

NATRUM MURIATICUM KIT REFILL- sodium chloride pellet
Washington Homeopathic Products

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.

DRUG FACTS

ACTIVE INGREDIENTS

NATRUM MURIATICUM 200C

USES

To relieve the symptoms of sneezing.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

NATRUM MURIATICUM Sneezing

STOP USE AND ASK DOCTOR

If symptoms persist/worsen or if pregnant/nursing, stop use and consult your practitioner.

DIRECTIONS

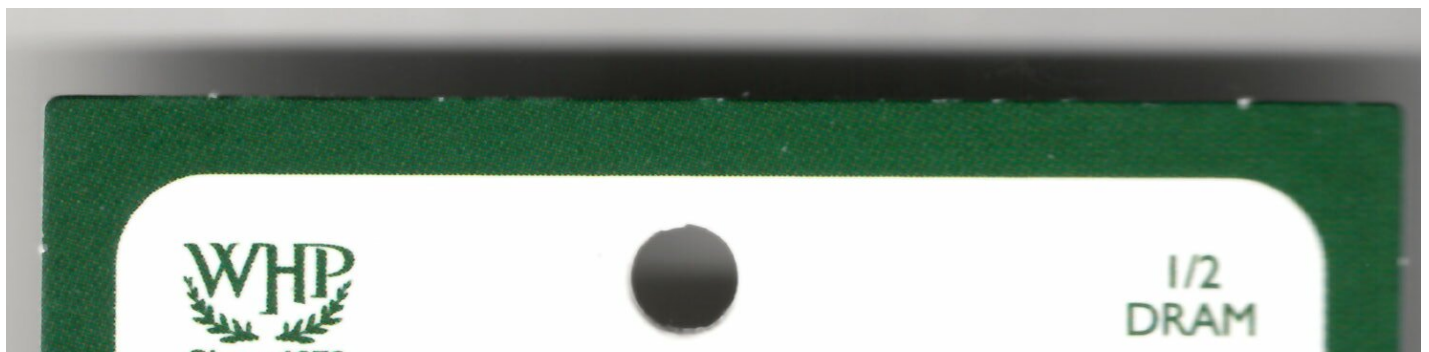
Adults: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides.

Children: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL



Since 1873

CONTENTS: **HOMEOPATHIC MEDICINE (HPUS*)**

DIRECTIONS: (Adults or Children): Dissolve 3 to 5 pellets under the tongue 3 times a day or as directed by a licensed practitioner. Take at greater intervals as condition subsides.

WARNING: If symptoms worsen or persist for more than 7 days, or if pregnant or nursing, consult your practitioner. Keep out of reach of children.

* The letters 'HPUS' indicate that the component in this product is officially monographed in the Homeopathic Pharmacopoeia of the United States.

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

DO NOT USE IF SEAL IS BROKEN.

SUCROSE/LACTOSE PELLETS



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EST. # 3006489197

NATRUM MURIATICUM KIT REFILL

sodium chloride pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68428-061
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:L4M0NH37)	SODIUM CHLORIDE	200 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics

Color	white (white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68428-061-01	750 in 1 VIAL, GLASS; Type 0: Not a Combination Product	05/03/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/03/2010	

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(68428-061)

