

ARNICA 20%- arnica 20% gel **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 20%

Directions: FOR TOPICAL USE ONLY.

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

Active Ingredient: 100 gm contains: 20 gm Arnica 1X

Inactive Ingredients: Spring water, Organic cane alcohol, Glycerin, Boric acid, Sodium alginate, Lavender oil, Sorbic acid, Tea tree oil, Sodium hydroxide, Grapefruit seed extract

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

prepared using rhythmical processes

KEEP OUT OF REACH OF CHILDREN.

Warnings: FOR EXTERNAL USE ONLY

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:

Uriel

Arnica 20%

Homeopathic Gel
net wt. 2 oz (60g)

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ARNICA 20%

arnica 20% gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1374
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	1 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
CITRUS PARADISI SEED (UNII: 12F08874Y7)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
TEA TREE OIL (UNII: VIF565UC2G)	
SORBIC ACID (UNII: X045WJ989B)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BORIC ACID (UNII: R57ZHV85D4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1374-5	60 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(48951-1374)