

TRAVEL ASSURE ADVANCED HAND SANITIZER- isopropyl alcohol spray
Dermapeutics

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.

TRAVEL ASSURE ADVANCED HAND SANITIZER SPRAY

Drug Facts

Active Ingredient

Isopropyl Alcohol 70%

Purpose

Antimicrobial

USE

- For hand washing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

- **Do not use** in the eyes. In case of contact rinse thoroughly with water
- **Do not use** on children less than 2 months of age.
- **For external use only.**
- **Flammable, keep away from heat and flame.**
- **Stop use and ask doctor if**
- Irritation and redness develop and persist for more than 72hrs
- Swallowed, get medical attention or contact poison control center right away.

Keep Out of Reach Of Children. Children must be supervised in use of this product.

Directions:

- Spray product into palm to cover hands and rub hands together until dry.
- Children under 6 years of age should be supervised when using this product.

Other Information:

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

Inactive Ingredients: Glycerin, Water, Coco-Caprylate, Citrus Aurantium Dulcis Peel Oil.

kills 99.9% of germs

Distributed By:

Dermapeutics Corp

Calabasas, CA 91302

NO ANIMAL TESTING

Made in the USA

WWW.DERMAPEUTICS.COM

Packaging



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isopropyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79416-556
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
COCOYL CAPRYLOCAPRATE (UNII: 8D9H4QU99H)	
ORANGE OIL (UNII: AKN3KSD11B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79416-556-01	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/07/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/07/2020	

Labeler - Dermapeutics (859082237)

Revised: 7/2020

Dermapeutics