## NA-ZONE - sodium chloride lotion Snuva, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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### ACTIVE INGREDIENT

**SODIUM CHLORIDE** 

### **INACTIVE INGREDIENTS**

BENZALKONIUM CHLORIDE DISODIUM EDTA DEIONIZED WATER SODIUM BICARBONATE

### **PURPOSE**

STERILE SALINE NASAL SPRAY

#### **USE**

SAFETY TO USE AS OFTEN AS NEEDED

### KEEP OUT OF REACH OF CHILDREN

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### **DIRECTIONS**

DIRECTIONS - WITH BOTTLE UPRIGHT SQUEEZE FIRMLY AND QUICKLY TO ATOMIZE NA-ZONE INTO EACH NOSTRIL. RELEASE BOTTLE TO VENT AND REPEAT 2 OR 3 TIMES

### **WARNINGS**

STERILITY GUARANTEED UNTIL THE SEAL IS BROKEN. NO SIDE EFFECTS ONCE SEAL IS BROKEN USE BY ONLY ONE PERSON IS RECOMMENDED

### **NA-ZONE LABEL**

NA-ZONE STERILE SALINE NASAL SPRAY
MANUFACTURED FOR SNUVA INC 715 SOUTH BLVD OAK PARK IL 2 FL OZ (59 ml)

# NA-ZONE STERILE SALINE NASAL SPRAY

STERILITY GUARANTEED UNTIL THE SEAL IS BROKEN NO SIDE EFFECTS SAFE TO USE AS OFTEN AS NEEDED

NDC 58291-0011-01



2 FL. OZ. (59 mL)

# DIRECTIONS -

With bottle upright squeeze firmly and quickly to atomize NA-ZONE into each nostril. Release bottle to vent and repeat 2 or 3 times. Once seal is broken, use by only one person is recommended.

### **ACTIVE INGREDIENTS**

Sodium Chloride

Benzylkonium Chloride

Disodium EDTA

Inert Ingredients

(inlcuding Sodium Bicarbonate as a buffer)

0.65%
0.01%
0.01%
99.33%



### **NA-ZONE**

sodium chloride lotion

P	ro	duct	<b>Information</b>
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:58291-0011

Route of Administration TOPICAL

### **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
SODIUM CHI ODIDE (LINII: 4513/47109 V) (SODIUM CATION LINII: I VD4M0 NID 7)	SODILIM CHI ODIDE	65 mI in 100 mI

### **Inactive Ingredients**

Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
EDETATE DISO DIUM (UNII: 7FLD91C86K)		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
SODIUM BICARBONATE (UNII: 8 MDF5 V39 QO)		

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:58291-0011-1	59 mL in 1 BOTTLE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	0 1/0 1/19 8 9	

## **Labeler -** Snuva, Inc. (807310008)

### **Registrant** - Snuva, Inc. (807310008)

Establishment			
Name	Address	ID/FEI	Business Operations
Snuva, Inc.		807310008	manufacture

Revised: 11/2010 Snuva, Inc.