

FORMICA APIS- formica apis 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.

Formica Apis

Use: Temporary relief of sore joints.

FOR ORAL USE ONLY

Directions: Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow. Follow inside instructions to open ampule.

Warnings:

NOT FOR INJECTION

Do not use if allergic to any ingredient. Consult a physician if symptoms worsen. If pregnant or nursing, consult a physician before use.

Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

KEEP OUT OF REACH OF CHILDREN.

Active Ingredients: Apis (Honeybee) 3X, Formica (Red wood ant) 3X

Inactive Ingredients: Water, Salt

Prepared using rhythmical processes.

Questions? Call 866.642.2858

Uriel, East Troy, WI 53120

shopuriel.com Lot:

Use: Temporary relief of headache.

Directions: Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow. Follow inside instructions to open ampule.

Warnings:
NOT FOR INJECTION
Do not use if allergic to any ingredient. Consult a physician if symptoms worsen. If pregnant or nursing, consult a physician before use.
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Uriel, East Troy, WI 53120
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Lot:



Formica Apis

Homeopathic Ampules
FOR ORAL USE ONLY
net vol. 0.3 fl. oz (10 x 1 ml)

Formica Apis

FORMICA APIS

formica apis liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-4193
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)		APIS MELLIFERA	3 [hp_X] in 1 mL
FORMICA RUFA (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)		FORMICA RUFA	3 [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:48951-4193-1	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(48951-4193)