ADVANCED HAND SANITIZER- ethyl alcohol gel
RITE AID

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.

Sea Mineral Advanced Hand Sanitizer
955.000/955AA-AB

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only-hands

Flammable. Keep away from heat and flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children.

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
- for children under 6, use only under adult supervision

- not recommended for infants

Other information

- do not store above 105º F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glyceryl, caprylate/caprate, glycerin, tocopheryl acetate, aloe barbadensis leaf juice, isopropyl myristate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4, blue 1, ext. violet2

*Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

DISTRIBUTED BY: RITE AID

30 HUNTER LANE

CAMP HILL, PA 17011

Principl display panel

daylogic

Advance Hand Sanitizer

Sea mineral scent

more effective formula

kills more than 99.99% of germs

Moisturizing Formula with Vitamin E

8 FL OZ (236 mL)

ADVANCED HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-0955
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)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0955-1	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2013	
2	NDC:11822-0955-2	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/13/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/13/2013	

Labeler - RITE AID (014578892)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11822-0955)

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Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(11822-0955)

Revised: 4/2023