PREMIUM HAND SANITIZER- benzalkonium chloride 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

Benzalkonium Chloride 0.13%

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 a ask 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

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

Inactive Ingredients

Water (aqua), Propylene Glycol, Polysorbate 20, Phenoxyethanol, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Lavandula Angustifolia (Lavender) Oil, Aloe barbadensis (Aloe Vera).

PRINCIPAL DISPLAY PANEL - 237 ml Bottle Label

M.D. Science Lab LLC

PREMIUM hand

sanitizer with Tea Tree Oil & Aloe Kills 99.99% of Germs Antimicrobial - Alcohol Free moisturizes & softens skin 8 fl oz 237ml e





with Tea Tree Oil & Aloe

Kills 99.99% of Germs Antimicrobial - Alcohol Free



moisturizes & softens skin

8 fl oz 237mle



Kills 99.99% of Germs Antimicrobial - Alcohol Free

Swiss Navy® Premium Hand Sanitizer is alcohol-free, yet powerful and gentle. This formula with Tea Tree Oil and Aloe, moisturizes and softens the skin, and it's safe to use everyday.

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MATE USAT









Manufactured for: M.D. Science Lab, LLC 2131 Blount Road, Pompano Beach, FL 33069 Phone: 1-954-725-5502 USA www.swissnavy.com #swissnavy



PREMIUM HAND SANITIZER

benzalkonium chloride liquid

Product	Inform	ation
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:69898-501

TOPICAL Route of Administration

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	1.3 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Propylene Glycol (UNII: 6DC9Q167V3)			
Phenoxyethanol (UNII: HIE492ZZ3T)			
LAVENDER O IL (UNII: ZBP1YXW0 H8)			
TEA TREE O IL (UNII: VIF565UC2G)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
Polysorbate 20 (UNII: 7T1F30V5YH)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69898-501- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2020			
2	NDC:69898-501- 12	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2020			
3	NDC:69898-501- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2020			
4	NDC:69898-501- 01	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2020			
5	NDC:69898-501- 07	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC MONOGRAPH NOT FINAL	part333E	04/05/2020			

Labeler - MD Science Lab LLC (054484289)

Revised: 4/2020 MD Science Lab LLC