

TOTAL SANITIZER ALCOHOL FREE 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 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
FOAM HAND SANITIZER

Eliminates 99.99%
OF MOST COMMON GERMS THAT MAY CAUSE ILLNESS

Environmentally
Friendly

7.5 fl oz (220ml)

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

TOTAL SANITIZER ALCOHOL FREE FOAM HAND SANITIZER

benzalkonium chloride liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:80463-001 |
| 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 |
|--|----------|
| WATER (UNII: 059QF0K00R) | |
| DIHYDROXYPROPYL PEG-5 LINO LEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1) | |
| GLYCERETH-2 CO COATE (UNII: JWM00VS7HC) | |
| BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T) | |
| DIHYDROXYETHYL CO CAMINE O XIDE (UNII: 8AR51R3BL5) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:80463-001-00 | 220 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