ANTIBACTERIAL HAND SANITIZER- ethyl alcohol gel Top Trenz

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: Ethyl Alcohol 62%

Purpose Antiseptic

Warnings For external use only. Flammable. Keep away from fire or flame.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a doctor right away.

Inactive Ingredients Water, Aloe Barbadensis Leaf Juice, Fragrance, Glycerin, Propylene Glycol, Carbomer, Aminomethyl Propanol, Lactose, Microcrystalline Cellulose, Sucrose, Zea Mays (corn) Starch, Ultramarine Blue CI 77007, Tocopheryl Acetate, Hydroxpropyl Methyl Cellulose, FD&C Red No.4, FD&C Yellow No.5.

Uses To help reduce bacteria on the skin.

Directions Pump as needed into your palms to cover hands. Rub hands together briskly unitl dry.





ANTIBACTERIAL HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70827-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		
LACTOSE (UNII: J2B2A4N98G)		
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
SUCROSE (UNII: C151H8 M554)		
STARCH, CORN (UNII: O8232NY3SJ)		
ULTRAMARINE BLUE (UNII: I39 WR9 9 8 BI)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
HYPROMELLOSES (UNII: 3NXW29 V3WO)		
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70827-001-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/18/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/18/2016	

Labeler - Top Trenz (040225026)

Revised: 7/2016 Top Trenz