ARNICA CEPA SYMPHYTUM- arnica cepa symphytum 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.

Arnica Cepa Symphytum

Directions: FOR TOPICAL USE ONLY.

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

Active Ingredients: 100gm contains: 2gm Allium cepa (Onion) 1X, 5gm Arnica montana 1X, 3gm Symphytum ex herba (Comfrey) 1X

Inactive Ingredients: Organic olive oil, Lanolin, Sunflower seed oil, Spring water, Yellow beeswax, Lanolin alcohol

Prepared using rhythmical processes.

Uses: Temporary relief of bruises, sprains, and minor cuts.

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:

ARNICA CEPA SYMPHYTUM

arnica cepa symphytum ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1390
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
COMFREY ROOT (UNII: M9VVZ08EKQ) (COMFREY ROOT - UNII:M9VVZ08EKQ)	COMFREY ROOT	1 [hp_X] in 1 g
ONION (UNII: 492225Q21H) (ONION - UNII:492225Q21H)	ONION	1 [hp_X] in 1 g
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	1 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
BEESWAX (UNII: 2ZA36H0S2V)	
LANOLIN ALCOHOL (UNII: 884C3FA9HE)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
LANOLIN (UNII: 7EV65EAW6H)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1390-2	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(48951-1390)

Revised: 1/2025 Uriel Pharmacy, Inc