PURIFI HAND SANITIZER- benzalkonium chloride solution
Purifi LLC

Purifi Moisturizing Hand Sanitizer

INSTANTLY KILLS 99.99% OF TESTED GERMS

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

Hand sanitizing to help reduce bacteria on the skin

Warnings

For external use only.

When using this product avoid contact with the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness appears and lasts

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

Directions

Apply liberally to hands and gently rub until dry. Reapply every 6 hours.

Other information

Store above 32°F (0°C) and below 100°F (38.8°C)

Inactive ingredients

purified water, glycerin, aloe barbadensis leaf juice

Product label

PURIFI HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80659-301
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
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80659-301-01	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2020	
2	NDC:80659-301-02	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2020	
3	NDC:80659-301-03	3800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2020	

4	NDC:80659-301-04	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2020	
5	NDC:80659-301-05	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	12/10/2020	

Labeler - Purifi LLC (117568842)

Revised: 1/2024

Purifi LLC