

## RESEARCH ARTICLE

# Antiulcer Activity of *Trianthema portulacastrum* Linn. in Gastric Ulcers Caused by Alcohol

Sumeet Dwivedi<sup>1\*</sup>, Paras Gupta<sup>2</sup>, Divyansh Kateja<sup>3</sup>, Smriti<sup>4</sup>

<sup>1</sup>Acropolis Institute of Pharmaceutical Education and Research, Indore, Madhya Pradesh, India.

<sup>2</sup>United Institute of Pharmacy, Prayagraj, Uttar Pradesh, India.

<sup>3</sup>Department of Pharmacy Practice, NIMS Institute of Pharmacy, NIMS University Rajasthan, Jaipur, Rajasthan, India.

<sup>4</sup>Department of Pharmacy Practice, JSS College of Pharmacy, Mysuru, Karnataka, India.

Received: 20<sup>th</sup> March, 2024; Revised: 18<sup>th</sup> July, 2024; Accepted: 06<sup>th</sup> August, 2024; Available Online: 31<sup>st</sup> August, 2024

## ABSTRACT

*Trianthema portulacastrum* Linn. is an ancient Indian medicinal plant widely used in the treatment of ulcers, wounds, inflammation and pain. The leaves, flowers, and roots are the most important parts that are used traditionally. In the present investigation, antiulcer screening of etOH and aqueous leaves extract was screened in ethanol-induced gastric ulcers in rats at the selected doses. The results thus obtained were compared with standard drugs and it was found that EATPL exhibited significant results when compared with AETPL.

**Keywords:** Extract, Ethanol, Ulcer, *Trianthema portulacastrum*, Leaves.

International Journal of Pharmaceutical Quality Assurance (2024); DOI: 10.25258/ijpqa.15.3.12

**How to cite this article:** Dwivedi S, Gupta P, Kateja D, Smriti. Antiulcer Activity of *Trianthema portulacastrum* Linn. in Gastric Ulcers Caused by Alcohol. International Journal of Pharmaceutical Quality Assurance. 2024;15(3):1171-1172.

**Source of support:** Nil.

**Conflict of interest:** None

## INTRODUCTION

The most common gastrointestinal ailment, ulcers, are characterized by breaks in the stomach, small intestine, and occasionally the lower esophagus' lining. The ulcer is primarily linked to symptoms including nausea, vomiting, bloody or dark feces, burning pain in the abdomen that radiates to the chest, loss of appetite, and unexplained weight loss.<sup>1,2</sup> Together, ulcers, which is also known as gastric ulcers and duodenal ulcers, are referred to as peptic ulcers PU. *Trianthema portulacastrum*, commonly known as black pigweed (E) and patharchata (H) belongs to the family Lamiaceae is a flowering medicinal plants. The plant contains various bioactive compounds such as trianthenol, leptormol, cinnamic acid, beta cyanin and ecdysterone. Traditionally, almost every part of the plants is used for the treatment of inflammation, fungal infection, pain, liver disorders etc.<sup>3</sup>

## METHODOLOGY

### Gathering and Verification of Leaves

Leaves of *T. portulacastrum* Linn. were gathered from the Indore area, MP and were verified by Prof Dr. S.N. Dwivedi, Botanist, Rewa, MP and voucher specimen no J/Bot/TPL-011 was assigned.

### Extract Preparation

Approximately 500 gm of coarsely dried powdered leaves of selected plants were loaded in soxhlet using ethanol or water as solvent to obtain the ethanolic and aqueous extract.<sup>4</sup>

### Preparation of Niosomes

#### Acute toxicity of extract

The dose of ethanolic and aqueous extract of the selected plant was determined using OECD-423.<sup>5</sup>

#### Antiulcer activity ethanol induced

Rats having 150 to 200 gm weight were separated in five groups of six animals in each. Group I is considered as control and receive normal saline, i.e., 5 mL/kg bw. Group II is standard and received ranitidine, i.e., 150 mg/bw. Groups III and IV are treated group received 200 mg/kg bw extract dose of EETPL and AETPL.<sup>6,7</sup> The animals were given the test dose and after that, they were slaughtered to take stomach and finally, ulcer score was determined.

#### Statistical analysis

The Dunnet comparison test was used after one-way ANOVA to determine the statistical significance. ANOVA was utilized to compare nonparametric ulcer scores, and then the nonparametric Dunn post-test was employed.

\*Author for Correspondence: sumeet\_dwivedi2002@yahoo.com

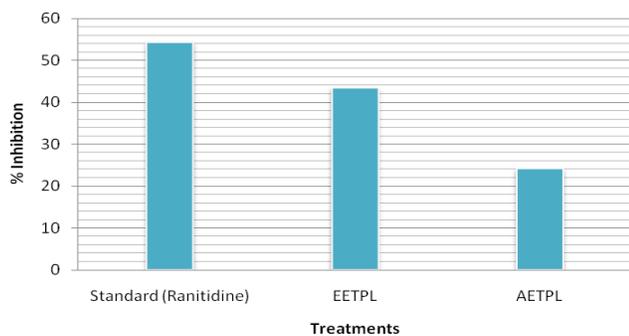
**Table 1:** Dose calculation of selected extract

Animals	Dose of extract in mg/kg	Death of animals in numbers	
		EETPL	AETPL
3	5	0	0
3	50	0	0
3	300	0	0
3	2000	0	0

**Table 2:** Gastric UI and %inhibition of the extract

Groups	Dose in mg/kg	Gastric UI	%Inhibition
Control	-	5.44 ± 0.22	-
Standard	150	2.10 ± 0.18	54.29**
EETPL	200	3.11 ± 0.20	43.44*
AETPL	200	4.22 ± 0.11	24.11**

Note: Reading are X[Mean] ± SEM, n = 6; \*  $p < 0.01$ , \*\* $p < 0.001$  is significant

**Figure 1:** Comparative %inhibition

## RESULTS AND DISCUSSION

The toxicity of EATPL and AETPL was determined using OECD-423 via acute oral toxicity experiments in order to ascertain the LD<sub>50</sub>. According to the findings, the extracts fit under group 5 (unclassified). Consequently, the ED<sub>50</sub> was 200 mg/kg and the LD<sub>50</sub> was 2000 mg/kg. Thus, 200 mg doses were taken in consideration for the study. Table 1 presents the detailed results.

In the ethanol-induced ulcer model, EETPL and HAESCS had a significantly lower ulcerogenic effect than the control group. The percentage of inhibition in the animal group treated with extract with 200 mg/kg dose was determined to be 54.29, 43.44, and 24.11%, respectively, when compared to the standard, EETPL, and AETPL. These findings suggest that EETPL produces better results than AETPL. Table 2 and Figure 1 show the results.

## CONCLUSION

From the results, it was clearly depicted that EETPL was more effective in reducing ulcers than AETPL when compared to the reference.

## ACKNOWLEDGMENT

The author (s) are thankful to Dr. S.N. Dwivedi, Retd. Prof. & HOD, APS University, Rewa, (M.P.) for his necessary supervision.

## REFERENCES

- Dwivedi S. Status survey of medicinal plants wealth of Malwa region of Madhya Pradesh with special reference to conservation of vulnerable and endangered species. *Journal of economic and taxonomic botany*. 2009; 33(2): 443-452.
- Loffeld RJLF, Liberov B, Dekkers PEP. Peptic ulcer disease: A vanishing disease. *Journal of Gastric Disorder and Therapy*. 2016;2:4. <http://dx.doi.org/10.16966/2381-8689.123>
- Kumar A, Ashwlayan VD, Verma M. Diagnostic approach and pharmacological treatment regimen of peptic ulcer disease. *Pharmacy and Pharmaceutical Research* 2019;1(1):1-12. DOI: 10.15406/ghoa.2019.10.00352
- Kokate CK. *Practical Pharmacognosy*, Vallabh Prakashan, Delhi., 4<sup>th</sup> Edition, 107 – 111, 1997.
- OECD. Guidelines for the testing of chemicals revised draft guideline 423: Acute oral toxicity. France: Organization for Economic Cooperation and Development; 2000.
- Bhajoni PS, Meshram GG, Lahkar M. Evaluation of Antiulcer Activity of the Leaves of *azadirachta indica*: An Experimental Study. *International Journal of Integrative Medicine*. 2016;3:10-16. <https://doi.org/10.1159/000442750>
- Prabhu K, Rahan S. Assessment of Antiulcer Activity of Ethanolic Extract of *Magnifera indica* seed Kernel using Acid Ethanol Induced Ulcer Model. *International Journal of Current Microbiology and Applied Science*. 2015;4(4): 854-860.