WINNING HANDS FOAMING ANTIBACTERIAL- triclosan soap Betco Corporation, Ltd.

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.

Winning Hands Foaming Antibacterial

Winning Hands Foaming Antibacterial

Active Ingredient

□ □Triclosan 0.30%

Winning Hands Foaming Antibacterial

Uses

- Use in a variet of public facilities including daycare centers, hospitals, nursing homes, physicians offices.

Winning Hands Foaming Antibacterial

Warnings

- For external use only.
- Avoid contact with eyes.
- Children under the age of 6 should be supervised by an adult when using this product.
- Discontinue use is irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
- KEEP OUT OF REACH OF CHILDREN.

Winning Hands Foaming Antibacterial

Directions

- Read the entire label before using this product.
- Dispense 0.8 mL of product onto wet palm.
- Rub hands together to distribute product, then rinse hands with clean.

Winning Hands Foaming Antibacterial

Inactive Ingredients

UWater, Potasium cocoate (contains coconut), Propylene Glycol, Glycerine, DMDM Hydantoin, Fragrance, FD&C Yellow #5, Aloe Barbadensis Leaf Juice. FD&C Red #40.

Winning Hands Foaming Antibacterial

Questions or Comments?Phone: (800) 777-9343

MDS information: (800) 891-4965

Winning Hands Foaming Antibacterial

Purpose

Antibacterial

Winning Hands Foaming Antibacterial

KEEP OUT OF REACH OF CHILDREN

Winning Hands Foaming Antibacterial Winning Hands Foaming Antibacterial 75155-00

Foaming Skin Cleanser with Triclosan
NET CONTENTS: 55 gallons (U.S./E.U.) 208 L





WINNING HANDS FOAMING ANTIBACTERIAL

tricolsan soap

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:65601-751
Route of Administration	Topical	DEA Schedule	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
TRICLOSAN (TRICLOSAN)	TRICLOSAN	3.0 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER	
POTASSIUM COCOATE	
PROPYLENE GLYCOL	
GLYCERIN	
EDETATE SO DIUM	
DMDM HYDANTO IN	
FD&C YELLOW NO.5	
ALOE VERA Leaf	
FD&C RED NO. 40	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65601-751-54	1000 mL in 1 BOTTLE, PLASTIC		
2	NDC:65601-751-55	208000 mL in 1 DRUM		
3	NDC:65601-751-04	3780 mL in 1 JUG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/12/2012	

Labeler - Betco Corporation, Ltd. (005050158)

Registrant - Betco corporation, Ltd. (005050158)

Establishment				
Name	Address	ID/FEI	Business Operations	
Betco Corpo, Ltd.		005050158	manufacture(65601-751)	

Revised: 12/2012 Betco Corporation, Ltd.