AIR AND WATER HAND SANITIZER - SNOW MINT- ethyl alcohol gel
Pearl World 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.

AIR&WATER Snow Mint Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

• To decrease bacteria on the skin that could cause disease. • When water, soap and towel are not available. • Recommended for repeated use.

Warnings

For external use only: hands.

Flammable, Keep away from fire or flame.

When using this product • Keep out of eyes, ears, or mouth. • In case of eye contact, flush eyes thoroughly with water • Avoid contact with

broken skin • Do not inhale or ingest.

Stop use and ask a doctor if • Redness or irritation develop • condition persists for more than 72 hours.

Keep out of reach of children. • Children should be supervised by an adult when using this product. • If swallowed, get medical help or contact a Poison Control Center right away.

Directions • Wet hands thoroughly with product and allow to dry without wiping. • No rinsing required. • For children under 6, use only under adult supervision. • Not recommended for infants.

Other information • Store below 105°F (40°C) • May discolor certain fabrics. • Harmful to wood finishes & plastics.

Inactive ingredients: Water (Aqua), Triethanolamine, Carbomer, Aloe Barbadensis Leaf Extract, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E). Main Contain: FD & C Red 40 (CI 16035), FD & C Yellow 6 (CI 15985), FD & C Violet 2 (CI 60725), FD & C Blue 1 (CI 42090), FD & C Yellow 5 (CI 19140), FD & C Red 33(CI 17200).

70% Alcohol

Manufactured for and distributed by Pearl World Inc, NY, NY 10019

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Packaging

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ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69933-402
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
D&C VIOLET NO. 2 (UNII: 350KA7O6HK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69933-402-50	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/17/2020	
2	NDC:69933-402-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/17/2020	

Labeler - Pearl World Inc. (043130142)

Revised: 9/2020 Pearl World Inc.