TOTAL SANITIZER ALCOHOL FREE HAND SANITING FOAM- benzalkonium chloride liquid

Total Sanitizer, 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.

Total Sanitizer Alcohol Free Hand Saniting Foam

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.12%

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin.
- After changing diapers.
- After assisting ill persons.
- Before contact with a person under medical care or treatment.
- Recommended for repeat use.

Warnigns

For external use only.

Do not use

in the eyes.

When using this product

do not get into eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctorif

irritation and redness develop. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands thoroughly with product and allow to dry.
- Children under six years of age should be supervised when using this product.

Other information

• Store in a cool, dry place.

Inactive Ingredients

Water, Aloe Barbadensis Leaf Juice, Polysorbate 20, Glycerin, Potassium Sorbate, Fragrnace, Citric Acid, Disodium EDTA, Tocopheryl Acetate (Vitamin E).

Package Labeling:

DESCRIPTION

This non-rinse, alcohol-free formula cleans hands while killing bacteria, and can safely be used in places where alcohol-based products are a safety risk.

This formula ideal for use in a health care environment, food processing or anywhere else bacterial contamination may be a problem.

Moisturizes the skin with Aloe, Glycerin and Vitamin E.





Elmhurst, IL 60126

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0000-100-410

TOTAL SANITIZER ALCOHOL FREE HAND SANITING FOAM

benzalkonium chloride liquid

Product Information

Inactive Ingredients

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80463-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Benzalko Nium Chloride (Unii: F5UM2KM3W7) (BENZALKONIUM - BENZALKONIUM CHLORIDE (Unii: F5UM2KM3W7) (BENZALKONIUM - CHLORIDE in 1 mL

Ingredient Name WATER (UNII: 059QF0KO0R) ALOE VERA LEAF (UNII: ZY81Z83H0X) POLYSORBATE 20 (UNII: 7T1F30V5YH) GLYCERIN (UNII: PDC6A3C0OX) POTASSIUM SORBATE (UNII: 1VPU26JZZ4) CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

EDETATE DISODIUM ANHYDROUS (UNII: 8 NLQ36 F6 MM)

.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:80463-003-	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020		
Marketing Information				
Marketing Inf	ormation			
Marketing Info		Marketing Start Date	Marketing End Date	
	y Application Number or Monograph Citation	Marketing Start Date 07/01/2020	Marketing End Date	

Labeler - Total Sanitizer, LLC (080216348)

Revised: 9/2020 Total Sanitizer, LLC