MERCI HANDY SANCTUARY HAND SANITIZER ARIES- alcohol gel
MERCI HANDY CORPORATION

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.

Merci Handy Sanctuary Hand Sanitizer ARIES

Drug Facts

Active ingredient

Alcohol 67% (v/v)

Purpose

Antiseptic

Use

- for handwashing to decrease bacteria on the skin

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

in the eyes. In the case of contact, flush eyes with water.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away

Directions

wet hands thoroughly with product and allow to dry without wiping

Other information

store at a temperature below 110° F (43° C)

Inactive Ingredients

Aqua (Water), Parfum (Fragrance), Aloe Barbadensis Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Zea Mays (Corn) Starch, Denatonium Benzoate, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, FD&C Red No.4, D&C Red No.33, D&C Red No.30

QUESTIONS OR COMMENTS?

(646)-358-3432

Package Labeling:

MERCI HANDY SANCTUARY HAND SANITIZER ARIES

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72866-018
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	67 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MANNITOL (UNII: 3OWL53L36A)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
CORN (UNII: 0N8672707O)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
D&C RED NO. 30 (UNII: 2S42T2808B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-018-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2021	

Labeler - MERCI HANDY CORPORATION (118006306)

Revised: 12/2022 MERCI HANDY CORPORATION