## OXALIS 1X- oxalis 1x ointment 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 1X

Directions: FOR TOPICAL USE ONLY.

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

Active Ingredients: Oxalis extract (Wood sorrel) 1X

Inactive Ingredients: Sesame oil, Lanolin, Yellow beeswax, Lanolin alcohol

Prepared using rhythmical processes.

Uses: Temporary relief of cramps.

KEEP OUT OF REACH OF CHILDREN.

Warnings: 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 Uriel, East Troy, WI 53120 shopuriel.com Lot:



OXALIS 1X oxalis 1x ointment								
Product Information								
Product Type	HUMAN OTC DRUG	ltem Code (Source)	NDC:48951-7214					
Route of Administration	TOPICAL							
Active Ingredient/Active Moiety								

		Ingredient Name		Stren		Strength			
	OXALIS STRICTA WHOLE (UNII: 3STW243384) (OXALIS STRICTA WHOLE - OXALIS S UNII:3STW243384) WHOLE				ICTA	1 [hp_X] in 1 g			
Inactive Ingredients									
		Ingredient Name			Strength				
YE	LLOW WAX (UNI	I: 2ZA36H0S2V)							
LANOLIN ALCOHOL (UNII: 884C3FA9HE)									
SE	SAME OIL (UNII:	QX10HYY4QV)							
LANOLIN (UNII: 7EV65EAW6H)									
	ickaging Item Code	Package Description	Marketin Dat			eting End Date			
#		Package Description 30 g in 1 TUBE; Type 0: Not a Combination Product				-			
#	Item Code NDC:48951- 7214-2	30 g in 1 TUBE; Type 0: Not a Combination Product	Dat			-			
#	Item Code NDC:48951- 7214-2	30 g in 1 TUBE; Type 0: Not a Combination Product Information Application Number or Monograp	Dat 09/01/2009			Date			
# 1 M	Item Code NDC:48951- 7214-2 arketing Marketing Category	30 g in 1 TUBE; Type 0: Not a Combination Product	Dat 09/01/2009 h Market	e		Date			
# 1 M	Item Code NDC:48951- 7214-2 arketing Marketing	30 g in 1 TUBE; Type 0: Not a Combination Product Information Application Number or Monograp	Dat 09/01/2009 h Market	ing Start tate		Date keting End			

## Labeler - Uriel Pharmacy, Inc (043471163)

Establishment								
Name	Address	ID/FEI	<b>Business Operations</b>					
Uriel Pharmacy, Inc		043471163	manufacture(48951-7214)					

Revised: 1/2025

Uriel Pharmacy, Inc