GERM FREE 24 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.

germfree 24[®] FOAMING 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, Caprylyl Glucoside, Laureth-4, Polyhexanide, Phenoxyethanol, Triethoxysilylpropyl Steardimonium Chloride.

Ouestions?

+1(888) 331-8332, M-F, 9:00AM-5:00PM (EST)

Long-lasting, alcohol-free protection from germs

FOAMING HAND SANITIZER

KILLS 99.99% OF GERMS

Formulated and enhanced with **Zetrisil**®

FAST ACTING 15 SEC FORMULA

WITH SOOTHING ALOE VERA

FORMULATED IN THE USA

MADE IN CHINA

Distributed by: ESC Brands, LLC. 1060 Blue Prince Road Bluefield, WV 24701 www.germ-free24.com

Packaging



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GERM FREE 24 HAND SANITIZER								
-	nzalkonium cl		JANIIZER					
Ρ	roduct Info	rmation						
Ρ	Product Type HUMAN OTC DRUG Item Code (Source) NDC						NDC:7	1884-006
	oute of Administration TOPICAL							
Active Ingredient/Active Moiety								
		Ingre	dient Name			Basis of Str	ength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - BENZALKONIUM UNII: 7N6JUD5X6Y) BENZALKONIUM CHLORIDE							1.3 mg in 1 mL	
Inactive Ingredients								
Ingredient Name Strength								
ALOE VERA LEAF (UNII: ZY81Z83H0X)								
WATER (UNII: 059QF0KO0R)								
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)								
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)								
LAURETH-4 (UNII: 6HQ855798J)								
POLIHEXANIDE (UNII: 322U039GMF)								
PHENOXYETHANOL (UNII: HIE492ZZ3T) TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)								
					1010070	1		
Packaging								
#	ltem Code	Pa	ckage Description		Mark	eting Start Date	Mark	ceting End Date
1	NDC:71884- 006-01	50 mL in 1 BOT Product	TLE; Type 0: Not a Combir	nation	08/22/2			
2	NDC:71884- 006-07	207 mL in 1 BC Product	OTTLE; Type 0: Not a Comb	ination	08/22/2	018		
3	NDC:71884- 006-08	245 mL in 1 BC Product	OTTLE; Type 0: Not a Comb	ination	08/22/2	018		

5 NDC:71884- 006-10 1000 mL in 1 BOTTLE; Type 0: Not a Combination Product 08/22/2018 6 NDC:71884- 006-11 3785 mL in 1 PAIL; Type 0: Not a Combination Product 08/22/2018	4	NDC:71884- 006-09	475 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2018	
	5	NDC:71884- 006-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2018	
	6			08/22/2018	

	Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not p	part333A	08/22/2018				

Revised: 2/2022

Enviro Specialty Chemicals Inc