DOCTOR INKS HAND SANITIZER- benzalkonium chloride liquid
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 HAND SANITIZER

Drug Facts

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose:

Antiseptic, Hand & Skin Sanitizer

Uses:

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

Warnings:

Do not freeze. For external use only. Do not use in ears, eyes or mouth.

- When using this product, avoid contact with the eyes. In case of contact, flush eyes with water.
- Stop use and ask a doctor if redness or irritation develop and persist for more than 72 hours.

- Keep out of reach of children.
- Children should be supervised when using this product.

Directions:

Apply liberally to the palms of the hands. Rub into skin until dry. Recommended for repeat use.

Inactive Ingredients:

Aloe Barbadensis leaf extract, Aqua, Citric Acid, Laureth-4, Phenoxyethanol, Triethoxysilylpropyl Steardimonium chloride.

Questions?

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

Long-lasting, alcohol-free protection from germs

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

Kills 99.9% of Germs

Alcohol Free

Contains Soothing Aloe Vera

Fast-acting
15-Second Formula

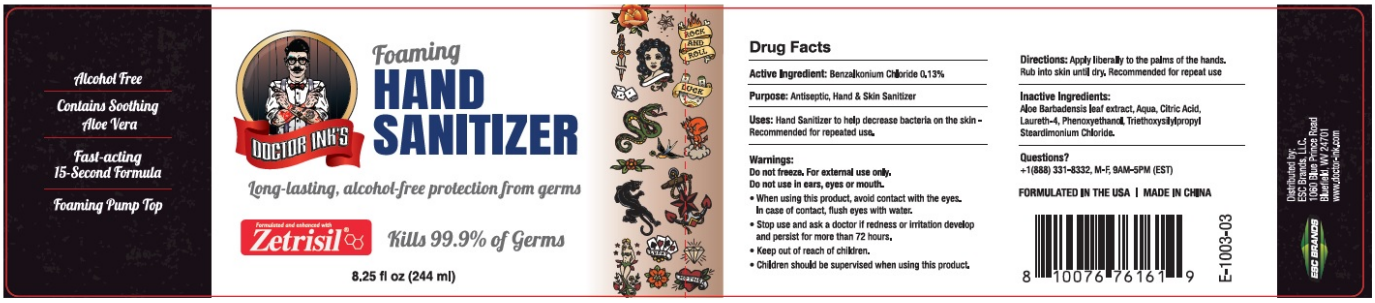
Foaming Pump Top

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

FORMULATED IN THE USA

MADE IN CHINA

Packaging



DRUG FACTS TABLE

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FORMULATED IN THE USA I MADE IN CHINA

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DOCTOR INKS HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71884-201
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
LAURETH-4 (UNII: 6HQ855798J)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71884-201-01	58 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/01/2022	
2	NDC:71884-201-02	118 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/01/2022	
3	NDC:71884-201-03	244 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/01/2022	
4	NDC:71884-201-04	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/01/2022	
5	NDC:71884-201-05	1182 mL in 1 BAG; Type 0: Not a Combination Product	09/01/2022	
6	NDC:71884-201-06	3785 mL in 1 JUG; Type 0: Not a Combination Product	09/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/01/2022	

Labeler - Enviro Specialty Chemicals Inc (202621850)

Revised: 2/2023 Enviro Specialty Chemicals Inc