

**OROXYLUMINDICUM: A REVIEW**Dr. Ved Parkash Sharma<sup>1\*</sup> and Dr. Om Prakash Sharma<sup>2</sup><sup>1</sup>PG Scholar Department of Dravyagunavigyan, Sri Ganga Nagar College of Ayurvedic science and Hospital, Tantia University, Sri GangaNagar-33500, India.<sup>2</sup>Professor and HOD Department of Dravyagunavigyan, Sri Ganga Nagar College of Ayurvedic science and Hospital, Tantia University, Sri GangaNagar-33500, India.**\*Corresponding Author: Dr. Ved Parkash Sharma**

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**ABSTRACT**

*Oroxylum indicum* (Bignoniaceae), also known as Sonapatha or Shyonaka is commonly used herbal medicine in Ayurvedic system. Roots, leaves and stems of *Oroxylum indicum* have been used as a single drug or as a component of certain compound drug preparations in the Indian Ayurvedic system of medicine for treatment of various disorders as well as used as atonic and Rasayana drug. It contains flavonoids like chrysin, baicalin and Oroxylin-A. Various studies indicated that sonapatha possesses anticancer, antioxidant, hepato protective and immune modulatory properties mainly. Various other effects like antibacterial, analgesic and gastro-protective properties of sonapatha have also been reported. It is a tree that is found generally in damp region. In the present review an attempt has been made to compile and critically analyse various published reports on *Oroxylum indicum*.

**INTRODUCTION**

*Oroxylum indicum* also known as 'Sonapatha' is an important herb in Ayurvedic medicine and indigenous medical system for over thousands of years.<sup>[1]</sup> *Oroxylum indicum* has been used as a single drug or as a component of certain poly-herbal drug preparations in Indian system of medicine i.e. Ayurveda. It is active ingredient of well known Ayurvedic formulations like Chyavanprash, Dashmularista etc.<sup>[2]</sup> The root bark and stem bark possess anti-allergic properties and are used in treating allergic disease, urticaria, jaundice, asthma, sore throat, laryngitis, hoarseness, gastralgia, diarrhoea, dysentery, infantile, erythema and measles.<sup>[3-4]</sup> The normal dose is reported 8 to 16 g of bark in the form of decoction, extract or powder.<sup>[4-5]</sup> The seeds are active in chronic cough and gastralgia: 5 to 10 g daily in the form of decoction or powder and also used as purgative. Alcohol maceration of fresh bark is applied externally for lacquer allergic dermatitis. The fruits of *Oroxylum indicum* are acrid, sweet, stomachic, anthelmintic, and good in diseases of the heart and the throat, piles, bronchitis, used as an expectorant, improves the appetite, useful in leucoderma.<sup>[2,6-10]</sup>

**BOTANICAL DESCRIPTION**

It is a tree which can attain a height of 12 meter

(40 feet). The large leaf stalks wither and fall off the tree and collect near the base of the trunk, appearing to look like a pile of broken limb bones. The tree is a night-bloomer and flowers are adapted to natural pollination by bats. They form enormous seed pods that hang down from bare branches. Those long fruits curve downward and resemble the wings of a large bird or dangling sickles or swords in the night. The seeds are round with papery wings. Bark is off brown in color. Leaves are 2 to 4 inch long, broad, leaflets are 5 inch long and 3 to 4 inch broad having harpedges. The flowers stalks are one foot long. The flowers are purple in color. Fruits are 1 to 3 foot long, 2 to 4 inch broad. Seeds are flat and are 3 inch in length and 2 inch in width. The flowers are born in rainy season and fruit appears in December to March.<sup>[2-3,5]</sup>

**GEOGRAPHICAL DISTRIBUTION**

*Oroxylum indicum* is native to the Indian subcontinent, in the Himalayan foothills with a part extending to Bhutan and southern China, in Indo-China and the Malaysia ecozone. It is diversely available in the forest of National Park in Assam, India, reported from Sri Lanka (Ceylon).<sup>[5]</sup>

**TAXONOMICAL CLASSIFICATION<sup>[4]</sup>**

Kingdom	:	Plantae
Division	:	Magnoliophyta
Class	:	Magnoliopsida

Order : Lamiales  
 Family : Bignoniaceae  
 Genus : *Oroxylum*  
 Species : *indicum*

#### SYNONYMS<sup>[9-10]</sup>

Sansk : Prthsuimba, Katvanga  
 Hindi : Sonapatha, Syonak, Tentoo  
 Eng : Indian trumpet flower  
 Beng : Sonagachh  
 Guj : Tentoo  
 Punj : Tatpaling, Talvarphali  
 Mar : Tentoo  
 Tamil : Peruvaagai

According to Ayurveda it contains<sup>[11-13]</sup>

Gunna (Properties) – laghu (light), tikshan (sharp) and ruksha (dry). Rasa (Taste) – madhur (sweet), tikta (bitter)

Virya (Potency) – ushan (hot)

#### CHEMICAL CONSTITUENTS

The chemical constituents of *Oroxylum indicum* are always of an interest for the researcher. A number of secondary metabolites like flavonoids, glycosides, alkaloids, tannins, terpenoids etc. have been reported from various parts of the plant.

The leaves have been reported containing flavones and their glycosides baicalein and scutellarein. Leaves also contain an anthraquinone, aloemodin.<sup>[9,17]</sup>



FIGURE1: LOWER SOF OROXYLUM INDICUM.



FIGURE2: LEAVES OF OROXYLUM INDICUM.



FIGURE3: FRUIT OF OROXYLUM INDICUM.

- Bark of the root is reported with chrysin, baicalein and oroxylin-A. Bark also gave dihydrobaicalein. Heart wood yield ed beta-sitosterol and anisoflavone, prunetin. The bark also contains traces of alkaloid, tannic acid, sitosterol and galactose.<sup>[14-15]</sup>
- Its root and stem contains three flavones named oroxylin A (5, 7-dihydroxy-6-methoxyflavone), baicalein (5, 6, 7-trihydroxyflavone) and chrysin (5, 7-dihydroxyflavone). It also contains pterocarpan and rhodioside with p-hydroxyphenylethanol and cyclohexanol.<sup>[16-18]</sup>
- Four flavonoids, chrysin, baicalein, baicalein-7-O-glucoside, baicalein-7-O-diglucoside (Oroxilin B) and one unknown flavonoid have also been isolated from the seeds of *Oroxylum indicum*.<sup>[19]</sup> Seeds also contain hiny oil, the yield of which was 20%.<sup>[2]</sup>
- In Indian system of medicine the root, bark, stem and leaf are prescribed for snake bite.<sup>[2]</sup>
- Leaves are used externally to treat an enlarged spleen and also to alleviate headaches and ulcers and also reported for its analgesic and antimicrobial activity.<sup>[20]</sup>
- In various tribes of India, bark and seeds of the plant are used in fever, pneumonia and repair of troubles.<sup>[21-22]</sup> It is also used to cure various stomach disorders.<sup>[23]</sup>
- In Nepal a root decoction is used in diarrhea and dysentery. Seeds are used as a digestive. A seed paste is applied to treat boils and wounds. The root is used as astringent, anti-inflammatory, aphrodisiac, expectorant, anthelmintic and tonic. The bark is diuretic and stomachic and useful in diarrhea and dysentery. Root bark and seeds are carminative, stomachic, tonic, diaphoretic and astringent. Root bark is also used to treat bile problems, cough, diarrhea, and dysentery.<sup>[24]</sup> It is also used in a formulation used for nootropic activity.<sup>[25]</sup>

## PHARMACOLOGICAL REPORTS

Although all of the pharmacological and nonpharmacological investigations have been carried out on the plant and its phytoconstituents. A summary of the findings of the studies is presented below.

## ETHNOMEDICINAL USES

- The root bark of the plant is astringent, bitter, pungent; astringent to the bowels, cooling, aphrodisiac, tonic, increases appetite, useful in "vata", biliousness, fevers, bronchitis, intestinal worms, vomiting, dysentery, leucoderma, asthma, inflammation, anal troubles. It is used to treat diarrhea, dysentery, diaphoretic, and rheumatism.<sup>[2-3]</sup> Paste prepared from sesame oil (*Sesamum indicum*) and the powdered bark of the root is given as a digestive tonic. The seeds are purgative and take no rest to treat throat infections and hypertension.<sup>[20]</sup>
- The fruits are astringent, sweet; stomachic, anthelmintic; effective in diseases of the throat and heart, piles, bronchitis, used as an expectorant; improves the appetite; useful in leucoderma.<sup>[6-9]</sup>

## ANTI-INFLAMMATORY ACTIVITY

The aqueous extract of leaves of *Oroxylum indicum* has been reported to possess significant anti-inflammatory activity. The anti-inflammatory activity has been studied *in vivo* in a carrageenan induced rat paw edema model and it was reported that the aqueous extract of *Oroxylum indicum* leaves exhibited significant anti-inflammatory activity at a dose level of 150 mg/kg body weight and 300 mg/kg body weight. *Oroxylum indicum* aqueous extract at a dose of 300 mg/kg body weight showed maximum anti-inflammatory activity. However, the activity produced by both the doses was less effective than the reference standard diclofenac sodium. Extract at both doses showed significant anti-inflammatory activity at 5 hr. against carrageenan injection suggesting that the extract predominantly inhibits the release of prostaglandin-like substances. In conclusion, leaves of *Oroxylum indicum* showed anti-inflammatory activity which may be attributed to the presence of different chemical constituents present within.<sup>[26]</sup> A number of flavonoid compounds have also been reported in previous studies. Anti-inflammatory flavonoids present in the plant may be responsible for this activity.

Aqueous and alcoholic extracts were tested using three different *in vitro* systems for effect relevant to anti-inflammatory activity of stem bark of *Oroxylum indicum*. The aqueous extracts of *O. indicum* significantly reduced myeloperoxidase release. In the rat hind paw edema test, extracts also showed significant activity.<sup>[27]</sup> All these findings

suggest, *Oroxylum indicum* may be useful in management of chronic inflammatory conditions like arthritis.

## ANTI-HEPATOTOXIC ACTIVITY

Leaves of *Oroxylum indicum* are widely used as a prophylaxis for liver disorders in the Indian system of medicine. Tenpeet *et al.* reported anti-hepatotoxic activity of various extracts of *Oroxylum indicum* against CCl<sub>4</sub> induced hepatotoxicity. Petroleum ether, chloroform, ethanolic and aqueous extracts were administered to diseased animals (rats) at a dose of 300 mg/kg body weight and serum enzyme levels were observed. All the test groups showed a significant reduction in SGOT, SGPT, ALP, total bilirubin content and a significant increase in the level of total protein was observed in CCl<sub>4</sub> and *Oroxylum indicum* treated rats. Among all the extracts, the ethanolic extract was found to be more effective.<sup>[28]</sup> Free radical scavenging activity was also reported and the hepatoprotective action of these extracts was likely to be due to its ability to scavenge free radicals and induce microsomal enzymes thereby inhibition of the lipid peroxidation induced by CCl<sub>4</sub>. The study scientifically proved the folklore use of *Oroxylum indicum* in liver disorders and as an ingredient in various Ayurvedic formulations used in liver disorders.

## ANTHELMINTIC ACTIVITY

Jessica *et al.* evaluated the anthelmintic activity of *Oroxylum indicum* against the strongyle eggs *in vitro* and compared it to ivermectin, one of the most effective deworming agents. At a dose of 2 × 10<sup>-5</sup> g/mL and greater, hatching of the strongyle eggs was delayed using *Oroxylum indicum*. 0% hatching was achieved at 2 × 10<sup>-1</sup> g/mL *Oroxylum indicum*. At a dose of 2 × 10<sup>-4</sup> g/mL and greater, 0% viability of the strongyle eggs and larvae was achieved. The results of the study suggested that *Oroxylum indicum* may be an appropriate anthelmintic against strongyles.<sup>[29]</sup>

## ANTI-CANCER ACTIVITY

Various studies have proved the anticancer potential of *Oroxylum indicum* using various models. Narisa *et al.* extracted *Oroxylum indicum* with 95% ethanol and tested for cytotoxic effects determining the anti-proliferative effects on Hep2 cell lines. Cell proliferation was measured using a colorimetric method based on the ability of metabolic active cells to cleave the yellow tetrazolium salt XTT to an orange formazan dye and soluble formazan dye was directly quantified using a scanning multi-well spectrophotometer (ELISA plate reader). Ethanol exhibited cytotoxic activity against the Hep2 cell lines at a concentration of 0.05%.<sup>[30]</sup>

Roy *et al.* Reported the *in vitro* effects of baicalein on the viability and induction of apoptosis in the HL-60 cell line was investigated. The cell viability after treating with baicalein for 24h was quantified by counting viable cells using trypan blue staining. The results showed that baicalein caused a 50% inhibition of HL-60 cell growth on concentrations of 25–30  $\mu\text{M}$ . The inhibition of proliferation of HL-60 cells due to 36–48h exposure to 10 or 20  $\mu\text{M}$  baicalein was associated with the accumulation of cells in G2/M phases. However, proliferation inhibition at a higher dose may be associated with induction by apoptosis and terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL). The results indicate that baicalein has anti-tumor effect on human cancer cells, and *Oroxylum indicum* extract could be used in supplementary cancer therapy.<sup>[31]</sup>

Nakahara *et al.* reported that methanolic extract of *Oroxylum indicum* strongly inhibited the mutagenicity of Trp-P-1 in an Ames test. The major antimutagenic constituent was identified as baicalein with an  $\text{IC}_{50}$  value of  $2.78 \pm 0.15 \mu\text{M}$ . The potent antimutagenicity of the extract was correlated with the high content ( $3.95 \pm 0.43\%$ , dry weight) of baicalein. Baicalein acted as an antimutagen since it inhibited the N-hydroxylation of Trp-P-2.<sup>[32]</sup>

Tepsuwan *et al.* reported the *in vivo* genotoxic activity and cell proliferative activity in stomach mucosa of male F344 rats by *in vivo* short-term methods after oral administration of a nitrosated *Oroxylum indicum* Ventrifraction, which had been found to be mutagenic with out S9 mix to Salmonella typhimurium TA 98 and TA 100. Administration of the nitrosated *Oroxylum indicum* Ventrifraction at doses of 1 and 2 g/kg body weight induced dose-dependent DNA single-strand scission in the stomach pyloric mucosa 2 h after its administration: a dose of 2 g/kg body weight induced an 18-fold increase in the DNA elution rate constant. Administration of the nitrosated *Oroxylum indicum* fraction at doses of 0.7–2.8 g/kg body weight also induced dose-dependent increases, up to 11-fold, in replicative DNA synthesis in the stomach pyloric mucosa 16 h after its administration. Moreover administration of the nitrosated *Oroxylum indicum* fraction at doses of 0.25–2.0 g/kg body weight induced dose-dependent increases, up to 100-fold, in ornithine decarboxylase activity in the stomach pyloric mucosa within a maximum 4h after its administration. These results demonstrate that the nitrosated *Oroxylum indicum* fraction has genotoxic and cell proliferative activity in the pyloric mucosa of rat stomach *in vivo*.<sup>[33]</sup>

Leticia *et al.* reported that extract of *Oroxylum indicum* showed the toxicity on tumor cell lines tested, with

an  $\text{IC}_{50}$  value 19.6  $\mu\text{g/ml}$  for CEM, 14.2  $\mu\text{g/ml}$  for HL-60, 17.2  $\mu\text{g/ml}$  for B-16 and 32.5  $\mu\text{g/ml}$  for HCT-8. On the search in eggs, it also inhibits the progression of cell cycles since the first cleavage ( $\text{IC}_{50} = 13.5 \mu\text{g/ml}$ ). On the basis of all these findings it can be concluded that extracts of *Oroxylum indicum*, could be considered as potential sources of anti cancer compounds.<sup>[34]</sup>

#### IMMUNOSTIMULATING ACTIVITY

The immune modulatory activity and the mechanism of action of then-butanol fraction (100mg/kg body weight, per os, once daily for 22 consecutive days) of the root bark of *Oroxylum indicum*, was reported by Zaveri *et al.* in rats using measures of immune responses to sheep red blood cells (SRBC haemagglutinating antibody [HA] titer) and delayed-type hypersensitivity (DTH) reactions. In response to SRBC, treatment with then-butanol fraction caused a significant rise in circulating HA titers during secondary antibody responses, indicating potentiation of certain aspects of the humoral response. The treatment also resulted in a significant rise in paw edema formation, indicating increased host DTH response. Additionally, the antioxidant potential of the drug was exhibited by significant reductions in whole blood malondialdehyde content along with a rise in the activities/levels of superoxide dismutase, catalase and reduced glutathione. Furthermore, histopathological analysis of lymphoid tissues showed an increase in cellularity, e.g., T-lymphocytes and sinusoids, in the treatment group. In a triple antigen-mediated immunological edema model, the extent of edema raised in drug-treated rats was greater compared to that in control rats, thus confirming enhanced DTH reactions in response to the drug treatment. Based on all these findings, the reported immunomodulatory activity of an active fraction of *O. indicum* might be attributed to its ability to enhance specific immune responses (both humoral and cell-mediated) as well as its antioxidant potential.<sup>[35]</sup> This study also justifies the use of plant in various immunomodulatory formulations of Ayurveda like Chyavanprash etc.

#### ANTI MICROBIAL ACTIVITY

The anti-microbial activity of various extracts of *Oroxylum indicum* has been screened against fourteen pathogenic bacteria (five gram-positive and nine gram-negative) and seven pathogenic fungi by Kawsar *et al.* using disk diffusion method. The crude ethyl acetate extract showed mild to moderate activity against all bacteria and fungi whereas the methanolic extract showed little activity against bacteria but moderate activity against fungi. The minimum inhibitory concentration of two isolated flavonoid compounds from *O. indicum* were

determined against *Bacillus subtilis*, *Staphylococcus aureus*, *Escherichia coli* and *Shigella dysenteriae* and the values were found to be between 64–128 µg/ml. A study by Thatoi et al. further confirmed the activity by using different strains.<sup>[36–37]</sup> Ali et al. (1998) studied the effect of dichloromethane extract of *Oroxylum indicum* against dermatophytes and wood rot fungi and reported a strong antifungal activity of dichloromethane extract of *Oroxylum indicum*.<sup>[38]</sup>

#### GASTRO-PROTECTIVE ACTIVITY

Zaveri et al. reported the gastroprotective activity of 50% alcoholic extract of root bark of *Oroxylum indicum* and its different fractions viz. petroleum ether, chloroform, ethylacetate and *n*-butanol fractions in ethanol-induced gastric mucosal damage. *n*-butanol fraction was also studied in Water Immersion Plus Restraint Stress (WIRS)-model. Alcoholic extract (300 mg/kg) and its different fractions (at a dose of 100–300 mg/kg) showed significant reduction in gastric ulceration against ethanol-induced gastric damage. Out of all these fractions, *n*-butanol fraction showed significant maximum inhibition of gastric lesions. In WIRS-model, pretreatment with *n*-butanol fractions showed significant anti-ulcer and antioxidant activity in gastric mucosal homogenates, where it reversed the increase in ulcer index, lipid peroxidation and decrease in superoxide dismutase, catalase and reduced glutathione levels induced by stress. This study reveals significant gastroprotective effect of *n*-butanol fraction against both ethanol and WIRS-induced gastric ulcers in rats.<sup>[39]</sup> Flavonoids present in *Oroxylum indicum* Vent. was found to be responsible for its gastro-protective activity.<sup>[40]</sup>

#### CONCLUSION

*Oroxylum indicum* is a highly placed drug in the Ayurvedic medicine. It is one of the most versatile plants having a wide spectrum of medicinal activities. This medicinal plant is the unique source of various types of compounds having diverse chemical structure and nature. Quite less scientific work has been conducted on the possible medicinal applications of these compounds and hence extensive investigation is desirable to exploit their therapeutic utility. Although crude extracts from various parts of *Oroxylum indicum* have been assigned various medicinal applications from time immemorial, the probability of converting these promising activities into modern drugs can be explored further only after extensive investigation of its bio activity of responsible constituents, mechanism of action, and toxicity and after proper standardization. As this approach would be in line with the global scenario which is now changing towards the use of plant products, that are backed by ethno traditional medicinal use, which are comparatively non-toxic than currently available

marketed drugs of other systems.

#### REFERENCES

- Joshi KC, Prakash L, Shah RK. Chemical examination of the roots of *Tabebuia rosea* and heart wood of *Oroxylum indicum*. *Plant Med*, 1977; 31: 257–8.
- Kirtikar KR, Basu BD. *Indian Medicinal Plants*. Oriental Enterprises, Dehradun, 42001; 1105–1107.
- Paranjpe Prakash. *Indian Medicinal Plants*. (Chaukhamba Sanskrit Pratishthan, Delhi, 2005; 248–9.
- Oroxylum indicum: herb and benefit*. Available at <http://www.w3.org/1999/xlink> <http://www.ayushveda.com/herbs/Oroxylum-indicum.html#1> (accessed November 25, 2009).
- Oroxylum indicum* available at <http://www.w3.org/1999/xlink> [http://en.wikipedia.org/wiki/Oroxylum\\_indicum.html](http://en.wikipedia.org/wiki/Oroxylum_indicum.html) (Accessed March 15, 2010).
- Chopra RN, Nayar SL, Chopra IC. *Glossary of Indian Medicinal Plants*. National Institute of Science Communication and Information Resources, New Delhi, 2002; 182.
- Drury CH. *Ayurvedic Useful Plants of India*. Asiatic Publishing House, 2006; 360.
- Nadkarni AK. *Indian Materia Medica*. Bombay Popular Prakashan, Mumbai, 1982; 876–77.
- Khare CP. *Indian Medicinal Plants*. Springer Science Business Media, LLC, 2007; 453.
- Ayurvedic Pharmacopoeia of India, Part 1<sup>st</sup>*, Vol.3, Government of India, Ministry of Health and Family Welfare, 183–4.
- Sharma M. *Shushrut Samhita: Chikitsa Sthan*. Vol.2, Khemraj Shrikrishnadass Press, Bombay, 2003; 924–926.
- Sharma S. *Ashtang Hridaya: Chikitsa Sthan*. Khemraj Shrikrishnadass Press, Bombay, 1996; 506–10.
- Mishra KN. *Bhavprakash*. Khemraj Shri Krishna dass Press, Bombay, 2004; 229: 1021.
- The Wealth of India (Raw Materials)*, Vol.3, Council of Scientific and Industrial Research, New Delhi, 316–7.
- Yin WG, Li ML, Kang C. Advances in the studies of *Oroxylum indicum*. *Zhongguo Zhong Yao ZaZhi*, 2007; 32(19): 1965–70.
- Vasanth S, Natarajan M, Sundaresan R, Rao RB, Kundu AB. Ellagic acid from *Oroxylum indicum* Vent. *Indian Drugs*, 1990; 28(11): 507.
- Dey AK, Mukherjee P, Das PC, Chatterjee A. Occurrence of Aloe-emodin in the leaves of *Oroxylum indicum* Vent. *Indian Journal of*

- Chemistry*, 1978; 16: 1042.
18. Theobald WL, Dassanayake MD, Fosberg MR. *A Revised Hand book to the Flora of Ceylon*. Amerind Publishing Co. Pvt. Ltd., New Delhi, 1981.
  19. Chen LJ, David EG, Jones J. Isolation and identification of four flavonoid constituents from the seeds of *Oroxylum indicum* by high-speed counter-current chromatography. *Journal of Chromatography A*, 2003; 988(1): 95–105.
  20. Singh HB, Prasad P, Rai LK. Folk Medicinal Plants in the Sikkim Himalayas of India. *Asian Folklore Studies*, 2002; 61: 295–310.
  21. Panghal M, Aryal V, Yadav S, Kumar S, Yadav JP. Indigenous knowledge of medicinal plants used by Saperas community of Khetawas, Jhajjar District, Haryana, India. *Journal of Ethnobiology and Ethnomedicine*, 2010; 6: 4.
  22. Patil GG, Mali PY, Bhadane VV. Folk remedies used against respiratory disorders in Jalgaon district, Maharashtra. *Natural Product Radiance*, 2008; 7(4): 354–8.
  23. Rout SD, Panda T, Mishra N. Ethno-medicinal Plants Used to Cure Different Diseases by Tribals of Mayurbhanj District of North Orissa. *EthnoMedicine*, 2009; 3(1): 27–32.
  24. Kunwar RM, Uprety Y, Burlakoti C, Chowdhary CL, Bussmann RW. Indigenous Use and Ethnopharmacology of Medicinal Plants in Far-west Nepal. *Ethnobotany Research & Applications*, 2009; 7: 5–28.
  25. Maciuk A, Bouchet MJ, Mazars G, UM BH, Anton R. *Nootropic (med-hya) plants from ayurvedic Pharmacopoeia*. *Etudes chimiques et pharmacologiques*, 402–11.
  26. Upananlawar A, Tenpe CR, Yeole YG. Anti-inflammatory activity of aqueous extract of *Oroxylum indicum* vent. Leaves extract-preliminary study. *Pharmacologyonline*, 2009; 1: 22–6.
  27. Laupattarakasem P, Houghton PJ, Houlst JR, Itharat A. An evaluation of the activity related to inflammation of our plants used in Thailand to treat arthritis. *Journal of Ethnopharmacology*, 2003; 85(2–3): 207–15.
  28. Tenpe CR, Aman Upananlawar, Sushil Burle, Yeole YG. *In vitro* antioxidant and preliminary hepato protective activity of *Oroxylum indicum* vent leaf extracts. *Pharmacologyonline*, 2009; 1: 35–43.
  29. Downing JE. Anthelmintic Activity of *Oroxylum indicum* Against *Equine Strongyles* *in vitro* Compared to the Anthelmintic Activity of Ivermectin. *Journal of Biological Research*, 2000; Vol.1:
  30. Narisa K, Jenny MW, Heather MAC. Cytotoxic Effect of Four Thai Edible Plant on Mammalian Cell Proliferation. *Thai Pharmaceutical and Health Science Journal*, 2006; 1(3): 189–95.
  31. Roy MK, Nakahara K, Na TV, Trakoontivakorn G, Takenaka M, Isobe S et al. Baicalein- A flavonoid extracted from a methanolic extract of *Oroxylum indicum* inhibits proliferation of a cancer cell line *in vitro* via induction of apoptosis. *Pharmazie*, 2007; 62(2): 149–53.
  32. Nakahara K, Onishi KM, Ono H, Yoshida M, Trakoontivakorn G. Antitumorigenic activity against trp-P-1 of the edible Thai Plant: *Oroxylum indicum* Vent. *Biosci Biotechnol Biochem*, 2001; 65(10): 2358–60.
  33. Tepsuwan A, Furihata C, Rojanapo W, Matsuhima T. Genotoxicity and cell proliferative activity of a nitrosated *Oroxylum indicum* Vent fraction in the pyloric mucosa of rat stomach. *Mutat Res*, 1992; 281(1): 55–61.
  34. Lotufo LVC, Khan MTH, Ather A, Wilke DV, Jimenez PC, Pessoa C et al. Studies of the anticancer potential of plants used in Bangladesh folk medicine. *Journal of Ethnopharmacology*, 2005; 99: 21–30.
  35. Zaveri M, Gohil P, Jain S. Immunostimulant Activity of n-Butanol Fraction of Root Bark of *Oroxylum indicum* Vent. *Journal of Immunotoxicology*, 2006; 3(2): 83–99.
  36. Kawsar U, Sayeed A, Islam A, Abdur RA, Khatun S, Khan A et al. Biological activity of Extracts and two Flavonoids from *Oroxylum indicum* Vent. (Bignoniaceae). *Online journal of Biological science*, 2003; 3(3): 371–5.
  37. Thatoi HN, Panda SK, Rath SK, Dutta SK. Antimicrobial activity and ethnomedicinal uses of some medicinal plants from Similipal biosphere reserve Orissa. *Asian Journal of Plant Sciences*, 2008; 7(3): 260–7.
  38. Hari Babu T, Manjulatha K, Suresh Kumar G, Hymavathi A, Tiwari AK, Purohit M et al. Gastro protective flavonoid constituents from *Oroxylum indicum* Vent. *Bioorganic & Medicinal Chemistry Letters*, 2010.