MAD BEAUTY HAND SANITIZER CANDY CANE- alcohol spray Mad Beauty USA 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.

Mad Beauty Hand Sanitizer, Candy Cane

DRUG FACTS

Active Ingredient

Ethylalcohol 69%

Purpose

Antimicrobial

Uses

Reduces bacteria on hands.

Warnings

Flammable. Keep away from source of ignition or flame.

For external use only.

When using this product

keep out of eyes.

Stop use and ask a doctor

if irritation or redness develops.

Keep out of reach of children.

If swallowed, get medical help or contact a doctor immediately.

Inactive Ingredients

Water(Aqua)/Eau, Propylene Glycol, Isopropyl Alcohol, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Isopropyl Myristate, Tocopheryl Acetate(Vitamin E), Fragrance(Parfum), Butylene Glycol, Aloe Barbadensis (Aloe Vera) Leaf Extract.

Directions

Spray into hands. Rub until absorbed.

Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

Store between 15-30 C (59-86 F).

Avoid freezing and excessive heat above 40 C (104 F).

Questions or Comments

MAD BEAUTY USA LLC 1030 SALEM ROAD UNION NJ 07083 MARYLAND TEL (844) 995 1701

Package Labeling:







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MAD BEAUTY HAND SANITIZER CANDY CANE

alcohol spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78789-028

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.69 mL in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)		
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:78789-028- 00	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2020	12/31/2024		

Marketing In	formation	on		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/20/2020	12/31/2024	

Labeler - Mad Beauty USA LLC (117508758)

Revised: 12/2021 Mad Beauty USA LLC