NATRUM MURIATICUM- sodium chloride tablet Hyland's 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.

#9 Natrum Muriaticum 6X

DIRECTIONS

Adults & Children ages 7 - 12: 4 tablets. **Children ages 2 - 6:** 2 tablets. Dissolve under tongue 3 times a day. Use more frequently (every 15 minutes for up to 8 doses) with acute conditions.

INDICATIONS

Relief of symptoms of headaches, colds, heartburn, gastric upset and distress

FORMULA

Natrum Muriaticum 6X HPUS

In a base of Acacia Gum, Lactose N.F.

"HPUS" indicates that the active ingredients are in the official Homeopathic Pharmacopœia of the United States.

Warnings

Do not use if imprinted cap band is broken or missing.

If symptoms persist for more than 7 days or worsen, contact a licensed health care provider.

If you are pregnant or nursing, seek the advice of a licensed health care provider before using this product.

Keep all medications out of the reach of children.

QUESTIONS?

(800) 624-9659

PRINCIPAL DISPLAY PANEL - 500 Tablet Bottle Label HOMEOPATHIC

NDC 54973-6433-1

Hyland's[®] #9

Nat. Mur. 6X

Headaches, Colds,
Heartburn, Gastric Upset
and Distress*

500 TABLETS



NATRUM MURIATICUM

sodium chloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54973-6433
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	6 [hp_X]	

Inactive Ingredients		
Ingredient Name	Strength	
ACACIA (UNII: 5C5403N260)		

LACTOSE (UNII: J2B2A4N98G)			
Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	9mm

Imprint Code

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:54973- 6433-1	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1960		
2	NDC:54973- 6433-2	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1960	12/29/2021	

Marketing Information			
eting End Date			

Labeler - Hyland's Inc. (008316655)

Flavor Contains

Establishment			
Name	Address	ID/FEI	Business Operations
Hyland's Inc.		008316655	manufacture(54973-6433) , pack(54973-6433) , label(54973-6433)

Revised: 12/2022 Hyland's Inc.