

TECHNI-PRO ANTIBACTERIAL HAND CLEANER- benzalkonium chloride soap R & R Lotion, Inc

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.

Techni-Pro Antibacterial Hand Cleaner

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

For handwashing to decrease bacteria on the skin

Warnings

For external use only

When using this product avoid contact with eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if irritation or redness develops.

Keep out of reach of children. If swallowed, get medical help or contact a poison Control Center right away.

Directions

- Pump into DRY hands.
- Lather vigorously for at least 15 seconds
- Rinse and dry thoroughly

Inactive Ingredients

Water, Triethanolamine Lauryl Sulfate, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Cocamide Monoethanolamine, Sodium Chloride, Gluconolactone (and) Sodium Benzoate, Citric Acid, Sodium Hydroxide

PRINCIPAL DISPLAY PANEL - 946 mL Bottle Label

TECHNI-PRO

ANTIBACTERIAL HAND CLEANER

TestEquity Part #:
TNP-ICS-32

PACKAGING SIZE: 32oz.

TECHNI-PRO

ANTIBACTERIAL HAND CLEANER

Kills Germs & Paraben Free

A Hand Cleaner, formulated especially for the Electronic/Cleanroom/Pharmaceutical/ Food Service Industry.

Our advanced cleaning formula softens & protects your skin while effectively removing soils, makeup, and other contaminants.

Antibacterial Hand Cleaner helps to eliminate germs on your hands and is for external use only.

The Center for Disease Control states that "the most important measure for preventing the spread of pathogens is effective hand washing."

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RoHS Compliant  FDA Registered Manufacturer

Gov't Contractor GSA/VADOD



66080 85150 7

Packaged by R&R Lotion for Test Equity LLC
6100 Condon Dr.
Moorepark, CA 93021
800-950-3457
NDC # 59555-703-10

TECHNI-PRO ANTIBACTERIAL HAND CLEANER

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59555-703
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)	

Triethanolamine Lauryl Sulfate (UNII: E8458C1KAA)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59555-703-10	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/10/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	03/10/2022	

Labeler - R & R Lotion, Inc (062979000)

Revised: 3/2022

R & R Lotion, Inc