



## FORMULATION AND EVALUATION OF ANTIULCER GEL FROM GUAVA LEAVES AND CORDIA DICHOTOMA SEEDS

**Prof. Shital K. Datir<sup>1\*</sup>, Reshma R. Ugale<sup>1</sup>, Raksha B. Madhe<sup>2</sup>, Kiran S. Wagh<sup>3</sup>**

NGSPM'S College of Pharmacy Anjaneri, Trimbakeshwar Nashik, Maharashtra, India.



**\*Corresponding Prof. Shital K. Datir**

Department of Pharmaceutics, NGSPM'S College of Pharmacy Anjaneri, Trimbakeshwar Nashik, Maharashtra, India.

**DOI:** <https://doi.org/10.5281/zenodo.20642203>

**How to cite this Article:** Prof. Shital K. Datir<sup>1\*</sup>, Reshma R. Ugale<sup>1</sup>, Raksha B. Madhe<sup>2</sup>, Kiran S. Wagh<sup>3</sup>. (2026). Formulation and Evaluation of Antiulcer Gel from Guava Leaves and Cordia dichotoma Seeds. European Journal of Biomedical and Pharmaceutical Sciences, 13(6), 493-495.

This work is licensed under Creative Commons Attribution 4.0 International license.



Article Received on 15/05/2026

Article Revised on 05/06/2026

Article Published on 10/06/2026

### ABSTRACT

Peptic ulcer disease is a common gastrointestinal disorder characterized by erosion of the gastric mucosal lining. Herbal medicines have gained considerable attention due to their safety, efficacy, and reduced side effects. The present study aimed to formulate and evaluate an antiulcer gel containing extracts of Guava leaves (*Psidium guajava*) and *Cordia dichotoma* seeds. Guava leaves possess antioxidant, anti-inflammatory, and gastroprotective activities, while *Cordia dichotoma* seeds contain mucilage and bioactive constituents that aid in mucosal protection and wound healing. The herbal gel was prepared using Carbopol 940 as the gelling agent and evaluated for physical appearance, pH, viscosity, spreadability, extrudability, and stability. The formulated gel showed satisfactory physicochemical characteristics and demonstrated potential antiulcer activity due to the synergistic action of both herbal extracts. The study suggests that the developed formulation may serve as a promising herbal alternative for the management of gastric ulcers.

**KEYWORDS:** Antiulcer gel, *Psidium guajava*, *Cordia dichotoma*, Herbal formulation, Carbopol 940, Gastroprotective activity.

### INTRODUCTION

Peptic ulcer disease is a chronic disorder affecting the stomach and duodenum and is commonly associated with *Helicobacter pylori* infection, excessive acid secretion, stress, alcohol consumption, and prolonged use of non-steroidal anti-inflammatory drugs (NSAIDs).<sup>[1]</sup> Conventional antiulcer therapies, including proton pump inhibitors and H<sub>2</sub> receptor antagonists, are effective but may cause adverse effects upon long-term use.<sup>[2]</sup> Medicinal plants have been extensively investigated for their gastroprotective potential due to the presence of flavonoids, tannins, alkaloids, and phenolic compounds.<sup>[3]</sup> *Psidium guajava* (Guava) leaves contain quercetin, tannins, and essential oils that exhibit antioxidant and anti-inflammatory activities contributing to ulcer healing.<sup>[4]</sup> *Cordia dichotoma* is a medicinal plant whose seeds are rich in mucilage, polysaccharides, and bioactive compounds that help protect the gastric mucosa and promote tissue regeneration.<sup>[5]</sup> Combining these herbal ingredients in a topical oral gel system may provide prolonged mucosal contact and enhanced therapeutic benefits.

Therefore, the present work was undertaken to formulate and evaluate an antiulcer gel containing Guava leaf extract and *Cordia dichotoma* seed extract for potential gastroprotective activity.

### Antiulcer Gel

Antiulcer gels are semisolid formulations designed to provide localized protection and therapeutic action on ulcerated mucosal tissues. They form a protective layer over the affected area, reduce irritation, and enhance healing by maintaining prolonged contact with bioactive compounds.<sup>[6]</sup> Herbal antiulcer gels have gained popularity due to their natural origin and minimal side effects.

### Advantages of Antiulcer Gel

1. Provides prolonged contact with ulcerated tissues.
2. Improves patient compliance.
3. Easy application and administration.
4. Reduced systemic side effects.
5. Enhanced bioavailability of herbal constituents.
6. Cost-effective and eco-friendly formulation.<sup>[7]</sup>

**Disadvantages of Antiulcer Gel**

1. Limited drug loading capacity.
2. Possibility of microbial contamination.
3. Stability may be affected by environmental conditions.
4. Requires suitable preservatives.
5. Shorter shelf life compared to some conventional dosage forms.<sup>[7]</sup>

**Formulation of Antiulcer Gel****Table 1: Composition of Antiulcer Gel (20 g).**

| Sr.No | Ingredients                   | Quantity   |            |            |
|-------|-------------------------------|------------|------------|------------|
|       |                               | F1 (gm)    | F2 (gm)    | F3 (gm)    |
| 1.    | Guava Leaf Extract            | 1g         | 1g         | 1g         |
| 2.    | Cordia dichotoma Seed Extract | 1g         | 1g         | 1g         |
| 3.    | Carbopol 940                  | 0.4g       | 0.4g       | 0.4g       |
| 4.    | Glycerin                      | 2ml        | 2ml        | 2ml        |
| 5.    | Propyl Paraben                | 0.02g      | 0.02g      | 0.02g      |
| 6.    | Triethanolamine               | q.s        | q.s        | q.s        |
| 7.    | Purified water                | up to 20 g | up to 20 g | up to 20 g |

**Method of Preparation**

1. Carbopol 940 was dispersed slowly in purified water and allowed to hydrate for 24 hours.<sup>[8]</sup>
2. Guava leaf extract and Cordia dichotoma seed extract were dissolved in a small quantity of water.
3. Methyl paraben and propyl paraben were dissolved in propylene glycol.
4. The preservative solution was added to the hydrated Carbopol gel with continuous stirring.
5. Herbal extracts were incorporated gradually into the gel base.
6. Triethanolamine was added dropwise to adjust the pH and obtain the desired gel consistency.
7. The final gel was mixed thoroughly to ensure uniform distribution of ingredients and stored in airtight containers.<sup>[8]</sup>

**Evaluation of Antiulcer Gel****1. Physical Appearance**

The gel was visually examined for color, homogeneity, consistency, and presence of any particulate matter.<sup>[9]</sup>

**2. pH Determination**

The pH of the formulation was measured using a calibrated digital pH meter at room temperature<sup>[9]</sup>

**3. Viscosity**

Viscosity was determined using a Brookfield viscometer at appropriate spindle speed and temperature<sup>[9]</sup>

**4. Spreadability**

Spreadability was determined by placing the gel between two glass slides and measuring the time required for separation under specified weight<sup>[10]</sup>

**5. Extrudability**

Extrudability was evaluated by determining the force required to extrude the gel from a collapsible tube.<sup>[10]</sup>

**6. Homogeneity**

The formulation was examined visually after application on a glass slide to ensure uniformity<sup>[10]</sup>

**4. Transparency**

The 1 gm gel formulation taken in 10 ml test tube and visually checked for its transparency.

**7. Stability Study**

The prepared gel was stored at room temperature and accelerated conditions for three months and evaluated periodically for changes in appearance, pH, and viscosity.<sup>[10]</sup>

**Fig. 1: Spreadability.****Fig. 2: Irritancy.****RESULT AND DISCUSSION**

Evaluation Parameter of Herbal gel was performed.

## 1. Physical Appearance

| Sr. no. | Characters | Observation       |
|---------|------------|-------------------|
| 1.      | Nature     | Gel               |
| 2.      | Colour     | Greenish- Brown   |
| 3.      | Appearance | Semi- transparent |

## 2. Determination of pH.

| Sr. no. | Formulation batches |              |       |
|---------|---------------------|--------------|-------|
|         | F1                  | F2           | F3    |
| 1.      | Yellowish orange    | Yellow green | Green |

## 3. Determination Irritancy Test.

| Formulation | Irritancy      |
|-------------|----------------|
| F1          | Irritancy      |
| F2          | Mild irritancy |
| F3          | No irritancy   |

## 4. Determination of Stability.

| Sr.No. | Parameter | Formulation batches |        |        |
|--------|-----------|---------------------|--------|--------|
|        |           | F1                  | F2     | F3     |
| 1.     | Stability | Passed              | Failed | Passed |

According to above table stability of F1 and F3 passed and F2 was failed the test .

## 5. Determination of Transparency

| Sr. No. | Parameter    | Formulation batches |                      |                   |
|---------|--------------|---------------------|----------------------|-------------------|
|         |              | F1                  | F2                   | F3                |
| 1       | Transparency | Opaque              | Slightly Transparent | Semi- Transparent |

## 6. Determination of Spreadability.

| Sr. No. | Parameter     | Formulation batches |                       |                   |
|---------|---------------|---------------------|-----------------------|-------------------|
|         |               | F1                  | F2                    | F3                |
| 1.      | Spreadability | Slightly spreadable | Moderately spreadable | Easily spreadable |

## CONCLUSION

The present study was successfully carried out to formulate and evaluate an herbal anti-ulcer gel using Guava leaves extract and Cordia dichotoma seed extract. The prepared formulations were evaluated for different parameters such as pH, viscosity, homogeneity, spreadability, and transparency. The results showed that the gel possessed satisfactory physicochemical properties and good stability. Among the formulations, the optimized batch exhibited better consistency and spreadability suitable for oral application. Hence, the formulated herbal gel can be considered a safe, effective, and economical treatment option for mouth ulcers with minimal side effects.

## REFERENCES

- Rang HP, Dale MM, Ritter JM, Flower RJ. *Rang and Dale's Pharmacology*. 9th ed. Elsevier, 2019.
- Brunton LL, Hilal-Dandan R, Knollmann BC. *Goodman & Gilman's The Pharmacological Basis of Therapeutics*. 13th ed. McGraw-Hill Education, 2018.
- Kokate CK, Purohit AP, Gokhale SB. *Pharmacognosy*. 56th ed. Nirali Prakashan, 2021.
- Gutierrez RM, Mitchell S, Solis RV. Psidium guajava: A review of its traditional uses, phytochemistry and pharmacology. *J Ethnopharmacol.*, 2008; 117(1): 1-27.
- Kirtikar KR, Basu BD. *Indian Medicinal Plants*. Vol. 3. International Book Distributors, 2006.
- Aulton ME, Taylor K. *Aulton's Pharmaceutics: The Design and Manufacture of Medicines*. 6th ed. Elsevier, 2022.
- Lachman L, Lieberman HA, Kanig JL. *The Theory and Practice of Industrial Pharmacy*. 4th ed. CBS Publishers, 2013.
- Allen LV, Popovich NG, Ansel HC. *Ansel's Pharmaceutical Dosage Forms and Drug Delivery Systems*. 11th ed. Wolters Kluwer, 2020.
- Indian Pharmacopoeia Commission. *Indian Pharmacopoeia*. Ghaziabad: IPC., 2022.
- Sinko PJ. *Martin's Physical Pharmacy and Pharmaceutical Sciences*. 7th ed. Lippincott Williams & Wilkins, 2017.