

AMORAY PREMIUM HAND SANITIZER- alcohol gel

My Imports 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.

Amoray Premium Hand Sanitizer

Drug Facts

Active ingredients

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- Hand sanitizer to help decrease bacteria on the skin.
- When water, soap & towel are not available.
- Recommended for repeated use.

Warnings

For external use only. Flammable. Keep away from fire or flame. Do not apply around eyes.

Do not use

in ears & mouth.

When using this product,

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.

Children must be supervised in use of this product.

Directions

- Pump as needed into your palms and thoroughly spread on both hands.
- Rub into skin until dry.

Other information

- Store at 20°C (68 to 77°F).
- May discolor fabrics.

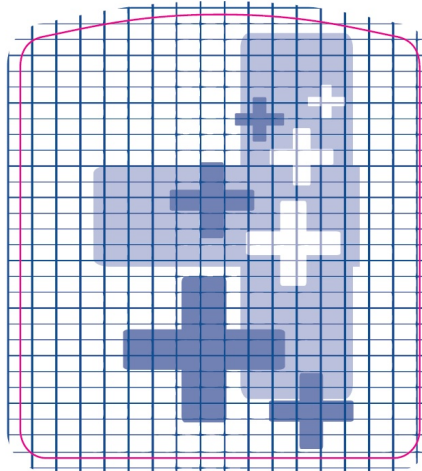
Inactive ingredients

- Water, triethanolamine, glycerin, carbomer, PEG-40 hydrogenated castor oil, propylene glycol, fragrance, tocopheryl acetate, aloe barbadensis gel.

Package Labeling:



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AMORAY PREMIUM HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51628-4221
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51628-4221-0	110 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/29/2020	

Labeler - My Imports USA LLC (195767988)

Revised: 10/2020

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