AGARICUS PHOSPHORUS- agaricus phosphorus 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.

Agaricus Phosphorus

Directions: FOR ORAL USE.

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

Active Ingredients: Filix mas (Male fern) 3X, Argentum fluoratum (Silver fluoride) 6X, Conchae (Oyster shells) 6X, Phosphorus (Yellow phosphorus) 6X, Agaricus (Fly agaric mushroom) 8X

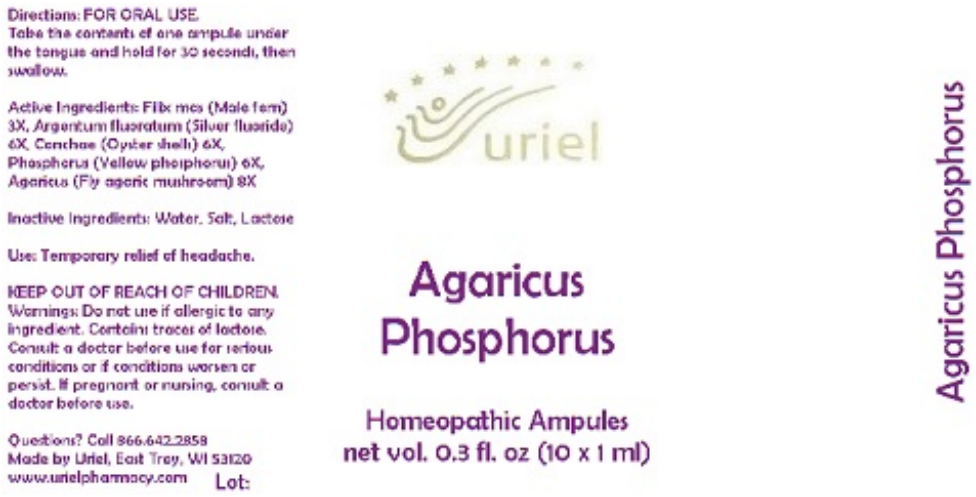
Inactive Ingredients: Water, Salt

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. Contains traces of lactose. 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 Made by Uriel, East Troy, WI 53120 www.urielpharmacy.com



AGARICUS PHOSPHORUS

agaricus phosphorus liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1022
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DRYOPTERIS FILIX-MAS ROOT (UNII: C0ZK0RRF5X) (DRYOPTERIS FILIX-MAS ROOT - UNII:C0ZK0RRF5X)	DRYOPTERIS FILIX-MAS ROOT	3 [hp_X] in 1 mL

SILVER FLUORIDE (UNII: 1Z00ZK3E66) (SILVER FLUORIDE - UNII:1Z00ZK3E66)	SILVER FLUORIDE	6 [hp_X] in 1 mL
OSTREA EDULIS SHELL (UNII: 49OY13BE7Z) (OSTREA EDULIS SHELL - UNII:49OY13BE7Z)	OSTREA EDULIS SHELL	6 [hp_X] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	6 [hp_X] in 1 mL
AMANITA MUSCARIA FRUITING BODY (UNII: DIF093I037) (AMANITA MUSCARIA FRUITING BODY - UNII:DIF093I037)	AMANITA MUSCARIA FRUITING BODY	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:48951-1022-1	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(48951-1022)

Revised: 4/2018 Uriel Pharmacy Inc.