DAYLOGIC FOAMING SANITIZER ALOE- benzalkonium chlolride liquid
RITE AID CORPORATION

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

Benzalkonium Chloride 0.1%

Purpose

Antiseptic

Uses

to help decrease bacteria on the skin.

Warnings

For external use only

When using this product

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

Stop use and ask a doctor if

irritation or redness develops and lasts.

Keep out of reach of children.

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

Directions

- Pump enough product on your palm to thoroughly cover your hands, rub together until dry.
- Children under 6 years should be supervised when using this product.

Inactive ingredients

Water (Aqua), Polysorbate 20, Ethylhexyl Methoxycinnamate, Buty Methoxydibenzoylmethane, Ethylhexyl Salicyalte, PPG-26-Buteth-26, PEG-40 Hydrogenated Castor Oil, Aloe Barbadensis Leaf Juice, Fragrance (Parfum), Camellia Sinensis Leaf Extract, DMDM Hydantoin, Sodium Hydroxide, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1 (CI 42090), Yellow 5 (CI 19140).

Label copy

DAYLOGIC FOAMING SANITIZER ALOE

benzalkonium chlolride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-2440
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
OCTINOXATE (UNII: 4Y5P7MUD51)	
AVOBENZONE (UNII: G63QQF2NOX)	
OCTISALATE (UNII: 4X49Y0596W)	
PPG-26-BUTETH-26 (UNII: 2II1K6TZ4P)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-2440-8	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/20/2020	

Labeler - RITE AID CORPORATION (014578892)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(11822-2440)

Revised: 3/2020 RITE AID CORPORATION