

**TOTAL SANITIZER ALCOHOL FREE ANTISEPTIC FOAM HAND SANITIZER-
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 Antiseptic Foam Hand Sanitizer

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

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

Warnings

- For external use only.

Do not use

- in eyes. If contact occurs, flush eyes with water.

Stop use and ask a doctor

- if irritation or redness develops. 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

- Pump a small amount of foam into palm of hand.
- Wet hands thoroughly with product and allow to dry without wiping.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide.

Package Labeling:

NO RINSE • MOISTURIZES • LEAVES SKIN SOFT

NDC # 49765-300-03



TOTAL SANITIZER ALCOHOL-FREE ANTISEPTIC FOAM HAND SANITIZER

Eliminates 99.999%
OF MOST COMMON GERMS THAT MAY CAUSE ILLNESS

NET WT 35.195 fl oz (1000mL)

Environmentally Friendly 

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Inactive Ingredients Water, dihydroxypropyl PEG -5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Manufactured for Total Sanitizer
5805 State Bridge Road, Duluth, GA 30097. Made in USA.



TOTAL SANITIZER ALCOHOL FREE ANTISEPTIC FOAM HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80463-002
Route of Administration	TOPICAL		

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
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WATER (UNII: 059QF0KO0R)	
DIHYDROXYPROPYL PEG-5 LINO LEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80463-002-00	1000 mL in 1 BOTTLE; 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 - Total Sanitizer, LLC (080216348)

Revised: 9/2020

Total Sanitizer, LLC