

**PULPA DENTIS 30 SPECIAL ORDER- pulpa dentis 30 special order 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.*

**Pulpa Dentis 30 Special Order**

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops.  
Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredient: Pulpa dentis (Bovine dental pulp) 30X

Inactive Ingredient: Distilled water

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. Do not use if safety seal is broken or missing.

REFRIGERATE AFTER OPENING.

BEST WHEN USED WITHIN 90 DAYS OF OPENING.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120

[www.urielpharmacy.com](http://www.urielpharmacy.com)

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**Pulpa dentis  
30X  
Special Order**  
**Homeopathic Liquid**  
**net vol. 2 fl. oz (60ml)**

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Take 3-4 times daily. Ages 12 and older: 10 drops.  
Ages 2-11: 5 drops. Under age 2: Consult a doctor.

**Active Ingredient:** Pulpa dentis (Bovine dental pulp) 30X

**Inactive Ingredient:** Distilled water

**Use:** Temporary relief of headache.

**Lot:**

**PULPA DENTIS 30 SPECIAL ORDER**

pulpa dentis 30 special order liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-8324
<b>Route of Administration</b>	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
SUS SCROFA TOOTH (UNII: V69U5FL51F) (SUS SCROFA TOOTH - UNII:V69U5FL51F)			SUS SCROFA TOOTH	30 [hp_X] in 1 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8324-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	
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(48951-8324)