

**DISNEY VILLAINS EVIL QUEEN HAND SANITIZER, POISON APPLE- 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 Villains Evil Queen Hand Sanitizer, Poison Apple

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

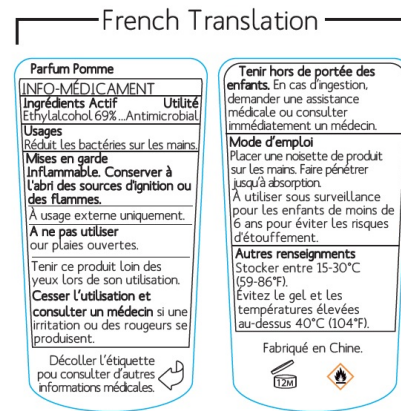
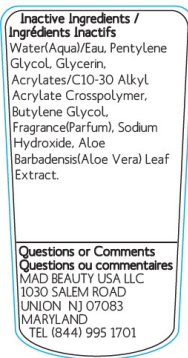
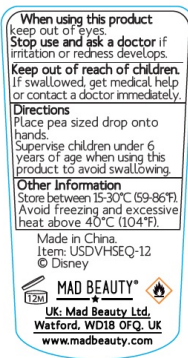
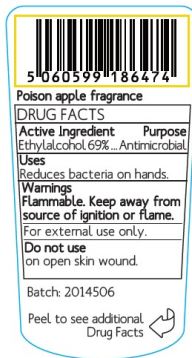
Water(Aqua)/Eau, Pentylene Glycol, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Butylene Glycol, Fragrance(Parfum), Sodium Hydroxide, Aloe Barbadensis(Aloe Vera) Leaf Extract.

Questions or Comments

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Package Labeling:

EVIL QUEEN - POISON APPLE



Front

DISNEY VILLAINS EVIL QUEEN HAND SANITIZER, POISON APPLE

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78789-004
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)	
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:78789-004-01	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