DOCTOR INKS SANITIZING- benzalkonium chloride soap
Enviro Specialty Chemicals 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.

DOCTOR INK'S SANITIZING SOAP

Drug Facts

Active Ingredient:

Benzalkonium Chloride 0.10%

Purpose:

Antiseptic Hand Soap

Uses:
- Hand wash to help reduce bacteria that potentially can cause disease
- Helps to prevent cross-contamination by hand contact
- Helps to prevent drying of skin • Recommended for repeat use

Warnings:
- For external use only
- When using this product do not use in or near eyes
- Stop use and ask doctor if irritation and redness develop

- Keep out of reach of children,
- If swallowed, seek medical attention or contact a Poison Control Center immediately.

Store in a cool dry place below 104° F (40°C)

Directions:
- Wet hands and wrists with clean running water
- Apply appropriate amount of product
- Lather & scrub hands, fingers, fingernails, cuticles & wrists thoroughly
- Rinse thoroughly with running water
- Dry hands completely (incomplete drying may result in chapped skin)

Inactive Ingredients:

Aloe Barbadensis Leaf Juice, Aqua, Benzyl Alcohol, Citric Acid, Glycerin, Lauramidopropyl

Betaine, Phenoxyethanol, Triethoxysilylpropyl Steardimonium Chloride.

Questions?

+1(888) 331-8332, M-F, 9AM-5PM (EST)

FDA Registered Anti-Microbial Soap for cleaning grime and germs from your hands

Kills 99.9% of Germs

*Formulated and enhanced with **Zetrisil®***

Alcohol Free

Contains Soothing Aloe Vera

Fast-acting
15-Second Formula

Foaming Pump Top

FORMULATED IN THE USA

MADE IN CHINA

Distributed by:
ESC Brands, LLC.
1060 Blue Prince Road
Bluefield, WV 24701
www.doctor-ink.com

Packaging

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FORMULATED IN THE USA | MADE IN CHINA

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E-1004-01

DOCTOR INKS SANITIZING
benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71884-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71884-200-01	118 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/30/2022	
2	NDC:71884-200-02	244 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/30/2022	
3	NDC:71884-200-03	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/30/2022	
4	NDC:71884-200-04	1182 mL in 1 BAG; Type 0: Not a Combination Product	06/30/2022	
5	NDC:71884-200-05	3785 mL in 1 JUG; Type 0: Not a Combination Product	06/30/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/30/2022	

Labeler - Enviro Specialty Chemicals Inc (202621850)

Revised: 2/2023

Enviro Specialty Chemicals Inc