

MY-SHIELD HAND SANITIZER GEL- 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.

MY-SHIELD HAND SANITIZER GEL

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 develops and persists for more than 72 hours.
- Children should be supervised when using this product.
- Keep out of reach of children.

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, Caprylyl Glucoside, Hydroxyethyl Cellulose, Polyhexanide, Phenoxyethanol, Triethoxysilylpropyl Steardimonium Chloride.

Questions?

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

FAST ACTING 15 SECOND FORMULA

WITH SOOTHING ALOE VERA

Long lasting, alcohol-free protection from germs for your hands.

KILLS 99.99% OF GERMS

Formulated and enhanced with **Zetrisil**

FORMULATED IN THE USA

MADE IN CHINA

DISTRIBUTED BY ESC BRANDS, LLC.

664 OLD HARGRAVES ROAD - SUITE B

LEXINGTON, NC 27295

www.myshield.com

Packaging

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E-1006-03



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30ML (1 FL OZ)

Drug Facts

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose:

Antiseptic/Hand & Skin Sanitizer

E-1006-01



Uses:

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2 FL OZ (57ML)

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4 FL OZ (118ML)

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6 FL OZ (177ML)



Alcohol-Free HAND SANITIZER

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8.25 fl oz (244 ml)



WITH SOOTHING
ALOE VERA

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630226 253284

E-1006-14

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MY-SHIELD HAND SANITIZER GEL

benzalkonium chloride gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:71884-011

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
POLYHEXANIDE (UNII: 322U039GMF)	
PHENOXYETHANOL (UNII: HE492ZZ3T)	
TRIETHOXYSYLILPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71884-011-01	3 in 1 PACKAGE	05/26/2020	
1		30 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:71884-011-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/26/2020	
3	NDC:71884-011-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/26/2020	
4	NDC:71884-011-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/26/2020	
5	NDC:71884-011-08	244 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/26/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/26/2020	

Labeler - Enviro Specialty Chemicals Inc (202621850)

Revised: 5/2020

Enviro Specialty Chemicals Inc