HAND SANITIZER- ethyl 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.

Active ingredient
Alcohol 67%

Purpose
Antiseptic

Uses

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

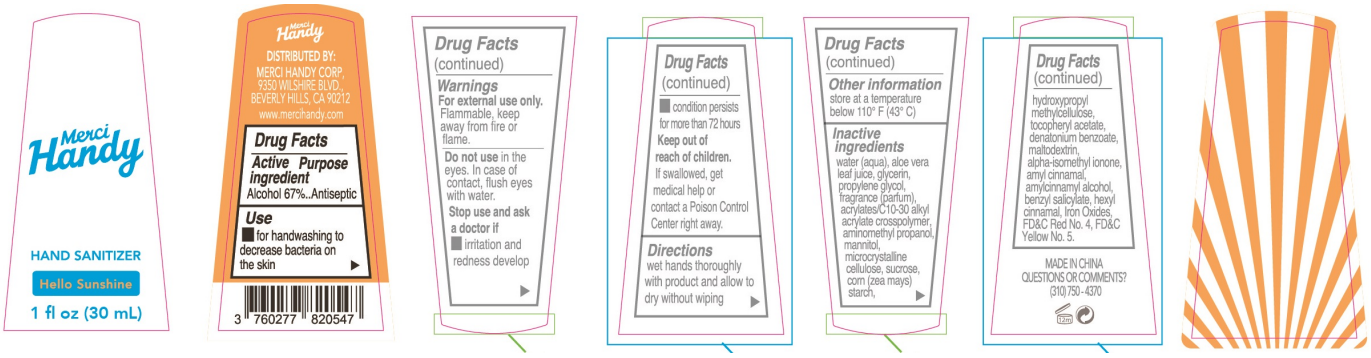
wet hand thoroughly with product and allow to dry without wiping

Other information

store at a temperature below 110 F(43 C)

Inactive ingredients

water(aqua), aloe vera leaf juice, glycerin, propylene glycol, fragrance(parfum), acrylates/C30-10 alkyl acrylate crosspolymer, aminomethyl propanol, mannitol, microcrystalline cellulose, sucrose, corn(zea mays) starch, hydroxpropyl methyl cellulose, tocopheryl acetate, denatonium benzonate, maltodextrin, alpha-isomethyl ionone, amyl cinnamal, amylcinnamyl alcohol, benzyl salicylate, hexyl cinnamal, Iron Oxides, FD&C Red No.4, FD&C Yellow No.5.



HAND SANITIZER
ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72866-103
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**
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
.ALPHA.-IONONE (UNII: I9V075M61R)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
.ALPHA.-AMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
MANNITOL (UNII: 3OWL53L36A)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
.ALPHA.-AMYLCINNAMYL ALCOHOL (UNII: DKB52S61GU)	
BROWN IRON OXIDE (UNII: 1N032N7MFO)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72866-103-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/18/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/18/2019	

Labeler - Merci Handy Corporation (116958007)