

ARNICA E RAD. 20X- arnica e rad. 20x 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 e rad. 20X

Directions: FOR ORAL USE.

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

Active Ingredient: Arnica e rad. (Arnica) 20X

Inactive Ingredients: Water, Salt

Use: Temporary relief of sore muscles.

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

Directions: FOR ORAL USE.
 Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredient: Arnica e rad. (Arnica) 20X

Inactive Ingredients: Water, Salt

Use: Temporary relief of sore muscles.

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 Lot:



**Arnica
 e rad. 20X**

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

Arnica e rad. 20X

ARNICA E RAD. 20X

arnica e rad. 20x liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1336
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA ROOT (UNII: MUE8 Y11327) (ARNICA MONTANA ROOT - UNII:MUE8 Y11327)	ARNICA MONTANA ROOT	20 [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:48951-1336-1	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(48951-1336)

Revised: 11/2019

Uriel Pharmacy Inc.