

DISNEY BAMBI MOISTURIZING HAND SANITIZER MELON- alcohol gel
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.

Disney Bambi Moisturizing Hand Sanitizer, MELON

DRUG FACTS

Active Ingredient

Ethylalcohol 69%

Purpose

Antimicrobial

Uses

Reduce bacteria on hands.

Warnings

Flammable. Keep away from source of Ignition or flame.

For external use only.

Do not use

on open skin wounds

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.

Directions

Spray into hands. Rub until absorbed.

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

Inactive Ingredients

Water(Aqua)/Eau, pentylene Glycol, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Butylene Glycol, Fragrance(parfum), Sodium Hydroxide, Aloe Barbadensis (Aloe Vera)Leaf Extract.

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:

BAMBI

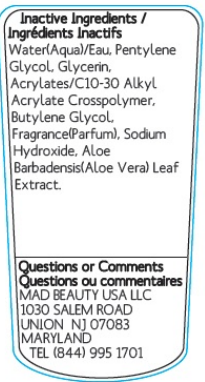
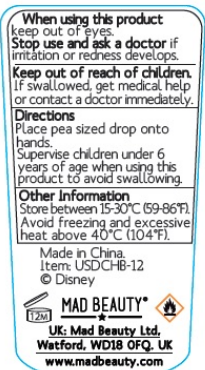
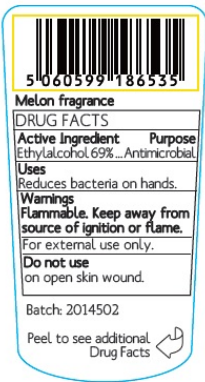
FRAGRANCE - MELON

LID **BLACK** METAL CLIP COLOUR 420C

French Translation



Front



DISNEY BAMBI MOISTURIZING HAND SANITIZER MELON

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78789-022
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)				
PENTYLENE GLYCOL (UNII: 50C1307PZG)				
GLYCERIN (UNII: PDC6A3C0OX)				
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78789-022-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2020	12/31/2024
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		07/20/2020	12/31/2024

Labeler - Mad Beauty USA LLC (117508758)