



# HILL COUNTRY ESSENTIALS ANTIBACTERIAL

Dosage form: liquid

Ingredients: TRICLOSAN 1.5mg in 1mL

Labeler: H E B

NDC Code: 37808-722

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

TRICLOSAN 0.15%

### PURPOSE

ANTIBACTERIAL

### USES

FOR HAND WASHING TO DECREASE BACTERIA ON THE SKIN

### WARNINGS

FOR EXTERNAL USE ONLY

### WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, FLUSH WITH WATER

### STOP USING THIS PRODUCT 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

USE AS A REFILL FOR DISPENSING BOTTLE. FROM DISPENSING BOTTLE, APPLY A SMALL AMOUNT ONTO WET HANDS. WORK INTO A LATHER AND RINSE THOROUGHLY

### OTHER INFORMATION

STORE AT ROOM TEMPERATURE

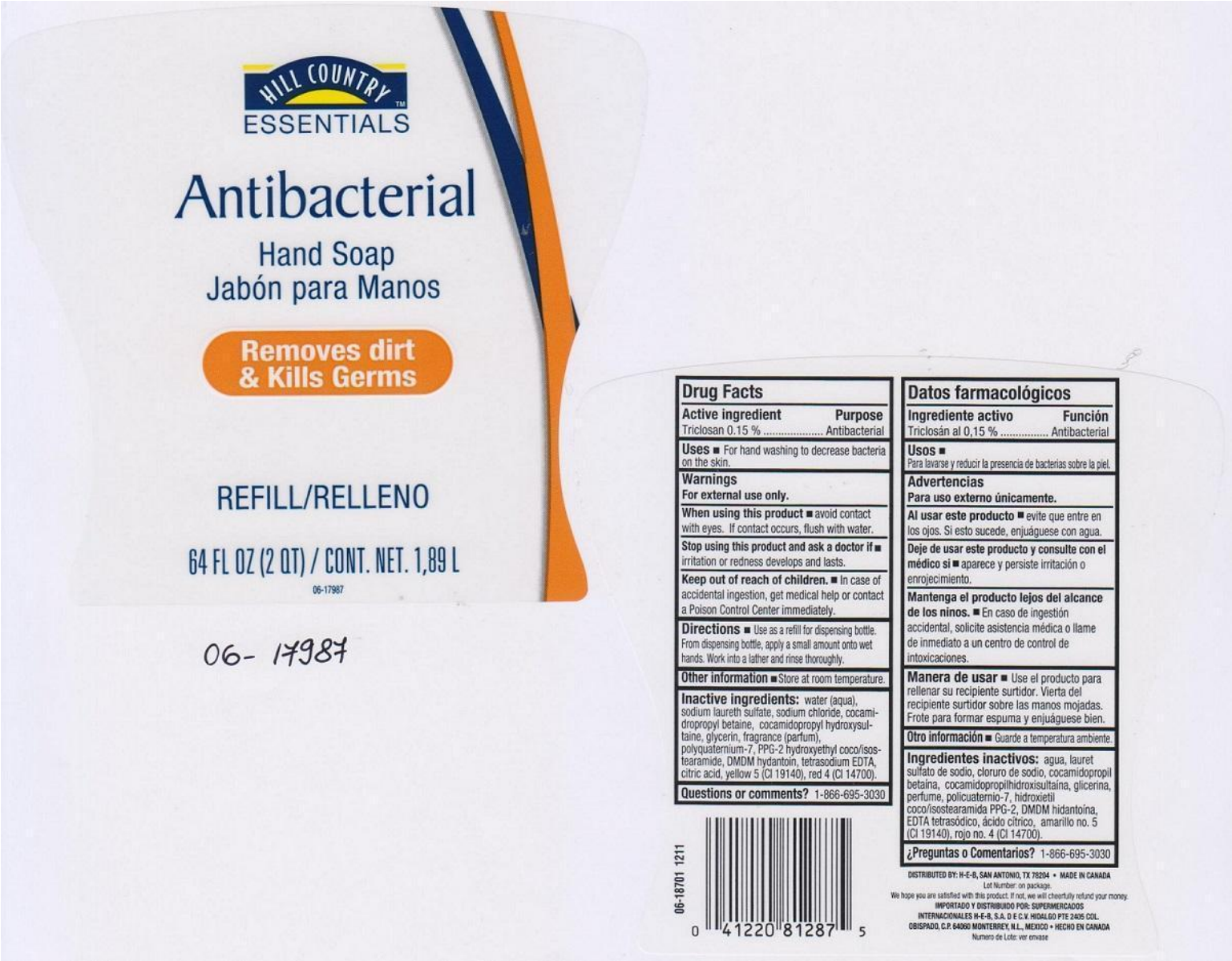
### INACTIVE INGREDIENTS

WATER (AQUA), SODIUM LAURETH SULFATE, SODIUM CHLORIDE, COCAMIDOPROPYL BETAINE, COCAMIDOPROPYL HYDROXYSULTAINE, GLYCERIN, FRAGRANCE (PARFUM), POLYQUATERNIUM-7, PPG-2 HYDROXYETHYL COCO/ISOSTEARAMIDE, DMDM HYDANTOIN, TETRASODIUM EDTA, CITRIC ACID, YELLOW 5 (CI 19140), RED 4 (CI 14700)

### QUESTIONS OR COMMENTS?

1-866-695-3030

### LABEL COPY



HILL COUNTRY ESSENTIALS ANTIBACTERIAL

triclosan liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-722
Route of Administration	TOPICAL	DEA Schedule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (TRICLOSAN)	TRICLOSAN	1.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER	
SODIUM LAURETH SULFATE	
SODIUM CHLORIDE	
COCAMIDOPROPYL BETAINE	
COCAMIDOPROPYL HYDROXYSULTAINE	
GLYCERIN	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)	
PPG-2 HYDROXYETHYL COCO/ISOSTEARAMIDE	
DMDM HYDANTOIN	
EDETATE SODIUM	
CITRIC ACID MONOHYDRATE	
FD&C YELLOW NO. 5	
FD&C RED NO. 4	

Packaging		
#	Item Code	Package Description
1	NDC:37808-722-64	1890 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	03/31/2015	

Labeler - H E B (007924756)
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Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)
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Establishment			
Name	Address	ID/FEI	Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(37808-722)

Revised: 04/2015

H E B

← See all HILL COUNTRY ESSENTIALS ANTIBACTERIAL brands

Disclaimer: Every effort has been made to ensure that the information provided here is accurate, up-to-date and complete, but no guarantee is made to that effect. Drug information contained herein may be time sensitive. This information has been compiled for use by healthcare practitioners and consumers in the United States. The absence of a warning for a given drug or combination thereof in no way should be construed to indicate that the drug or combination is safe, effective or appropriate for any given patient. If you have questions about the substances you are taking, check with your doctor, nurse or pharmacist.