

BERBERIS QUARTZ- berberis quartz 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.

Berberis Quartz

Use: Temporary relief of sinus congestion.

FOR ORAL USE ONLY

Directions: Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow. Follow inside instructions to open ampule.

Warnings:

Do not use if allergic to any ingredient. Consult a physician if symptoms worsen. If pregnant or nursing, consult a physician before use.

KEEP OUT OF REACH OF CHILDREN.

Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

KEEP OUT OF REACH OF CHILDREN.

Active Ingredients: Berberis (Barberry) 3X, Tunica mucosa nasi (Bovine nasal mucosa) 7X, Quartz (Rock crystal) 20X

Inactive Ingredients: Water, Salt

"prepared using rhythmical processes"

Questions? Call 866.642.2858

Uriel, East Troy, WI 53120

shopuriel.com Lot:

Use: Temporary relief of sinus congestion.

Directions: Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow. Follow inside instructions to open ampule.

Warnings:
NOT FOR INJECTION
Do not use if allergic to any ingredient. Consult a physician if symptoms worsen. If pregnant or nursing, consult a physician before use.
KEEP OUT OF REACH OF CHILDREN.
Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

Active Ingredients: Berberis (Barberry) 3X, Tunica mucosae (Bovine nasal mucosa) 7X, Quartz (Rock crystal) 20X

Inactive Ingredients: Water, Salt

Prepared using rhythmical processes.

Questions? Call 866.642.2858
Uriel, East Troy, WI 53120
shopuriel.com
Lot:

Uriel
Berberis Quartz
Homeopathic Ampules
FOR ORAL USE ONLY
net vol. 0.3 fl. oz (10 x 1 ml)

Berberis Quartz

BERBERIS QUARTZ

berberis quartz liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-2054
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	3 [hp_X] in 1 mL
BOS TAURUS NASAL MUCOSA (UNII: 343455G79K) (BOS TAURUS NASAL MUCOSA - UNII:343455G79K)	BOS TAURUS NASAL MUCOSA	7 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	20 [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:48951-2054-1	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(48951-2054)

Revised: 1/2026

Uriel Pharmacy Inc.