BRYOPHYLLUM E FOL. 50% SPECIAL ORDER- bryophyllum e fol. 50% 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.

Bryophyllum e fol. 50% 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: 100g contains: 50g Bryophyllum e fol. 1X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

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

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e fol. IX
Inactive Ingredients: Datiled water, 20% Organic cane olasho
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Overstenn? Call 866.642.2659
Made with care by Unol. East Troy, WI 53120

www.unielpharmacy.com Lot:



Bryophyllum e fol. 50% Special Order

Homeopathic Liquid net vol. 2 fl. oz (60ml) REFRICERATE AFTER OPENING. BEST WHEN USED WITHIN 90 DAYS OF OPENING KEED OUT OF REACH OF CHILDREN.

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Todact Imornation			Product Information				
roduct Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-2125				
oute of Administration	ORAL						

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
KALANCHO E DAIGREMO NTIANA LEAF (UNII: L6 X13JKL8O) (KALANCHO E DAIGREMO NTIANA LEAF - UNII:L6 X13JKL8O)	KALANCHOE DAIGREMONTIANA LEAF	1 [hp_X] in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:48951-212	- 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(48951-2125)	

Revised: 5/2018 Uriel Pharmacy Inc.