HAND SANITIZER, HELLO SUNSHINE- 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.

Hand Sanitizer, Hello Sunshine

Drug Facts

Active ingredient

Alcohol 67%

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 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

Water (Aqua), Fragrance (Parfum), Aloe Vera Leaf Juice Powder, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Mannitol, Microcrystalline Cellulose, Sucrose, Corn (Zea Mays) Starch, DenatoniumBenzoate, Tocopheryl Acetate, Maltodextrin, Hydroxypropyl Methylcellulose, Potassium Sorbate, Sodium Benzoate, Amyl Cinnamal, Hexyl Cinnamal, Benzyl Salicylate, Alpha-Isomethyl Ionone, Amylcinnamyl Alcohol, FD&C Yellow No. 5, FD&C Red No. 4, Iron Oxides.

QUESTIONS OR COMMENTS?

(646)-358-3432

Package Labeling:

HAND SANITIZER, HELLO SUNSHINE

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72866-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	670 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MANNITOL (UNII: 3OWL53L36A)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
.ALPHA.-AMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
ISOMETHYL-.ALPHA.-IONONE (UNII: 9XP4LC555B)	
.ALPHA.-AMYLCINNAMYL ALCOHOL (UNII: DKB52S61GU)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-000-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2019	06/30/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/01/2019	06/30/2025

Labeler - Merci Handy Corporation (118006306)

Revised: 11/2022 Merci Handy Corporation