PREMIUM HAND SANITIZER- alcohol liquid
MD Science Lab LLC

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.

PREMIUM HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 80%

Purpose

Antimicrobial

Use

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

- **For external use only.**
- **When using this product** avoid contact with eyes. In case of eye contact, flush eyes with water.
- **Stop use and ask a doctor if** irritation or redness develops, or if 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

- Dispense a small amount into palm of hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inactive Ingredients

Deionized Water, Glycerin, Hydrogen Peroxide.

PRINCIPAL DISPLAY PANEL - 3.7 liter Bottle Label

M.D.
Science Lab
LLC

PREMIUM
hand
sanitizer

Kills 99.99% of Germs

moisturizes & softens skin

1 gal 128 fl oz 3.7 liter e

PREMIUM HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69898-801
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	80 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	

Glycerin (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
TEA TREE OIL (UNII: VIF565UC2G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69898-801-03	88 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	
2	NDC:69898-801-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	
3	NDC:69898-801-12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	
4	NDC:69898-801-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	
5	NDC:69898-801-01	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	
6	NDC:69898-801-55	208197 mL in 1 DRUM; Type 0: Not a Combination Product	07/01/2020	
7	NDC:69898-801-27	1040988 mL in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product	07/01/2020	
8	NDC:69898-801-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	07/01/2020	

Labeler - MD Science Lab LLC (054484289)

Revised: 6/2020 MD Science Lab LLC