COLCHICUM CHELIDONIUM- colchicum chelidonium 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.

Colchicum Chelidonium

Directions: FOR TOPICAL USE ONLY.

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

Active Ingredient: Colchicum e tub. Inc.(Meadow saffron) 2X, Spongia tosta (Roasted sea sponge) 2X, Chelidonium e flos. Inc. (Greater celandine) 3X

Inactive Ingredients: Lanolin, Sunflower seed oil, Water, Yellow beeswax, Lanolin alcohol

Prepared using rhythmical processes.

Use: Temporary relief of headache.

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
Made with care by Uriel, East Troy, WI 53120
shopuriel.com Lot:

COLCHICUM CHELIDONIUM

colchicum chelidonium ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-3271
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHELIDONIUM MAJUS FLOWERING TOP (UNII: O9HP59XU7F) (CHELIDONIUM MAJUS FLOWERING TOP - UNII:O9HP59XU7F)	CHELIDONIUM MAJUS FLOWERING TOP	3 [hp_X] in 1 g
COLCHICUM AUTUMNALE WHOLE (UNII: W79255C628) (COLCHICUM AUTUMNALE WHOLE - UNII:W79255C628)	COLCHICUM AUTUMNALE WHOLE	2 [hp_X] in 1 g
SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA OFFICINALIS SKELETON, ROASTED - UNII:1PIP394IID)	SPONGIA OFFICINALIS SKELETON, ROASTED	2 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
SUNFLOWER OIL (UNII: 3W1JG795YI)	
LANOLIN ALCOHOL (UNII: 884C3FA9HE)	
BEESWAX (UNII: 2ZA36H0S2V)	
LANOLIN (UNII: 7EV65EAW6H)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3271-5	30 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(48951-3271)

Revised: 12/2025 Uriel Pharmacy, Inc.