TORK HAND SANITIZING ALCOHOL-FREE FOAM- benzalkonium chloride liquid
Essity Professional Hygiene North America 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.

Essity Alcohol-Free Foam Hand Sanitizer
D19.001/D19AC

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antiseptic

Uses

- Hand Sanitizer to help decrease bacteria on the skin.
- Recommended for repeated use

Warnings

For external use only

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product

- keep out of eyes, ears, and mouth
- In case of contact with eyes, rinse eyes thoroughly with water
- do not ingest

Stop use and ask a doctor if

- irritation and redness develop
- condition persists more than 72 hours

Keep out of reach of children:

if swallowed, get medial help or contact a Posion Control Center right away.

Directions

Dispense enough to spread on both hands to wrists, then rub into skin for 30 seconds. Allow to dry.

Inactive ingredients

water, lactic acid, phenoxyethanol, sodium hydroxide, propylene glycol, cocamidopropyl betaine, sodium cocoamphoacetate

Questions or comments?

1-866-722-8675

essity

Manufactured for

Essity Professional Hygiene

North America LLC

2929 Arch Street

Philadelphia, Pennsylvania 19104

www.essity.com, www.torkusa.com

made in USA from Domestic & Imported Components

Patents/Brevets/Patentes: www.essisty.com/patents

NOT FOR SALE

Can be ordered as: 401813 for S4, 1000ml

Principal panel display

Kills 99.9% of Common Germs*

*based on majority of microorgnisms tested under ASTM E2783-11 in 30 seconds

NDC 49351-303-02

TORK

Tork Hand Sanitizing

Alcohol-Free Foam

Benzalkonium Chloride, 0.13%

1.5 FL OZ (44 mL)

TORK HAND SANITIZING ALCOHOL-FREE FOAM

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49351-303
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM -	Benzalkonium	1.3 mg

UNII:7N6JUD5X6Y)	Chloride	in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LACTIC ACID (UNII: 33X04XA5AT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49351-303-02	44 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2022	
2	NDC:49351-303-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/18/2022	

Labeler - Essity Professional Hygiene North America LLC (005694349)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(49351-303)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(49351-303)

Revised: 9/2023 Essity Professional Hygiene North America LLC