Peptic ulcer being one of the most uncontrolled gastrointestinal problems representing a chief health hazards in terms of morbidity and mortality. The etiology of gastroduodenal ulcers is influenced by diverse aggressive and defensive factors for example acid-pepsin secretion, mucosal barrier, mucus secretion, blood flow, cellular regeneration, and endogenous protective agents. Mucosal injury may happen when noxious factors "overwhelm" an intact mucosal protection or when the mucosal defense is somehow disrupted.

Medicinal plants are being used by mankind as a source of medicine since immemorial time. Medicinal plants are generally known as “Chemical Goldmines” as it contain a variety of natural chemicals, which are acceptable to human being and animal systems. A medicinal plant possesses curative properties due to the existence of various complex chemical substances of different composition known as secondary metabolites. According to World Health Organization more than 80% of the World’s population depends on traditional medicine for their primary healthcare requirements. Approximately 75% of the medicinally useful plant species produce in wild condition. Pueraria tuberose Roxb., commonly known as kudzu, is a climber with woody tuberous roots. The tubers are consumed as supplementary food and for control by assured Indian tribes. Pueraria tuberose Roxb., commonly known as kudzu, Indian kudzu, or Nepalese kudzu is a climber with woody tuberculated stem. It is a climbing, coiling and trailing vine with large tuberous roots. The tubers are globose or pot-like, about 25 centimeters (9.8 in) across and the insides are white, starchy and mildly sweet. Leaves are trifoliate and alternate, while the leaflets are egg-shaped, with round base and unequal sides. They are 18 cm (7.1 in) long and 16 cm (6.3 in) wide and are hairless above. Flowers are bisexual, around 1.5 cm (0.59 in) across and blue or purplish-blue in color. The fruit pods are linear, about 2–5 cm (0.79–1.97 in) long and constricted densely between the seeds. They have silky, bristly reddish-brown hair. Seeds vary from 3 to 6 in number. Indian Kudzu or Pueraria tuberosa Linn (Fabaceae) is an important medicinal plant of the Indian traditional system of medicine that is Ayurveda, and is mentioned in the Ayurvedic Pharmacopoeia of India under the name of Vidari. It is used in traditional medicine as a fertility control agent and as an aphrodisiac, cardiotonic, diuretic and galactagogue. It has exhibited antihyperglycemic, antihyperlipidemic, and antifertility in male rats, hepatoprotective, and anti-implantation activities. It is a constituent of various formulations used as nutritive, diuretic, expectorants, and for the management of rheumatism, fever, and
bronchitis. P. tuberosa tubers are rich in isoflavonoids and the important phytoconstituents are puercarin, daidzein, genistein, puertuberosanol, and tuberosin. During the past decade, interest in these isoflavonoids has increased considerably because of the beneficial effects proposed by epidemiologists, nutritionists, and food manufacturers. These isoflavonoids could interact with milk proteins, namely, bovine serum albumin, casein micelle, and β-lactoglobulin, as has been reported in case of certain food and drug preparation containing soya isoflavonoids.

The Present work is to frame antiulcer activity of petroleum ether and ethanolic extracts of tuber of *Pueraria tuberosa* Roxb in albino rats. The plant of tuber of *Pueraria tuberosa* Roxb. Figure 1.

![Figure 1: Tubers of Pueraria tuberosa Roxb](image)

**MATERIALS AND METHODS**

**Plant material**

The plant specimens for the proposed study were collected from Chopda Tehsil (Adawad) MS, India in the month of April 2019 care was taken to select healthy plants and for normal organs. The plant was authenticated by Botanical Survey of India (BSI), Pune, Maharashtra, India. A voucher specimen (No. NIPPUT1) was deposited at B.S.I., Pune, India.

**Animals**

Male wistar rats weighing about 120-180 gm were procured. The animals were kept under a conventional light regimen at room temperature (about 250 C) and humidity. Animals were housed in polypropylene cages and were allowed free access to standard laboratory feed and water. All the animals have been divided into four groups and placed in separate cages, each consisting of six animals. The animals were acclimatized to the laboratory condition for one week before the onset of experiment. The Institutional Animals Ethics Committee approved the protocol vid no. NIB/ IAEC/ 11-12/ 23

**Preparation of standard drug and extract solution:**

Solution of all extracts was prepared in Tween 80.

Test drug was dissolved in distilled water or in physiological salt solution.

Prednisolone was dissolved in physiological saline.

**Treatment of drug schedule (Table 1):**

<table>
<thead>
<tr>
<th>Group</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>Prednisolone Standard (5 mg/kg) p. o.</td>
</tr>
<tr>
<td>II</td>
<td>Petroleum ether (100 mg/kg) p. o.</td>
</tr>
<tr>
<td>III</td>
<td>Ethanol (100 mg/kg) p. o.</td>
</tr>
<tr>
<td>IV</td>
<td>Negative Control (Distilled water containing 20% Tween 80) p. o.</td>
</tr>
</tbody>
</table>

**Acetic acid-induced ulcer model**

Study comprises of four different groups (n=6) as summarized in treatment schedule. Test animals in group II to III receives seven day treatment of the different crude extract as mentioned in treatment schedule. On eighth day all animals receives 0.1 ml 6% acetic acid intrarectally. Prednisolone treatment in standard group was started on the day of acetic acid treatment. Drug treatment in all groups was continued up to 10th day. After 48 hrs. of colitis induction mice were sacrificed by cervical dislocation and dissected upon to remove colon. 5cm long piece of colon was flushed gently with saline, cut upon and scored for inflammation based on the macroscopic features. Tissues were fixed in 10% formalin saline and examined histopathologically. Biochemical evaluation of colon inflammation was done using assay of MPO activity and MDA activity.

Overnight fasted mice, anaesthetized by Pentobarbital sodium (55.00 mg/Kg i.p.) 0.1 ml of 6% acetic acid once intrarectally. Allow to hang in air by holding tails for 1 – 2 min.

**Assessment of colitis severity**

The colonic samples from mice with colitis were shown to have severe mucous damage with edema, deep ulceration and hemorrhages. Oral administration of the petroleum ether and ethanol extracts of tubers of *Pueraria tuberosa*, two days before infusion of acetic acid into the colon was found to prevent progression of colitis. In mice treated with extracts, colonic macroscopic scores and the total square of damage were significantly reduced compared with those in the vehicle treated colitis group.

**Histopathological observation:**

In the present study control group showed higher degree of pathological changes i.e. more damage noticed in this
group. Methanol extract group showed considerable good reaction as compared to the other treated groups. Results shown Table 2.

**Table 2: Histopathological observation**

<table>
<thead>
<tr>
<th>Treatment Group</th>
<th>Ulceration</th>
<th>Hyperemia</th>
<th>Necrosis</th>
<th>Edema</th>
<th>Cellular infiltration</th>
<th>Goblet cell hyperplasia</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard Prednisolone (5 mg/kg)</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++</td>
</tr>
<tr>
<td>Pet. Ether extract 100 mg/kg</td>
<td>+++</td>
<td>+++</td>
<td>++</td>
<td>++</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>Ethanol extract 100 mg/kg</td>
<td>+++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++</td>
<td>++</td>
</tr>
<tr>
<td>Control, Acetic acid (-ve)</td>
<td>++++</td>
<td>+++++</td>
<td>++++</td>
<td>++</td>
<td>+++</td>
<td>+++</td>
</tr>
</tbody>
</table>

+: damage/ active changes up to less than 25%; ++: damage/ active changes up to less than 50%; +++: damage/ active changes up to less 75%; ++++: damage/ active changes up to more than 75%

**Figure 2:** Histopathology results

1. Prednisolone treated group, Red arrow – Hemorrhages
3. Ethanol extract treated group, Red arrow – Hemorrhages, Blue arrow - Leucocytic infiltration
4. –Ve treated group, Red arrow - Hemorrhages & ulceration, Blue arrow - Leucocytic infiltration

**Table 3: Determination of Ulcer Index**

<table>
<thead>
<tr>
<th>Treatment Group</th>
<th>Ulcer Index</th>
<th>Percent Ulcer Protection</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prednisolone (5 mg/kg)</td>
<td>2.62</td>
<td>45.60</td>
</tr>
<tr>
<td>Pet. Ether extract (100 mg/kg)</td>
<td>3.58</td>
<td>20.61</td>
</tr>
<tr>
<td>Ethanol extract (100 mg/kg)</td>
<td>2.10</td>
<td>56.74</td>
</tr>
<tr>
<td>Acetic acid (negative)</td>
<td>3.96</td>
<td>00.00</td>
</tr>
</tbody>
</table>

**Figure 3:** Ulcer Index of various extracts
Table 4: Qualitative Myeloperoxidase activity of the colon tissues

<table>
<thead>
<tr>
<th>Treatment Group</th>
<th>Myeloperoxidase activity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard Prednisolone (5 mg/kg)</td>
<td>++</td>
</tr>
<tr>
<td>Pet. Ether extract 100 mg/kg</td>
<td>+++</td>
</tr>
<tr>
<td>Ethanol extract 100 mg/kg</td>
<td>++</td>
</tr>
<tr>
<td>Control Acetic acid (negative)</td>
<td>++++</td>
</tr>
</tbody>
</table>

+: damage/active changes up to less than 25%
++: damage/active changes up to less than 50%
+++ : damage/active changes up to less than 75%
++++: damage/active changes up to more than 75%

Table 5: Quantitative Myeloperoxidase activity of the colon tissues

<table>
<thead>
<tr>
<th>Treatment Group</th>
<th>Myeloperoxidase activity in Blood U/ ml</th>
<th>Myeloperoxidase activity in tissue U/ ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stand. Prednisolone (5 mg/kg)</td>
<td>242</td>
<td>322</td>
</tr>
<tr>
<td>Pet. Ether extract 100 mg/kg</td>
<td>324</td>
<td>360</td>
</tr>
<tr>
<td>Ethanol extract 100 mg/kg</td>
<td>260</td>
<td>332</td>
</tr>
<tr>
<td>Control Acetic acid (negative)</td>
<td>362</td>
<td>375</td>
</tr>
</tbody>
</table>

Observations: Significantly normal activity seen with ethanol extract than petroleum ether extract, showed lower the values of this parameter as compared to other groups.

RESULTS AND DISCUSSION

Histopathological observation showed ulceration, hyperemia, necrosis, edema, cellular infiltration and goblet cell hyperplasia in the colon of mice treated with acetic acid. Treatment with ethanol extract of Pueraria tuberosa showed least ulceration and hyperemia (Table 2, Figure 2). Result from ulcer index showed better protective effect by ethanol extract of Pueraria tuberosa (Table 3).

Acetic acid caused increase in MPO level in blood and tissue up to 362 U/ml and 375 U/mg, respectively. After treatment with ethanol extract of Pueraria tuberosa, the MPO level in blood and tissue was decreased significantly to 260 U/ml and 332 U/mg respectively. Significant dose dependent reduction was observed after treatment with individual extract.

As ethanol extracts of Pueraria tuberosa root reduced MPO levels significantly, it may have potential anti-inflammatory role in the treatment of colitis because MPO is involved in the inflammatory reaction in colitis. Cytokines are responsible for modulating intestinal inflammation and injury. Increased levels of TNF-α and PGE2 may cause epithelial cell necrosis, edema, and neutrophil infiltration, as proved by the histopathological study. Recently Stucchi et al. (2006) found that LITAF (lipopolysaccharide-induced TNF-α factor), which mediates TNF-α expression in human macrophages, is significantly elevated above controls in macrophages of ileal and colonic tissues from patients with either CD or UC. Elevated levels of PGE2, goes in harmony with Otani et al. (2006) who proved that the increased level of PGE2 is attributed to its enhanced synthesis rather than reduced catabolism, both of which are mediated by TNF-α. On the other hand extracts of M. oleifera roots decreased significantly the gross lesion scores, and may be production of TNF-α and PGE2. Inhibition of PGE2, on the other hand, may follow that of TNF-α, or may result from its ability to inhibit cyclooxygenase enzymes. Since the intestine is in a constant state of controlled inflammation, thus amplification of the inflammatory response activates infiltration of inflammatory cells that triggers pathological responses and symptoms of IBD. Our study showed that acetic acid raised the levels of colonic MPO, indicating infiltration of neutrophils and perturbation of the inflammatory system. This fact is documented in both animal models and patients with IBD. In IBD, oxidative stress plays a role in disease initiation and progression. Reactive oxygen species (ROS) attack the cellular macromolecules, thus disrupting epithelial cell integrity and hindering mucosal recovery, especially in case of impaired endogenous defense systems. In this work, acetic acid induced ROS formation is inhibited by ethanol extracts of Pueraria tuberosa root are proved as good antioxidant, thus, its ability to inhibit free radical generation, as was proven in this work by restoring the redox state of the colonic mucosa, offers another explanation of the anti-ulcerogenic activity of this plant.

Ethanol extracts of Pueraria tuberosa root ameliorated neutrophil infiltration as evidenced by suppression of colon MPO and improvement of histological features. This action lends pharmacological support to folkloric, ethno-medical uses of the plant in the management of inflammatory GIT disorders.

CONCLUSION

The present antiulcer activity of petroleum ether and ethanolic extracts of tuber of *pueraria tuberosa* roxb in albino rats showed significant effect.

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REFERENCES


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Conflict of Interest: None declared.

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