# ANTIBACTERIAL FOAMING ORIGINAL- triclos an liquid MEIJER DISTRIBUTION 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.

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#### **DRUG FACTS**

#### **ACTIVE INGREDIENT**

TRICLOSAN 0.46%

#### **PURPOSE**

ANTIBACTERIAL

#### **USES**

HELPS REDUCES 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 DOCTOR

IF IRRITATION AND REDNESS DEVELOP AND LAST.

#### KEEP OUT OF REACH OF CHILDREN

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

#### **DIRECTIONS**

PUMP ONTO DRY HANDS. WORK INTO A LATHER AND RINSE THOROUGHLY.

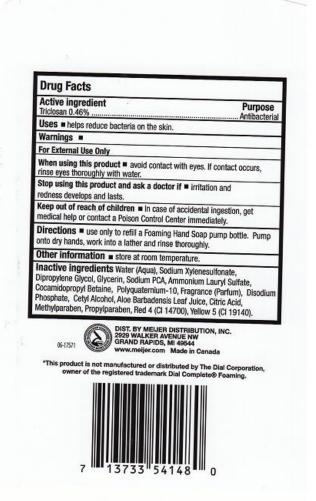
## OTHER INFORMATION

STORE AT ROOM TEMPERATURE.

## **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).





# ANTIBACTERIAL FOAMING ORIGINAL

triclosan liquid

### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-177

Route of Administration TOPICAL

#### Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039	Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.46 mL 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 CARBOXYLATE (UNII: 469OTG57A2)	
AMMO NIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3011KX)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
SODIUM PHO SPHATE (UNII: SE337SVY37)	
C12-16 ALCOHOLS (UNII: S4827SZE3L)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-177-32	936 mL in 1 BOTTLE, PUMP		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	11/18/2011		

# Labeler - MEUER DISTRIBUTION INC (006959555)

# Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 11/2011 MEIJER DISTRIBUTION INC