

PLUMBUM 8X- plumbum 8x 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.

Plumbum 8X

Directions: FOR ORAL USE.

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

Active Ingredient: Plumbum (Lead) 8X

Inactive Ingredients: Water, Salt, Lactose

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. Contains traces of lactose. 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

www.urielpharmacy.com

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www.urielpharmacy.com Lot:



**Plumbum
8X**

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

Plumbum 8X

PLUMBUM 8X

plumbum 8x liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8362
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEAD (UNII: 2P299V784P) (LEAD - UNII:2P299V784P)	LEAD	8 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8362-1	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(48951-8362)

Revised: 12/2019

Uriel Pharmacy Inc.