

**EQUATE ANTIBACTERIAL FOAMING HAND SPRING SHOWERS- benzalkonium
chloride liquid
WAL-MART STORE 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

BENZALKONIUM CHLORIDE 0.13%

PURPOSE

ANTIBACTERIAL

USES

HELPS REDUCE 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 USING THIS PRODUCT AND ASK A DOCTOR IF
IRRITATION AND 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

USE ONLY TO REFILL A FOAMING HAND SOAP PUMP BOTTLE. PUMP ONTO DRY HANDS,
WORK INTO A RICH FOAMY LATHER AND RINSE AND DRY THOROUGHLY

OTHER INFORMATION

STORE AT ROOM TEMPERATURE

INACTIVE INGREDIENTS

WATER (AQUA), COCAMIDOPROPYL BETAINE, GLYCERIN, DECYL GLUCOSIDE, ALOE
BARBADENSIS LEAF JUICE, CAMELLIA SINENSIS LEAF EXTRACT, FRAGRANCE
(PARFUM), POLYQUATERNIUM-7, XANTHAN GUM, TETRASODIUM EDTA, POLYSORBATE
20, SODIUM CITRATE, CITRIC ACID, METHYLCHLOROISOTHIAZOLINONE,
METHYLISOTHIAZOLINONE, BLUE 1 (CI 42090), EXT. VIOLET 2 (CI 60730)

QUESTIONS?

1-888-287-1915

LABEL COPY



EQUATE ANTIBACTERIAL FOAMING HAND SPRING SHOWERS

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)			BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)				
GLYCERIN (UNII: PDC6A3C0OX)				
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)				
XANTHAN GUM (UNII: TTV12P4NEE)				
EDETATE SODIUM (UNII: MP1J8420LU)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)				
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-014-32	946 mL in 1 BOTTLE, PLASTIC		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333E	08/17/2014	

Labeler - WAL-MART STORE INC (051957769)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(49035-014)