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

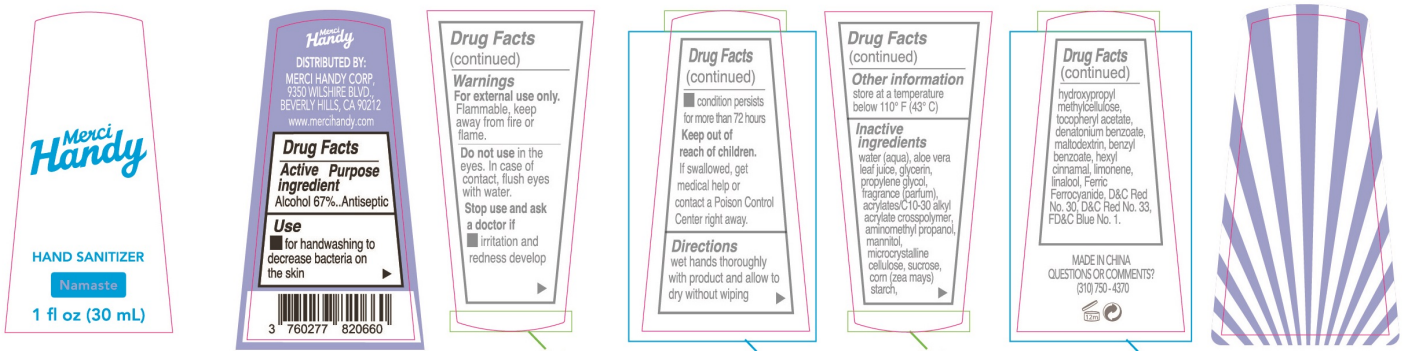
wet hand thoroughly with product and allow to dry without wiping

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, benzyl benzoate, hexyl cinnamal, limonene, linalool, Ferric Ferrocyanide, D&C Red No.30, D&C Red No.33, FD&C Blue No.1.



HAND SANITIZER
ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72866-105
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
D&C RED NO. 30 (UNII: 2S42T2808B)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
BENZYL BENZOATE (UNII: N863NB338G)	
FERRIC FERROCYANIDE (UNII: TLE294X33A)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
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)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
LINALOOL, (+)- (UNII: F4VNO44C09)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72866-105-01	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)