

ARNICA AURUM 20/30- arnica aurum 20/30 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.

Arnica Aurum 20/30

Directions: FOR ORAL USE.

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

Active Ingredients: Onopordon (Cotton thistle) 3X, Primula (Cowslip) 3X, Hyoscyamus (Henbane) 4X, Arnica 20X, Aurum (Metallic gold) 30X

Inactive Ingredients: Water, Salt

"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 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 shopuriel.com

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Lot:

Arnica Aurum 20X/30X

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Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

ARNICA AURUM 20/30

arnica aurum 20/30 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1106
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ONOPORDUM ACANTHIUM FLOWER (UNII: AP97AUF88E) (ONOPORDUM ACANTHIUM FLOWER - UNII:AP97AUF88E)	ONOPORDUM ACANTHIUM FLOWER	3 [hp_X] in 1 mL
PRIMULA VERIS FLOWER (UNII: W5BET37294) (PRIMULA VERIS FLOWER - UNII:W5BET37294)	PRIMULA VERIS FLOWER	3 [hp_X] in 1 mL
HYOSCYAMUS NIGER LEAF (UNII: 32IT7G8BAW) (HYOSCYAMUS NIGER LEAF - UNII:32IT7G8BAW)	HYOSCYAMUS NIGER LEAF	4 [hp_X] in 1 mL
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	20 [hp_X] in 1 mL
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	30 [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:48951-1106-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 0: Not a Combination Product		

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(48951-1106)

Revised: 1/2024

Uriel Pharmacy Inc.