NATRUM MURIATICUM 200CK- natrum muriaticum pellet
SEVENE USA

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.

Sevene USA (as PLD) - NATRUM MURIATICUM 200ck (76472-3026)

ACTIVE INGREDIENT

HOMEOPATHIC DILUTION OF HPUS NATRUM MURIATICUM 200ck **C,K, CK, AND X ARE HOMEOPATHIC DILUTIONS.

TRADITIONALLY USED FOR

Stinging edemas from stings or allergies, relieved by cold*

DIRECTIONS

(adults/children) Dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a physician.

USE

Condition listed above or as directed by a physician.

WARNINGS

Stop use and ask a physician if symptoms persist for more than 3 days or worsen.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

OTHER INFORMATION

Store at room temperature.

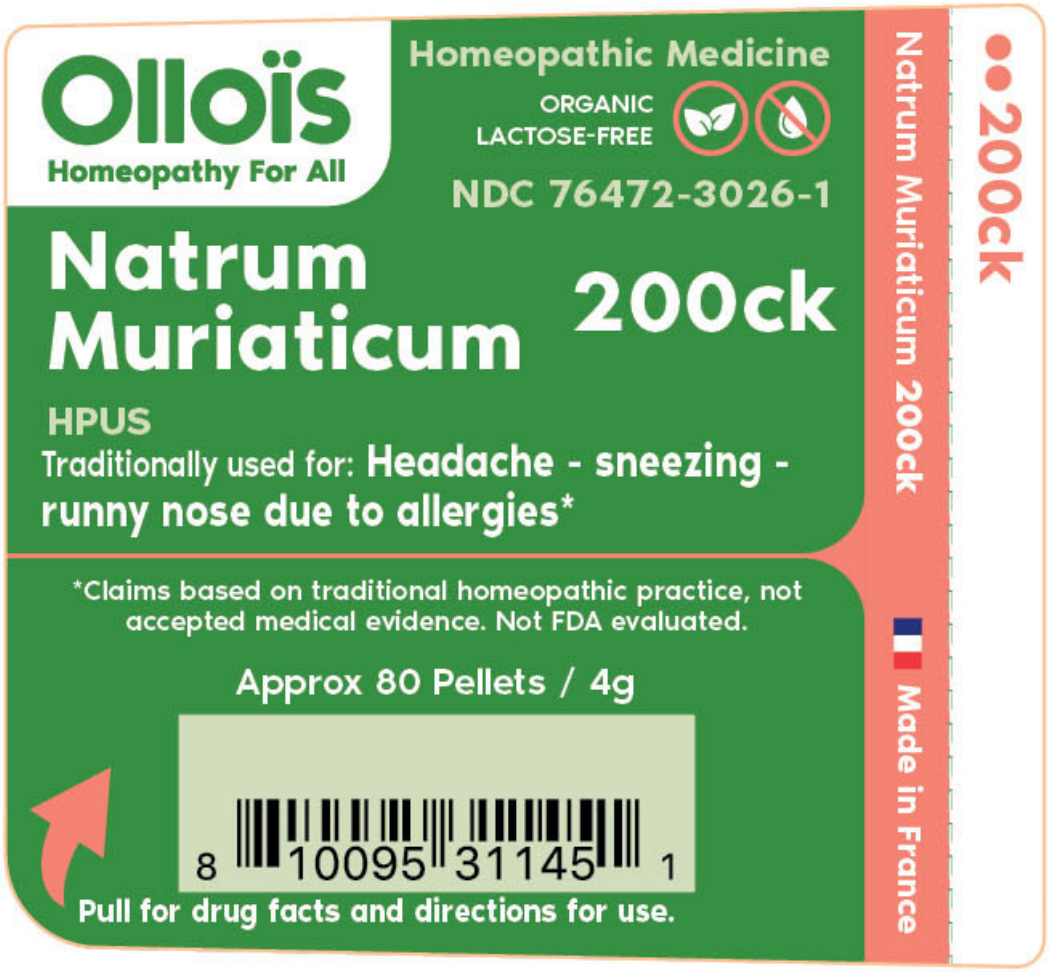
Do not use if pellet-dispenser seal is broken.

INACTIVE INGREDIENT

Sucrose.

QUESTIONS?

INFO@OLLOIS.COM * www.ollois.com * MADE IN FRANCE. NOT REVIEWED BY THE FDA AND NOT GUARANTEED TO BE EFFECTIVE. THIS HOMEOPATHIC DILUTION MAY NOT BE SUSCEPTIBLE TO SCIENTIFIC MEASUREMENT.



Drug Facts

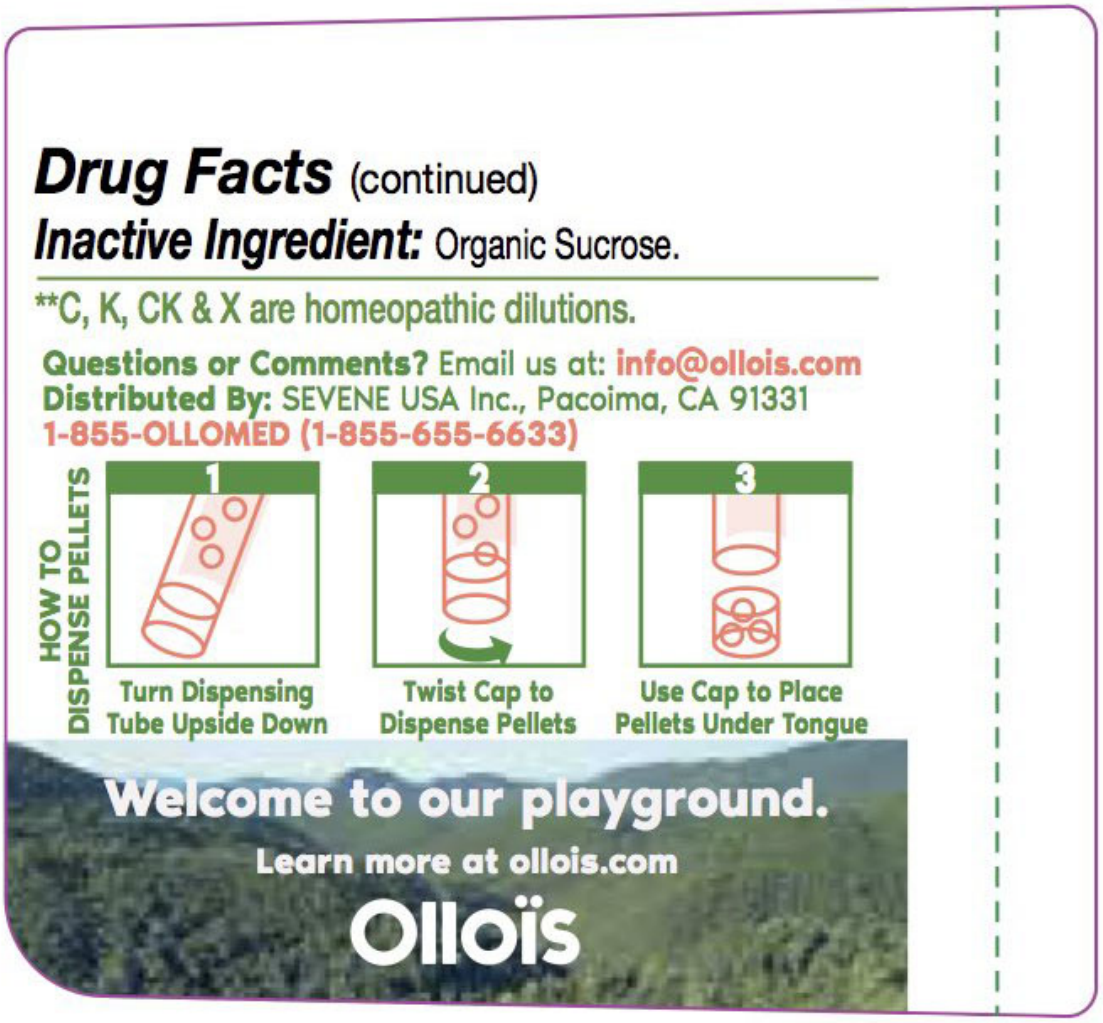
Active Ingredient: Natrum Muriaticum 200CK**

Uses: Headache - sneezing - runny nose due to allergies*.

Warnings: ■ **Stop use and ask a doctor** if symptoms persist for more than 3 days or worsen.
■ **If pregnant or breastfeeding**, ask a health professional prior to use. ■ **Keep out of reach of children.**
■ **In case of overdose**, get medical help.

Directions: ■ (Adults/Children) Dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

Other Information: ■ **Store at room temperature.**
■ **Do not use** if tamper-evidence seal is broken.
■ Organic verified by Alpes Contrôle 2021/A03M216K/1.



NATRUM MURIATICUM 200CK

natrum muriaticum pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76472-3026
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	200 [kp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Product Characteristics			
Color	white	**Score**	
Shape	ROUND	**Size**	4mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76472-3026-1	80 in 1 CYLINDER; Type 0: Not a Combination Product	11/23/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/23/2011	

Labeler - SEVENE USA (969332936)

Revised: 9/2023 SEVENE USA