

KALIUM CARBONICUM 12 SPECIAL ORDER- kalium carbonicum 12 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.

Kalium carbonicum 12 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: Kalium carbonicum 12X

Inactive Ingredient: Distilled water

Use: Temporary relief of skin rash.

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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**Kalium
carbonicum 12X
Special Order**

Homeopathic Liquid
net vol. 2 fl. oz (60ml)

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: Kalium carbonicum 12X

Inactive Ingredients: Distilled water

Use: Temporary relief of skin rash.

Lot:

KALIUM CARBONICUM 12 SPECIAL ORDER

kalium carbonicum 12 special order liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-6067
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION -	POTASSIUM	12 [hp_X]

UNII:7UJQ5OPE7D)			CARBONATE	in 1 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-6067-3	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(48951-6067)

Revised: 4/2018

Uriel Pharmacy Inc.