

**PHYTOLACCA E RAD 6X- phytolacca e rad 6x 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.*

**Phytolacca e rad 6X**

Directions: FOR ORAL USE.

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

Active Ingredient: Phytolacca e rad. 6X

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. 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 shopuriel.com

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

Active Ingredient: Phytolacca (Pokeweed)  
6X

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. Consult  
a doctor before use for serious conditions or  
if conditions worsen or persist. If pregnant or  
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Lot:



**Phytolacca  
e rad. 6X**

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

**Phytolacca e rad. 6X**

**PHYTOLACCA E RAD 6X**

phytolacca e rad 6x liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-8402
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name		Basis of Strength	Strength	
PHYTOLACCA AMERICANA ROOT (UNII: 11E6VI8VEG) (PHYTOLACCA AMERICANA ROOT - UNII:11E6VI8VEG)		PHYTOLACCA AMERICANA ROOT	6 [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:48951-8402-1	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(48951-8402)

Revised: 11/2023

Uriel Pharmacy Inc.