TOPCARE ANTIBACTERIAL WHITE TEA FOAMING HAND- benzalkonium chloride soap
Abaco Partners LLC DBA Surefil

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.

TopCare ® Antibacterial White Tea Foaming Hand Soap

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

for handwashing, reduces germs on the skin

Warnings

For external use only–hands only

When using this product

- Avoid contact with eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if

- irritation and redness develops.

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

Directions

- Pump into to dry hands, vigorously work into a lather and rinse thoroughly.

Inactive ingredients

Purified Water, Lauramine Oxide, Glycerin, Cocamidopropyl Betaine, Isostearamidopropyl Ethyldimonium Ethosulfate, Fragrance, DMDM Hydantoin, Tetrasodium EDTA, Hydroxypropyl Methyl Cellulose, Chlorhexidine Gluconate, Zinc Sulfate Monohydrate, Citric Acid, FD&C Red #33 (CI17200), FD&C Blue #1 (CI42090)

Other information

store at 20°C - 25°C (68°F - 77°F)

Questions?

Call 1-888-423-0139

DISTRIBUTED BY TOPCO ASSOCIATES LLC
ELK GROVE VILLAGE, IL 60007

PRINCIPAL DISPLAY PANEL - 221 mL Bottle Label

TopCare ®

ANTIBACTERIAL
foaming
hand soap

gentle enough
for every day

WHITE TEA

7.5 FL OZ (221 mL)

TOPCARE ANTIBACTERIAL WHITE TEA FOAMING HAND

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20890-0120
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	13 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
ISOSTEARAMIDOPROPYL ETHYLDIMONIUM ETHOSULFATE (UNII: U059JNZ17L)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
EDETATE SODIUM (UNII: MP1J8420LU)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
ZINC SULFATE MONOHYDRATE (UNII: PTX099XSF1)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20890-0120-1	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/01/2016	

Labeler - Abaco Partners LLC DBA Surefil (964809417)

Establishment

Name	Address	ID/FEI	Business Operations
Abaco Partners LLC DBA Surefil		964809417	manufacture(20890-0120)