SIGNATURE CARE ANTIBACTERIAL FOAMING MOISTURIZING- triclosan soap
SAFEWAY 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.

DRUG FACTS

Active ingredient

Triclosan 0.46%

Purpose

Antibacterial

Uses

for handwashing to decrease bacteria on the skin.

Warnings

For external use only.

When using this product

avoid contact with eyes. If eye contact occurs, rinse thoroughly with water.

Stop using this product and ask a doctor if

irritaiton or redness develops

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- Pump onto dry hands.
- Lather vigorously for at least 15 seconds.
- Rinse and dry thoroughly.

Other information

store at room temperature

Inactive ingredients

Water (Aqua), Sodium Xylenesulfonate, Dipropylene Glycol, Glycerin, Ammonium Lauryl Sulfate, Sodium PCA, Cocamidopropyl Betaine, Polyquaternium-10, Fragrance (Parfum), Disodium Phosphate, Cetyl Alcohol, Aloe Barbadensis Leaf Juice, Citric Acid, Methylparaben, Propylparaben, Red 4 (CI 14700), Yellow 5 (CI 19140).

Questions or comments?

1-888-723-3929

Label Copy

SIGNATURE CARE ANTIBACTERIAL FOAMING MOISTURIZING

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-177
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	4.6 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
SODIUM PHOSPHATE, DIBASIC ANHYDROUS (UNII: 22ADO53M6F)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-177-07	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/11/2016	

Labeler - SAFEWAY INC (009137209)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(21130-177)

Revised: 2/2016 SAFEWAY INC