NATURALL - ADVANCED HAND SANITIZER- alcohol spray Cosmetic Solutions LLC

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.

NaturAll - Advanced Hand Sanitizer

DRUG FACTS:

Active Ingredients

Ethyl Alcohol 71% v/v

Purpose

Antimicrobial

USES

Hand sanitizer to help reduce bacteria on the skin.

Recommended for repeated use.

WARNINGS

- Flammable. Keep away from fire or flame.
- For external use only.
- When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- **Stop use and seek a doctor** if redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children. Children under 6 years of age should be supervised when using this product. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

Put enough product in your palm to cover hands and rub hands together briskly until dry.

INACTIVE INGREDIENTS

Water (Aqua), Glycerin, Propanediol, Sodium Hyaluronate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Caprylic/Capric Triglyceride, Tocopherol, Carthamus Tinctorius Oleosomes, Triethanolamine, Phenoxyethanol, Ethylhexylglycerin, Fragrance.

OTHER INFORMATION

Store at 68°F to 77°F (20°C to 25°C). May discolor fabrics.

Distributed By:

NaturAll Club

PRINCIPAL DISPLAY PANEL - 60 mL Bottle Label

natur•all

ADVANCED HAND SANITIZER

Ethyl Alcohol Antimicrobial 71% Solution

2 fl. oz. | 60mL

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drug facts: (continued)

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Distributed By:

NaturAll Club Cleveland, OH 44114 www.naturallclub.com 800-683-8576 LB38351

NATURALL - ADVANCED HAND SANITIZER

alcohol spray

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:66163-1601

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength	
Alcohol (UNII: 3K9958 V90 M) (Alcohol - UNII: 3K9958 V90 M)	Alcohol	71 mL in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Glycerin (UNII: PDC6A3C0OX)			
Propanediol (UNII: 5965N8W85T)			
Hyaluronate Sodium (UNII: YSE9PPT4TH)			
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)			
Medium-Chain Triglycerides (UNII: C9H2L21V7U)			
Tocopherol (UNII: R0ZB2556P8)			
CARTHAMUS TINCTORIUS SEED OLEOSOMES (UNII: 9S60Q72309)			
Trolamine (UNII: 9O3K93S3TK)			
Phenoxyethanol (UNII: HIE492ZZ3T)			
Ethylhexylglycerin (UNII: 147D247K3P)			

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:66163-1601- 1	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020			
2	NDC:66163-1601- 2	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC MONOGRAPH NOT FINAL	part333A	04/01/2020			

Labeler - Cosmetic Solutions LLC (807907928)

Revised: 5/2020 Cosmetic Solutions LLC