RHEUMA COMP. 1- rheuma comp. 1 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.

Rheuma comp. 1

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Arnica montana 2X, Bryonia e rad. (White bryony) 3X, Aconitum e

tub. (Monkshood) 4X

Inactive Ingredients: Distilled water, 30% Organic cane 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 or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. 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:

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Homeopathic Liquid net vol. 2 fl. oz (60ml) Questions? Call 866.642.2858 Made with care by Uriel, East Tray, WI 53to shapuriel.com. Lot:

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:48951-8420

Route	of Ac	lmini	istra	tion

ORAL

Active Ingredient/Active Moiety							
Ingredient Name	Basis of Strength	Strength					
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII: OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	2 [hp_X] in 1 mL					
ACONITUM NAPELLUS (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	4 [hp_X] in 1 mL					
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	3 [hp_X] in 1 mL					

Inactive Ingredients					
Ingredient Name	Strength				
ALCOHOL (UNII: 3K9958V90M)					
WATER (UNII: 059QF0KO0R)					

l	Packaging								
	# Item Code Package Description		Marketing Start Date	Marketing End Date					
	1	NDC:48951- 8420-3	60 mL in 1 BOTTLE, DROPPER; 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(48951-8420)					

Revised: 11/2024 Uriel Pharmacy, Inc.