CVS PHARMACY COCONUT WATER- benzalkonium chloride liquid CVS PHARMACY

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

FOR WASHING TO DECREASE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE 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

APPLY ONTO DRY HANDS. WORK INTO A LATHER, RINSE AND DRY THOROUGHLY

OTHER INFORMATION

STORE AT ROOM TEMPERATURE

INACTIVE INGREDIENTS

ALOE BARBADENSIS LEAF JUICE, BAMBUSA VULGARIS LEAF EXTRACT, BLUE 1 (CI 42090), CITRIC ACID, COCAMIDOPROPYL BETAINE, COCOS NUCIFERA (COCONUT) EXTRACT, DECYL GLUCOSIDE, FRAGRANCE (PARFUM), GLYCERIN, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, POLYQUATERNIUM-7, POLYSORBATE 20, RED 33 (CI 17200), SODIUM CITRATE, TETRASODIUM EDTA, WATER (AQUA), XANTHAN GUM

QUESTIONS OR COMMENTS?

1-800-746-7287

LABEL COPY





CVS PHARMACY COCONUT WATER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-842

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
ı	Ingredient Name	Basis of Strength	Strength	
	BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
BAMBUSA VULGARIS LEAF (UNII: EMY54R518C)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
CITRIC ACID ACETATE (UNII: DSO 12WL7AU)		

COCAMIDO PRO PYL BETAINE (UNII: 50 CF3O 11KX)	
COCONUT OIL (UNII: Q9L0O73W7L)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE SO DIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:59779-842-32	946 mL in 1 POUCH		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	0 1/26/20 14		

Labeler - CVS PHARMACY (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(59779-842)	

Revised: 1/2014 CVS PHARMACY