MANDRAGORA ARNICA- mandragora arnica 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.

Mandragora Arnica

Directions: FOR ORAL USE.

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

Active Ingredients: Betula (Silver birch leaves) 3X, Mandragora (Mandrake) 3X, Meniscus genus (Bovine medial meniscus of the knee joint) 8X, Formica (Red wood ant) 10X, Arnica 15X, Equisetum (Common horsetail) 15X

Inactive Ingredients: Water, Salt

Use: Temporary relief of sore joints.

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.shopuriel.com Lot:

MANDRAGORA ARNICA mandragora arnica liquid			
Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7200

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA WHOLE (UNII: O80TY208ZW) (ARNICA MONTANA WHOLE - UNII:O80TY208ZW)	ARNICA MONTANA WHOLE	15 [hp_X] in 1 mL
BETULA PUBESCENS LEAF (UNII: 84SOH0O3OO) (BETULA PUBESCENS LEAF - UNII:84SOH0O3OO)	BETULA PUBESCENS LEAF	3 [hp_X] in 1 mL
MANDRAGORA OFFICINARUM ROOT (UNII: I2XCB174VB) (MANDRAGORA OFFICINARUM ROOT - UNII:I2XCB174VB)	MANDRAGORA OFFICINARUM ROOT	3 [hp_X] in 1 mL
EQUISETUM ARVENSE TOP (UNII: 1DP6Y6B65Z) (EQUISETUM ARVENSE TOP - UNII:1DP6Y6B65Z)	EQUISETUM ARVENSE TOP	15 [hp_X] in 1 mL
FORMICA RUFA (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)	FORMICA RUFA	10 [hp_X] in 1 mL
SUS SCROFA MENISCUS (UNII: 1C8T78BD9Q) (SUS SCROFA MENISCUS - UNII:1C8T78BD9Q)	SUS SCROFA MENISCUS	8 [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:48951-7200-1	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(48951-7200)

Revised: 5/2022 Uriel Pharmacy, Inc.