#### MERCI HANDY SANCTUARY HAND SANITIZER LIBRA- 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.

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#### Merci Handy Sanctuary Hand Sanitizer LIBRA

#### **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, Linalool, Hexyl Cinnamal, FD&C Blue No. 1, Ferric Ferrocyanide.

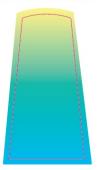
#### **QUESTIONS OR COMMENTS?**

(646)-358-3432

## **Package Labeling:**

#### LIBRA / US Production















## **MERCI HANDY SANCTUARY HAND SANITIZER LIBRA**

alcohol gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72866-024

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (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: 30WL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
<b>CORN</b> (UNII: 0N86727070)	
<b>DENATONIUM BENZOATE</b> (UNII: 4YK5Z54AT2)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.ALPHAHEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FERRIC FERROCYANIDE (UNII: TLE294X33A)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:72866-024- 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