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Research Article

Antiulcer Activity of the Root Bark of *Oroxylum indicum*. Against Experimental Gastric Ulcers

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Materials and Methods

Procurement of plant material and extraction procedure

The fresh root bark of *Oroxylum indicum*. was collected in January 2005 from Vanaushadhi Ektrikaran Udyan, Ahwa, Dang Forest, Gujarat, India. The authentication of this plant was established by the taxonomist of Gujarat Ayurved University, Jamnagar, India, and a voucher specimen (404) was deposited in the Department of Pharmacognosy and Phytochemistry, L. M. College of Pharmacy, Ahmedabad, India. The root bark was sun-dried and powdered to 60 mesh. The powder of root bark after defatting with petroleum ether (0.32% w/w) was dried and then moistened with ammonia solution and extracted with chloroform (0.78% w/w), ethyl acetate (1.52% w/w), and n.-butanol (1.68% w/w), successively. The dried fractions were stored in an air-tight borosil glass container until further use.

Drugs and chemicals

Omeprazole was obtained from Zyclus Research Centre, Ahmedabad, India. All different organic solvents and reagents used for the current study were of analytical grade (AR) and obtained from S.D. Chem. Pvt. Ltd. (Mumbai, India). The standard baicalein was obtained from Sigma-Aldrich (St. Louis, MO, USA). Fresh drug solutions were prepared in 1% carboxy methylcellulose (CMC) and were administered orally.

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experiments. The stomachs were removed, opened along the greater curvature, washed with saline, and examined using a 6.4 binocular magnifier. Lesions were assessed by two unbiased observers.

Methodology

The animals were divided into following groups of six.

- Group I (control): Rats received only aqueous suspension of 1% CMC vehicle with respect to the individual ulcerogenic procedure.
- Group II (drug treatment): Rats received the following treatments: 50% alcohol extract, petroleum ether, chloroform, ethyl acetate, and n.-butanol extracts (100–300 mg/kg, p.o.).
- Group III: Rats received standard omeprazole (20 mg/kg, p.o.) 1 h before the ulcerogenic procedure.

Ethanol-induced gastric mucosal damage

Gastric lesions were induced by 1 ml absolute ethanol in 24-h fasted rats as per the method of Robert ([1979](#)). In the treatment group, drug extracts in 1% CMC solution were administered orally 1 h before the administration of ethanol. Animals were

sacrificed and the stomachs were opened along the greater curvature and washed with ice-cold saline (pH 7.4). The stomachs were then examined for the presence of lesions. The area of the lesion was measured in terms of the diameter of the lesion (X). For circular lesions, five of the diameters were measured and the average diameter was used. The index = $10/X$, where X is the diameter of the lesion. The drug extracts were evaluated. The stomachs were washed with saline (10 mM, pH 7.4) using Remi-oximide for the estimation of superoxide dismutase (SOD) activity. The micro-



method of Mishra et al. ([1973](#)). Catalase (CAT) activity was measured according to the method of Aebi ([1974](#)). The reduced glutathione (GSH) was determined by the method of Beutler et al. ([1963](#)). The protein concentration in all samples was determined by the method of Lowry ([1951](#)).

Pylorus-ligation (PL) model

Rats fasted for 24 h were anesthetized with ether, and a portion of abdomen was opened by a small midline incision below the xiphoid. The pylorus portion of the stomach was lifted and ligated (with care being taken not to occlude blood vessels) by the method of Shay et al. ([1945](#)). The stomach was closed with interrupted sutures. Six hours after the pylorus ligation, animals were sacrificed. The stomach was dissected and the contents collected, measured, centrifuged, and subjected to biochemical analysis described below. Parameters investigated include: a ulcer index (UI) as described earlier, b acid secretory parameters, and c mucoprotective parameters. Acid secretory parameters include measurement of volume of gastric secretion, total acidity determined by titrating against 0.01 N sodium hydroxide to pH 8.0 using phenolphthalein as an indicator (Hawk et al., [1954](#)), and total acid output (product of total acidity and volume of gastric secretion). Further, pepsin activity was determined using hemoglobin as a substrate, according to the modified method of Debnath et al. ([1974](#)). Total carbohydrates (TC) (Nair, [1976](#)), total protein content (PR) (Lowry et al., [1951](#)), n

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studies (TLC, HPLC). TLC co-chromatography was performed on the petroleum ether fraction, the hydrolyzed n.-butanol fraction, and standard baicalein.

Method of TLC analysis

Ten microliters of each sample solution was spotted on the TLC plate (precoated with silica gel 60 F₂₅₄, thickness 0.2 mm, 20 × 20 cm) (E Merck, Darmstadt, Germany) along with a standard solution of baicalein. Chromatogram was developed using chloroform:ethyl acetate:formic acid (10:8:2) as a mobile phase and visualized using natural product poly ethylene glycol (NP/PEG) reagent. On the basis of the TLC study, an HPLC method was developed for the quantification of baicalein in both the active fractions and development of fingerprints of the same.

HPLC analysis

Chemicals: Methanol (HPLC grade), acetonitrile (HPLC grade), water (HPLC grade), trifluoroacetic acid (TFA) (analytical grade), baicalein (Sigma-Aldrich, Powai, Mumbai). HPLC was performed on a Shimadzu 2010 C (Tokyo, Japan), equipped with a C-18 Hypersil BDS column (250 × 4.6 mm, 5 μm). The instrument was operated under the following conditions: UV visible detector 254 nm, flow rate of 1.0 ml/min, retention time 40.5 min, injection volume 10 μl, and mobile phase A, water (pH = 2.70 adjusted with dilute H₂SO₄), mobile phase B, acetonitrile [diluent methanol: water pH = 3.0 with TFA (8:2)]. HPLC fingerprints were compared with the standard baicalein and also with the fingerprints of the samples.

Sample

A calibration curve was constructed by plotting the peak area (ml) at different concentrations (μg/ml) of each sample against the concentration of each extract.

Valid

The HPLC method was validated for linearity, accuracy, precision, recovery, and limit of detection.

Statisti



The results were expressed in terms of mean \pm SEM. The significance of difference between mean values for the various treatments was tested using one-way analysis of variance test (ANOVA test) followed by Tukey's multiple range tests (Bolton, [1997](#)) wherever applicable to assess statistical significance of difference between the groups.

Results

Ethanol-induced gastric mucosal damage

Alcohol extract and the different fractions (300 mg/kg) showed a significant reduction in the ulcer index when compared with the control group, and results were comparable with the omeprazole-treated rats ([Table 1](#)). Reduction in the ulcer index was found to be maximum with both the n.-butanol (99.5%) and petroleum ether (96.0%) fractions at 100 mg/kg dose level as compared with control and omeprazole (99.5%) treatment ([Table 2](#)).

Table 1. Effect of different extracts (300 mg/kg, p.o.) of *Oroxylum indicum*. on ethanol-induced gastric mucosal damage in rats.

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Table 3. Effect of different extracts of *Oroxylum indicum*. (p.o.) on lipid peroxidation and antioxidant enzymes against ethanol-induced gastric mucosal damage.



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Pylorus-ligation gastric ulcer model

The petroleum ether, n.-butanol fractions, and omeprazole pretreated rats showed significant reduction in the ulcer index when compared with the control group (Table 4).

Table 4. Effect of active fractions (100 mg/kg, p.o.) of *Oroxylum indicum*. on pylorus-ligated gastric ulcer model.



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Effect on acid secretory parameters

Both the active fractions of drug and omeprazole treatment showed significant decrease in the volume of gastric secretion along with significant increase in the gastric pH, as compared with control group. They also showed significant reduction in total acidity, compared with control group (Table 4).

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activity. Therefore, TC:PR ratio (mucin activity) was significantly increased by both fractions. Furthermore, the gastric mucus content was found increased in petroleum ether and n.-butanol fractions pretreated animals as compared with control group. Omeprazole treatment also showed significant rise in mucus content of gastric mucosa (Table 6).

Table 6. Effect of active fractions of *Oroxylum indicum*. (100 mg/kg, p.o.) on mucoprotective parameters in pylorus-ligated gastric ulcer model.



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Fingerprinting and estimation of baicalein

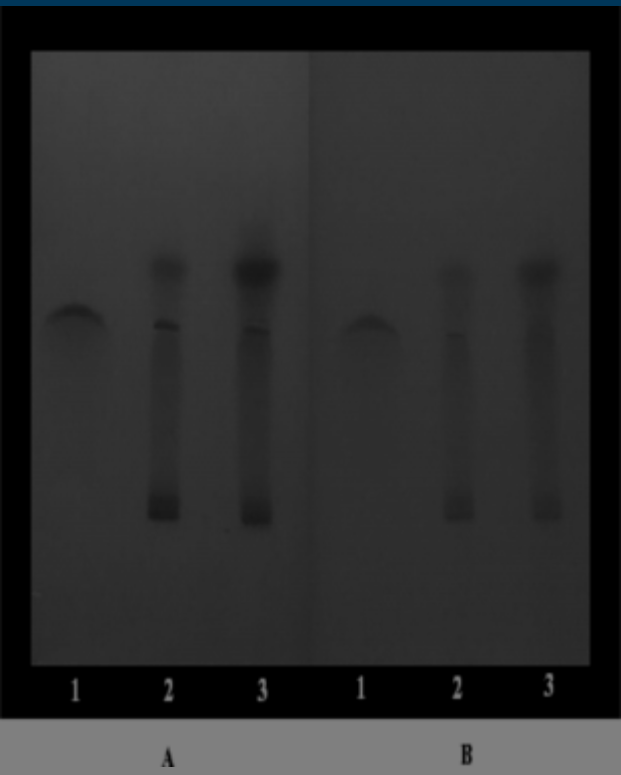
Based on the results of antiulcer activity, TLC study was aimed at checking the presence of baicalein in active fractions. Baicalein is reported to be present in stem bark and leaves of *Oroxylum indicum*. Our observations on TLC support the presence of baicalein, a major flavonoid (Fig. 1). Further, the authentic sample of baicalein resolved at 0.42 retention time (R_t), and nearly the same R_t was observed with the use of petroleum ether and hydrolyzed n.-butanol fractions (Fig. 2; Table 7). The

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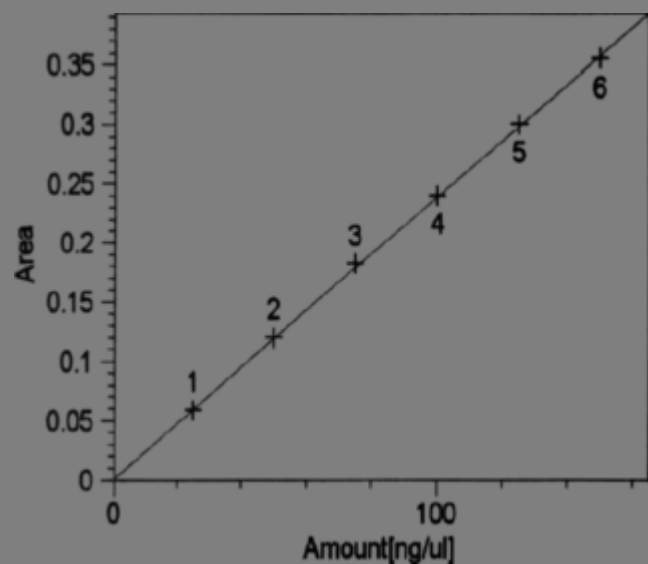
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Figure 2. HPLC chromatograms of active fractions and standard baicalein. (a) Standard baicalein, (b) petroleum ether fraction, (c) hydrolyzed n.-butanol fraction.



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Figure 3. Calibration curve of standard baicalein.



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Table 7. Percent of reference standard baicalein in active fractions of *Oroxylum indicum*..

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activity in this model along with alteration in antioxidant enzyme status. Preventive antioxidants, superoxide dismutase (SOD), and catalase (CAT) are the first line of defense against reactive oxygen species (Halliwell, [1995](#)). In addition, reduced glutathione (GSH) is a major low-molecular-weight scavenger of free radicals in the cytoplasm and an important inhibitor of free radical-mediated lipid peroxidation (Piper & Stiel, [1986](#)). It was observed in our study that the drug pretreatment resulted in significant reduction in MDA content, along with significant rise in SOD, CAT, and reduced GSH levels, suggesting their efficacy in preventing free radical-induced damage. The mechanism of antiulcer activity in this model, therefore, can be attributed to the free radical scavenging activity of this drug that in turn might lead to gastric cytoprotection.

Gastric acid and pepsin are important factors for the formation of ulcers in pylorus-ligated rats (Shay et al., [1945](#)). Increased synthesis of nucleic acids and metabolism of carbohydrates and other compensatory mechanism could also be responsible for the ulceration due to pylorus ligation (Robert et al., [1984](#)). We observed significant reduction in total acidity and pepsin activity along with significant increase in the gastric pH in drug-treated animals. Besides, there was a significant rise in mucin activity and mucus content. Therefore, it is suggested that the fractions suppressed the gastric damage caused by aggressive factors and cause increase in defensive factors in terms of gastroprotection.

Preliminary studies showed that the antiulcer activity of the flavonoid fractions was evaluated by HPLC analysis of the fractions. The HPLC analysis showed the presence of the baicalein and flavone in both the fractions. The antiulcer activity (Ng et al., [2000](#)) of the baicalein and flavone fractions was evaluated. Thus, the antiulcer activity of the baicalein and flavone fractions was evaluated. Therefore, it is suggested that the fractions suppressed the gastric damage caused by aggressive factors and cause increase in defensive factors in terms of gastroprotection.

Conclu



It is concluded that both the n.-butanol and petroleum ether fractions of *Oroxylum indicum*. possess significant antiulcer activity. There was an inhibitory effect on acid secretory mechanisms and free radical scavenging activity and a significant rise in gastric mucin activity. Further, with the help of HPLC-based profiling techniques, the antiulcer activity could be linked to a significant extent to the presence of baicalein in both fractions.

Acknowledgment

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[INFO

3. Alarcon De La Lastra C, Martin MJ, Motilva V, Jimenez M, La Casa C, Lopez A (1995): Gastro protection induced by silymarin, the hepatoprotective principle of *Silybum marianum*. in ischaemia reperfusion mucosal injury: Role of neutrophils. *Planta Med* 61: 116-119. [INFOTRIEVE], [CSA]

 | [PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

4. Anonymous (1998): *The Ayurvedic Pharmacopoeia of India*. New Delhi, Government of India, Ministry of Health and Family Welfare, Department of Indian System of Medicine and Homeopathy, 3: pp. 209-210. [CSA]

[Google Scholar](#)

5. Beutler E, Duron O, Kelly B (1963): Reduced glutathione estimation. *J Clin Med* 61: 882-889. [CSA]

[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

6. Bolton S (1997): Analysis of variance. In: Swarbrick J, ed., *Pharmaceutical Statistics: Practice and Clinical Application*, Basel, Marcel Dekker, Drug and Pharmaceutical Sciences Series, pp. 265-325.

[Google Scholar](#)

7. Bou-A... stasis
prece... rat. *Dig Dis*
Sci 33

8. Debn... on gastric
secret

9. Fridov... FOTRIEVE],
[CSA]

10. Goswa... L 34915), a



ulcers in rats and guinea-pigs. J Pharm Pharmacol 49: 195-199. [INFOTRIEVE], [CSA]

[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

1. Halliwell B (1995): Antioxidant characterization: Methodology and mechanism. Biochem Pharmacol 49: 1341-1345. [INFOTRIEVE], [CSA], [CROSSREF]

[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

2. Hawk P, Oser B, Summerson W (1954): Gastric analysis. In: Stevens E, ed., Practical Physiological Chemistry. Toronto, Blakiston, pp. 378-380.

[Google Scholar](#)

3. John AP (2001): Healing Plants of Peninsular India.. In: Bignoniaceae, Oroxyllum indicum. WallingfordUK, CABI publishing, p. 169-171.

[Google Scholar](#)

4. Kennouf S, Benabdallah H, Gharzouli K, Amira S, Ito H, Kim TH, Yoshida T, Gharzouli A (2003): Effect of tannins from Quercus suber. and species leaves on ethanol induced gastric lesions in mice. J Agri Food Chem 51: 1469-1473. [CSA], [CROSSREF]

[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

5. Kiso Y (1995): Mechanism of antihepatic damage by peroxidase. J Pharm Pharmacol 47: 1005-1010. [INFOTRIEVE], [CSA], [CROSSREF]



6. Lowry OH (1956): Protein measurement with Folin reagent. J Biol Chem 216: 1-13. [INFOTRIEVE], [CSA], [CROSSREF]



7. Mishra S, Mishra S (2007): Evaluation of the effect of epinephrine on gastric ulcers in rats. J Pharm Pharmacol 59: 3170-3175. [INFOTRIEVE], [CSA], [CROSSREF]

8. Nair BR (1976): Investigations on the venom of south Indian scorpion. *Heterometrus scaber*. Ph.D. thesis, University of Kerala, Trivendrum, p. 39.
[Google Scholar](#)


9. Ng TB, Lin F, Wang ZT (2000): Antioxidant activity of natural products from plants. *Life Sci* 68: 709–723. [CSA], [CROSSREF]
[PubMed](#) | [Google Scholar](#)

10. Oates PJ, Hakkinen JP (1988): Studies on the mechanism ethanol-induced gastric damage in rats. *Gastroenterology* 94: 9–21. [CSA]
[Web of Science ®](#) | [Google Scholar](#)

11. Peskar BM, Lange K, Hoppe U, Peskar BA (1986): Ethanol stimulates formation of leukotriene C4 in rat gastric mucosa. *Prostaglandins* 31: 283–293. [INFOTRIEVE], [CSA], [CROSSREF]
[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

12. Pihan G, Regellio C, Szabo S (1987): Free radicals and lipid peroxidation in ethanol- or aspirin-induced gastric mucosal injury. *Dig Dis Sci* 32: 1395–1401. [INFOTRIEVE], [CSA], [CROSSREF]
[PubMed](#) | [Web of Science ®](#) | [Google Scholar](#)

13. Piper I (1987): Gastric mucosal injury and its prevention by drinking and...
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14. Rober... (1987): Gastric mucosal injury...


15. Rober... (1987): Gastric mucosal injury...
 mucus... between...
 86: 670–674. [



26. Sankara S, Nair AGR (1972a): Flavonoids of the stem bark of *Oroxylum indicum*.. *Curr Sci* 41: 62-63. [CSA]

[Web of Science](#) [®] | [Google Scholar](#)

27. Sankara S, Nair AGR (1972b): Flavonoids from the leaves of *Oroxylum indicum*. and *Pajanelia longifolia*.. *Phytochemistry* 11: 439-440. [CSA], [CROSSREF]

[Web of Science](#) [®] | [Google Scholar](#)

28. Shay H, Komarov SA, Fels SS, Meraze D, Gruenstein M, Sipler H (1945): A simple method for the uniform production of gastric ulceration in rats. *Gastroenterology* 5: 43-61. [CSA]

[Web of Science](#) [®] | [Google Scholar](#)

29. Vasantha S, Natarajan M, Suderesan R, Bhima Rao R, Kundu AB (1991): Ellagic acid from root bark of *Oroxylum indicum*.. *Indian Drugs* 28: 507. [CSA]

[Google Scholar](#)

30. Warriar PK, Nambiar VPK, Ramankutty C (1995): *Oroxylum indicum*.. In: *A Compendium of 500 Species, Indian Medicinal Plants, Vol IV*. Madras, Orient Longman Ltd., pp. 186-190.

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