

DOCTOR INKS SANITIZING BODY- benzalkonium chloride spray
Enviro Specialty Chemicals Inc

DOCTOR INK'S *Sanitizing* BODY SPRAY

DRUG FACTS:

ACTIVE INGREDIENT:

Benzalkonium Chloride 0.10%

PURPOSE:

Antiseptic

USES:

Body sanitizer to help reduce bacteria that may cause infection to new tattoo and piercings.

DIRECTIONS:

- Apply liberally to skin.
- Children under 6 years of age should be supervised when using this product.

INACTIVE INGREDIENTS

Aloe Barbadensis Leaf Juice Extract, Agua, Citric Acid, Deceth-6, Phenoxyethanol, Triethoxysilylpropyl Steardimonium chloride.

WARNINGS:

- For external use only.
- When using this product do not use in or near the eyes.
- In case of contact with eyes, rinse eyes thoroughly with water.
- Stop use and ask a doctor if irritation or rash occurs.

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

QUESTIONS:

+1(888) 331-8332, Mon-Fri 9AM-5PM (EST)
Email: info@escbrands.com

**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

Distributed by:

ESC Brands, LLC.

1026 Blue Prince Road, suite B

Bluefield, WV 24701

www.escbrands.com

FORMULATED IN THE USA | MADE IN CHINA

Packaging

Package: 50 mL

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DOCTOR INK'S

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1.7 fl oz. (50ml)

Drug Facts

Active Ingredient: Benzalkonium Chloride 0.10%

Purpose: Antiseptic

Uses: Body sanitizer to help reduce bacteria that may cause infection to new tattoo and piercings.

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Questions?
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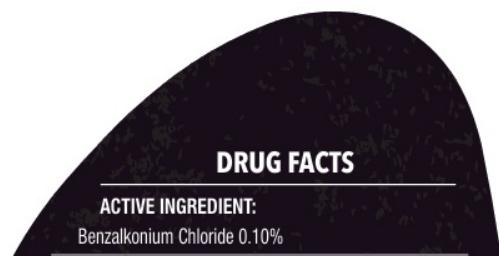
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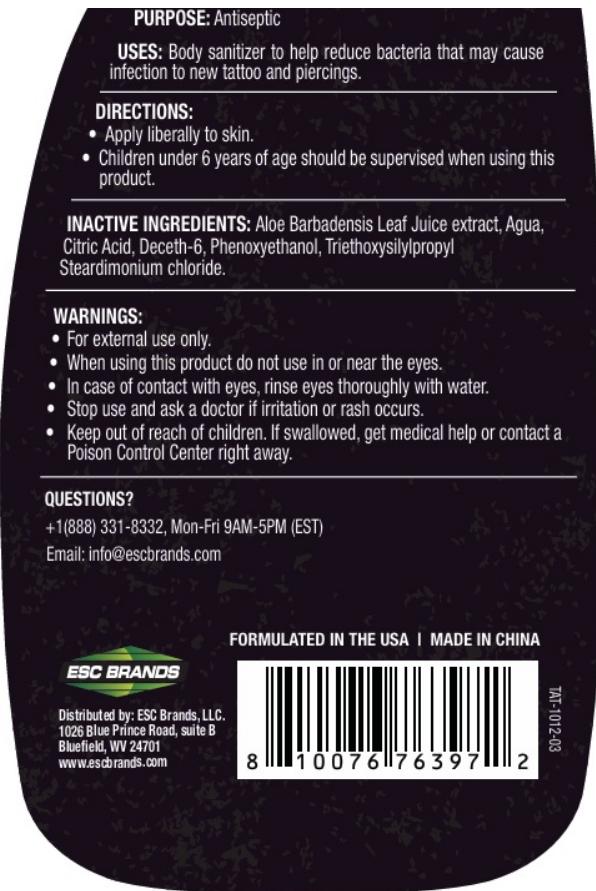


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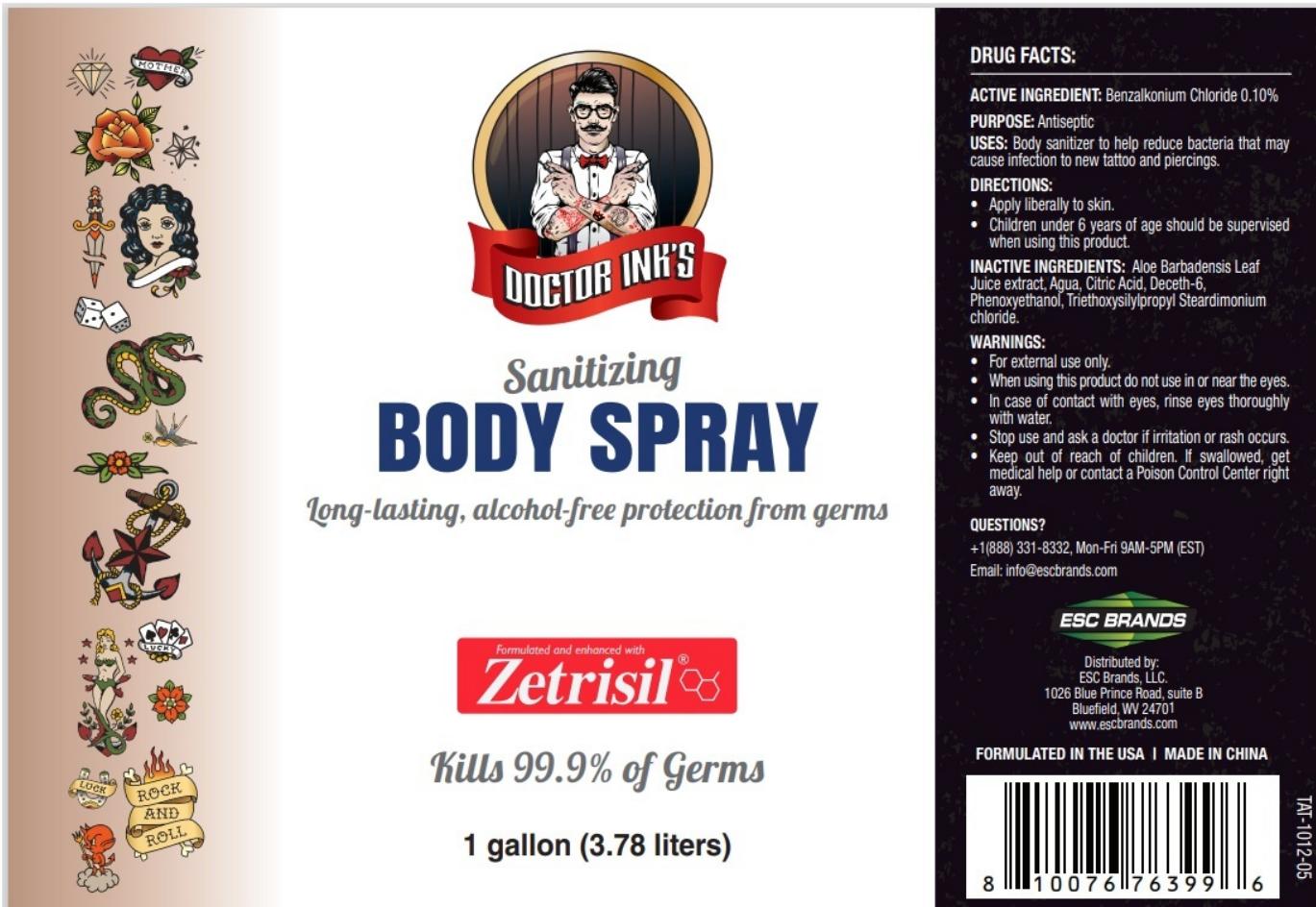
TAT-1012-01

Package: 473 mL





Package: 3780 ml



DOCTOR INKS SANITIZING BODY

benzalkonium chloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71884-202
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)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DECETH-6 (UNII: P4OYN7A209)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71884-202-01	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2026	
2	NDC:71884-202-03	473 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2026	
3	NDC:71884-202-05	3780 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2026	

Labeler - Enviro Specialty Chemicals Inc (202621850)

Revised: 1/2026

Enviro Specialty Chemicals Inc