

**MAD BEAUTY POM POM MOISTURISING HAND SANITIZER, WILD BERRIES-  
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.*

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**Mad Beauty Pom Pom Moisturising Hand Sanitizer, Wild Berries**

**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.

**Do not use**

on open skin wound.

**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**

Place pea sized drop onto hands.

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).

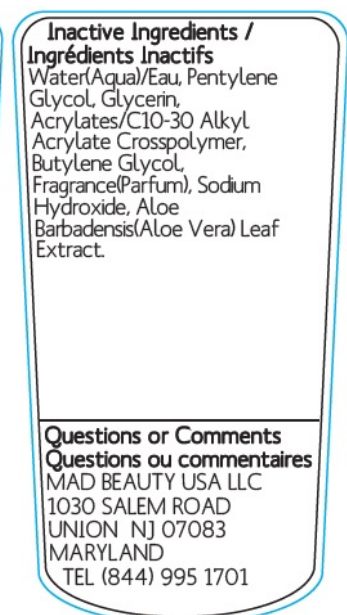
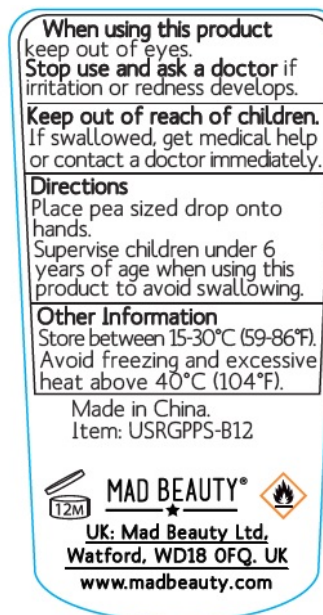
## Inactive Ingredients

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

## Questions or Comments

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## Package Labeling



## MAD BEAUTY POM POM MOISTURISING HAND SANITIZER, WILD BERRIES

alcohol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78789-008
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: 059QF0K00R)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
GLYCERIN (UNII: PDC6A3C00X)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
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:78789-008-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	12/31/2024

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2020	12/31/2024

**Labeler** - Mad Beauty USA LLC (117508758)

Revised: 12/2021

Mad Beauty USA LLC