PURELL PROFESSIONAL HEALTHY SP 0.5PCT BAK ANTIMICROBIAL FOAM-benzalkonium chloride liquid
GOJO Industries, 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.

PURELL Professional HEALTHY SOAP™ 0.5% BAK Antimicrobial Foam

Active ingredient

Benzalkonium Chloride 0.5%

Purpose

Antimicrobial

Uses

• Handwash to help decrease bacteria on the skin

Warnings

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

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

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

Directions

• Wet hands

• Apply a small amount of product and work into a lather

• Rinse well and dry hands completely

Inactive Ingredients

Water (Aqua), Propanediol, Glycerin, Cocamidopropyl Betaine, PEG-80 Sorbitan Laurate, Citric Acid, Ethylhexylglycerin, Lauramine Oxide, Polyquaternium-10, Trisodium Ethylenediamine Disuccinate, Fragrance (Parfum), Phenoxyethanol

PURELL PROFESSIONAL HEALTHY SP 0.5PCT BAK ANTIMICROBIAL FOAM

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-685
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	0.005 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	

PROPANEDIOL (UNII: 5965N8W85T)	
Glycerin (UNII: PDC6A3C0OX)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
PEG-80 Sorbitan Laurate (UNII: 239B50Y732)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Lauramine Oxide (UNII: 4F6FC4MI8W)	
POLYQUATERNIUM-10 (10000 MPA.S AT 2%) (UNII: PI1STR9QYH)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-685-23	222 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
2	NDC:21749-685-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	
3	NDC:21749-685-90	1250 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/02/2018	
4	NDC:21749-685-20	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/15/2017	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-685)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-685) , pack(21749-685)

Revised: 4/2018 GOJO Industries, Inc.