OXALIS BELLADONNA SPECIAL ORDER- oxalis belladonna special order 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 Special Order

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

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

Inactive Ingredients: Water, Salt

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.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 www.urielpharmacy.com

OXALIS BELLADONNA SPECIAL ORDER

oxalis belladonna special order liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7107
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
CHELIDONIUM MAJUS ROOT (UNII: FLT36UCF0N) (CHELIDONIUM MAJUS ROOT - UNII:FLT36UCF0N)	CHELIDONIUM MAJUS ROOT	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)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-7107-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

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(48951-7107)

Revised: 8/2017 Uriel Pharmacy Inc.