OXALIS BELLADONNA- oxalis belladonna liquid
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.

Oxalis Belladonna

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Atropa belladonna (Nightshade) 4X, Chamomilla (Chamomile) 4X, Gelsemium (Yellow jasmine) 4X, Oxalis (Wood sorrel) 4X, Sanguinaria (Bloodwort) 4X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

Use: 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 or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. 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 BELLADONNA

oxalis belladonna liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7108
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	4 [hp_X] in 1 mL

MATRICARIA RECUTITA (UNII: G0R4UBI2ZZ) (MATRICARIA RECUTITA - UNII:G0R4UBI2ZZ)	MATRICARIA RECUTITA	4 [hp_X] in 1 mL
GELSEMIUM SEMPERVIRENS ROOT (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	4 [hp_X] in 1 mL
OXALIS ACETOSELLA LEAF (UNII: U1W3U02EW0) (OXALIS ACETOSELLA LEAF - UNII:U1W3U02EW0)	OXALIS ACETOSELLA LEAF	4 [hp_X] in 1 mL
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	4 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-7108-3	60 mL in 1 BOTTLE, DROPPER; 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(48951-7108)

Revised: 4/2018 Uriel Pharmacy Inc.