

BETONICA ROSMARINUS- betonica rosmarinus 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.

Betonica Rosmarinus

Directions: FOR ORAL USE.

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

Active Ingredients: Betonica (Wood betony) 3X, Rosmarinus (Rosemary) 3X

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 www.urielpharmacy.com

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Made by Uriel, East Troy, WI 53120
www.urielpharmacy.com Lot:



**Betonica
Rosmarinus**

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

Betonica Rosmarinus

BETONICA ROSMARINUS

betonica rosmarinus liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-2062
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
STACHYS OFFICINALIS (UNII: UO9989Y17N) (STACHYS OFFICINALIS - UNII:UO9989Y17N)	STACHYS OFFICINALIS	3 [hp_X] in 1 mL
ROSMARINUS OFFICINALIS FLOWERING TOP (UNII: 8JM482TI79) (ROSMARINUS OFFICINALIS FLOWERING TOP - UNII:8JM482TI79)	ROSMARINUS OFFICINALIS FLOWERING TOP	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:48951-2062-1	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(48951-2062)

Revised: 5/2018

Uriel Pharmacy Inc.