

DOCTOR INKS TOPICAL ANTISEPTIC- benzalkonium chloride gel
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 TOPICAL ANTISEPTIC GEL

Drug Facts:

ACTIVE INGREDIENT:

Benzalkonium Chloride 0.13%

PURPOSE:

First aid antiseptic.
Topical skin treatment - Barrier Gel

Uses:

First aid soothing relief, reduced skin drying of newly applied tattoos, piercings, and germ protection for minor cuts, scrapes, burns, skin irritations and minor abrasions

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 develops and persists for more than 72 hours.

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

DIRECTIONS:

Apply liberally over new Tattoo artwork or piercings, recommended for repeat application regularly.

INACTIVE INGREDIENTS:

Aloe barbadensis leaf extract, aqua, laureth-4, citric acid, hydroxyethyl cellulose, phenoxyethanol, triethoxysilylpropyl Steardimonium chloride.

QUESTIONS?

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

Alcohol-Free

Long lasting, soothing protection from germs.

KILLS 99.9% OF GERMS

Formulated and enhanced with **Zetrisil®**

Contains Soothing Aloe Vera

***Fast-acting
15-second Formula***

DISTRIBUTED BY:

ESC Brands, LLC.
1060 Blue Prince Road
Bluefield, WV 24701
www.doctorink.com

Packaging

TAT-1001-1



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*Fast-acting
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2 FL OZ (59 ML)

DOCTOR INKS TOPICAL ANTISEPTIC

benzalkonium chloride gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:71884-203

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)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYISILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71884-203-01	30 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2022	
2	NDC:71884-203-02	59 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2022	
3	NDC:71884-203-03	118 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2022	
4	NDC:71884-203-04	237 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2022	
5	NDC:71884-203-05	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