ANVI- benzalkonium chloride liquid Anvi LLC

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.

Anvi

Active Ingredient(s)

Benzalkonium Chloride .13%..... Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use(s)

Hand and skin sanitizer to decrease microbes on the skin. For use when hand washing is not available.

Warnings

For external use only. Keep out of ears, eyes or mouth. In case of contact with eyes, flush with water.

Stop use and ask doctor: If redness or irritation develops and persists for more than 72 hours.

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

Do not use

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Directions

Directions: Dispense an adequate amount into palm to cover hands. Rub hands together to cover skin thoroughly until dry. Supervise young children while using this product.

Other Information: Skin cells may shed naturally or by abrasion prior to four hours, exposing fresh, unprotected skin. Reapply as needed.

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Inactive ingredients

Water. Colloidal Silicon Dioxide

Questions-Information

www.getanvi.com

Principal Display Panel



2 oz (59.14 mL) NDC 80201-100-01

ANVI benzalkonium chloride liquid **Product Information** Product Type HUMAN OTC DRUG Item Code (Source) NDC:80201-200 TOPICAL Route of Administration Active Ingredient/Active Moiety **Ingredient Name Basis of Strength** Strength BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -BENZALKONIUM 0.13 g UNII:7N6JUD5X6Y) **CHLORIDE** in 100 mL

Inactive Ingredients					
Ingredient Name	Strength				
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	0.2 g in 100 mL				
WATER (UNII: 059QF0KO0R)	99.67 g in 100 mL				

Product Characteristics					
Color	Score				
Shape	Size				
Flavor	Imprint Code				
Contains					

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80201-200- 01	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/26/2020		
2	NDC:80201-200- 02	108 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
3	NDC:80201-200- 04	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
4	NDC:80201-200- 03	167 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
5	NDC:80201-200- 05	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
6	NDC:80201-200- 06	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
7	NDC:80201-200- 07	3785 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		
8	NDC:80201-200- 09	208173 mL in 1 DRUM; Type 0: Not a Combination Product	08/26/2020		
9	NDC:80201-200- 10	1040864 mL in 1 CONTAINER; Type 0: Not a Combination Product	08/26/2020		
10	NDC:80201-200- 08	18925 mL in 1 PAIL; Type 0: Not a Combination Product	08/26/2020		
11	NDC:80201-200- 21	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/26/2020		

DRUG FACTS

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Inactive Ingredients: Water, Silicon Dioxide

Manufactured for: ANVI, LLC 640 International Parkway Richardson, TX 75081 KILLS 99.99% OF GERMS



Hand Sanitizer

A SOOTHING PROTECTANT WITH UP TO 500 SPRAYS

ANVI's specialized antiseptic formula sanitizes and protects hands against germs and harmful pathogens for up to 4 hours after applying. Perfect for everyday use and safer than alcohol based sanitizers.

- Moisturize and sanitize at the same time
- Forms a protective barrier on hands
- Sting-free on cuts and scrapes
- Non-flammable

Does not contain any dyes or fragrances.

ALCOHOL FREE

MADE IN THE USA



3 7 6 2 8 2 5 6 1 0 2

2.0 fl oz (59.14 mL)

NDC 80201-200-01

Marketing Information

Marketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateOTC monograph not finalpart333A08/26/2020

Labeler - Anvi LLC (117619678)

Registrant - Anvi LLC (117619678)

Establishment Name Address ID/FEI Business Operations Anvi LLC 117619678 manufacture(80201-200)

Revised: 11/2020 Anvi LLC