**OXALIS 10- oxalis 10 gel**
**Uriel Pharmacy Inc.**

*Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.*

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**Oxalis 10**

Directions: FOR TOPICAL USE ONLY.

Apply to skin as needed. Under age 2: Consult a doctor.

Active Ingredient: 100 gm contains: 10 gm Oxalis (Wood sorrel) 1X

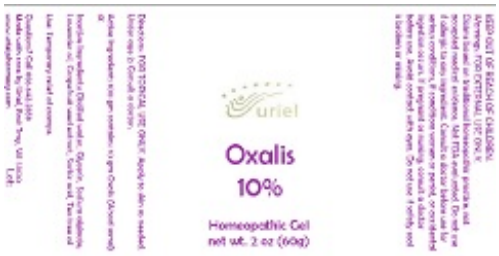
Inactive Ingredients: Distilled water, Glycerin, Sodium alginate, Lavender oil, Grapefruit seed extract, Sorbic acid, Tea tree oil

Use: Temporary relief of cramps.

KEEP OUT OF REACH OF CHILDREN.

Warnings: FOR EXTERNAL USE ONLY.
Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions, if conditions worsen or persist, or accidental ingestion occurs. If pregnant or nursing, consult a doctor before use. Avoid contact with eyes. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 www.urielpharmacy.com




## OXALIS 10
oxalis 10 gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:48951-7102 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **OXALIS ACETOSELLA LEAF** (UNII: U1W3U02EW0) (OXALIS ACETOSELLA LEAF - UNII:U1W3U02EW0) | OXALIS ACETOSELLA LEAF | 1 [hp_X] in 1 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |

| | |
|---|---|
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **SODIUM ALGINATE** (UNII: C269C4G2ZQ) | |
| **LAVENDER OIL** (UNII: ZBP1YXW0H8) | |
| **SORBIC ACID** (UNII: X045WJ989B) | |
| **CITRUS PARADISI SEED** (UNII: 12F08874Y7) | |
| **TEA TREE OIL** (UNII: VIF565UC2G) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:48951-7102-5 | 60 g in 1 TUBE; Type 0: Not a Combination Product | 09/01/2009 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved homeopathic | | 09/01/2009 | |

## Labeler - Uriel Pharmacy Inc. (043471163)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Uriel Pharmacy Inc. | | 043471163 | manufacture(48951-7102) |

Revised: 9/2009 Uriel Pharmacy Inc.