INSTA CLEAN HAND SANITIZER- ethyl alcohol gel Omega Packaging Corporation

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.

Insta Clean Hand Sanitizer

Active Ingredient

Ethyl alcohol 62%. w/w.....Antiseptic

Purpose

Antiseptic

For handwashing to decrease bacteria on the skin.

Warnings

Flammable, keep away from fire or flame. For external use only.

When using this produc

Avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor

If redness or irritation develop and persist 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

apply palmful to hand, rub hand together vigorously until dry, supervise children in the use of this product.

other information

store below 110 °F.(43.33 °C) may discolor certain fabrics or surfaces.

Inactive Ingredients: Water, Isopropyl Alcohol, Glycerin, Carbomer, Aminomethyl Propanol, Fragrance, Propylene Glycol, Isopropyl Myristate, Aloe Barbadensis Leaf Juice (Aloe Vera) Tocopheryl Acetate (Vitamin E)

Packaging



INSTA CLEAN HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57768-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		

ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)	
AMINO METHYLPRO PANO L (UNII: LU49 E6626Q)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	
ALOE (UNII: V5VD430 YW9)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:57768-001-17	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/13/2020	

Labeler - Omega Packaging Corporation (361281991)

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