration of extract was increased the delay in clot formation also increases. At 0.20 mg/dl concentration it showed the maximum delay (more than 90 min.) in clot formation. For thrombolytic activity, at concentration 1.00 mg/dl the clot dissolution time is minimum i.e. 58 and 66 min for aqueous and alcoholic extracts respectively<sup>[20]</sup>.

**6.19 Antipyretic Activity:** The antipyretic activity of ethanolic and aqueous extracts of *Terminalia bellirica* fruits (200 mg/kg, p.o.) was studied in brewer's yeast-induced fever models in mice and rats. Both extracts showed a significant inhibition of elevated body temperature when compared to corresponding control<sup>[21]</sup>.

**6.20 Antimutagenic Activity:** Water, acetone, and chloroform extracts of *Terminalia bellerica* were examined for their antimutagenic potency using the Ames Salmonella/microsome assay. Acetone extract exhibited variable inhibitory activity of 65.6%, and 69.7% with 4-O-nitrophenylenediamine (NPD) and sodium azide, respectively (as direct-acting mutagens), and 81.4% with 2-aminofluorene (2AF) (an S9-dependent mutagen), in the preincubation mode of experimentation. Inhibition with chloroform and water extracts was rather insignificant<sup>[22]</sup>.

## 7. Conclusion

Medicinal plants have been identified and used throughout human history. The study of traditional human uses of plants, is recognized as an effective way to discover future medicines. The use of herbs to treat diseases is almost universal among non- industrialized societies and is often more affordable than purchasing modern pharmaceuticals. Crude extracts of various parts of *Terminalia bellerica* plant have been found to contain constituents such as Glucoside, Gallo-tannic acid, colouring matter, resins and agreeenish yellow oil. Ellagic acid, gallic acid, lignans, 7-hydroxy 3'4' flavone and anolignan B. Tannins, ellagic acid, ethyl gallate, galloyl glucose and chebulic acid, phyllembin,  $\beta$ - sitosterol mannitol, glucose, fructose and rhamnose. These compounds are believed to be responsible for the pharmacological activities such as antimicrobial, antioxidant, antisalmonella, hepatoprotective, antispasmodic and bronchodilatory activities. Therefore, this plant is significantly used for the treatment and prevention of diseases. Further studies should be carried out for this plant to discover the unrevealed part of it which may serve for the welfare of mankind.

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