# CVS PHARMACY CLEAN ANTIBACTERIAL- triclos an 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.

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#### **DRUG FACTS**

#### **ACTIVE INGREDIENT**

TRICLOSAN 0.46%

#### **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 THOROUGHLY WITH WATER.

#### STOP USING THIS PRODUCT AND ASK A DOCTOR

IF IRRITATION OR REDNESS DEVELOPS AND LASTS MORE THAN 7 DAYS.

#### KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

#### **DIRECTIONS**

APPLY TO DRY HANDS, WORK INTO A RICH FOAMY LATHER AND RINSE THOROUGHLY.

### **INACTIVE INGREDIENTS:**

WATER (AQUA), SODIUM XYLENESULFONATE, DIPROPYLENE GLYCOL, GLYCERIN, SODIUM PCA, AMMONIUM LAURYL SULFATE, 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).

#### LABEL COPY





### CVS PHARMACY CLEAN ANTIBACTERIAL

triclosan liquid

Pro	duct	Info	rmati	nη
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:59779-179

Route of Administration TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
TRICLOSAN (UNII: 4NM5039 Y5X) (TRICLOSAN - UNII:4NM5039 Y5X)
TRICLOSAN

0.46 mg in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
SODIUM XYLENESULFONATE (UNII: G4LZF950 UR)			
DIPROPYLENE GLYCOL (UNII: E107L85C40)			
GLYCERIN (UNII: PDC6 A3C0 OX)			
SODIUM PYRROLIDONE CARBO XYLATE (UNII: 469 OTG 57A2)			
AMMO NIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)			
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)			
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)			
SODIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)			

CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
METHYLPARABEN (UNII: A218 C7H19 T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C RED NO. 4 (UNII: X3W0 AMIJLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:59779-179-08	221 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	07/02/2012	

## Labeler - CVS PHARMACY (062312574)

## Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
APOLLO HEALTH AND BEAUTY CARE		20 19 0 12 0 9	manufacture	

Revised: 7/2012 CVS PHARMACY