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

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

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.

Stop Use and ask a doctor

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

Keep out of Reach of Children

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

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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
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	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	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/26/2020	

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