ANTIBACTERIAL FOAMSOAP- benzalkonium chloride soap
Ferguson Enterprises

WC10007 - Antibacterial Foam Soap

Active ingredient

Benzalkonium Chloride 0.13% w/w

Uses

Antibacterial skin cleanser

Effective in destroying harful bacteria to prevent antibacterial cleansing

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 rash occurs.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Read entire label before using this product.

Apply 5 milliliters (teaspoonful) or palmful to hands and forearms.

Scrub thoroughly for 20 seconds and rinse with clean water.

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Citric Acid, FD&C Red 40, FD&C Yellow 5, Fragrance, Glycerin, Laurmine Oxide, Methylchloroisothiazolinone, Methylisothiazolinone, Tetrasodium Glutamate Diacetate, Water

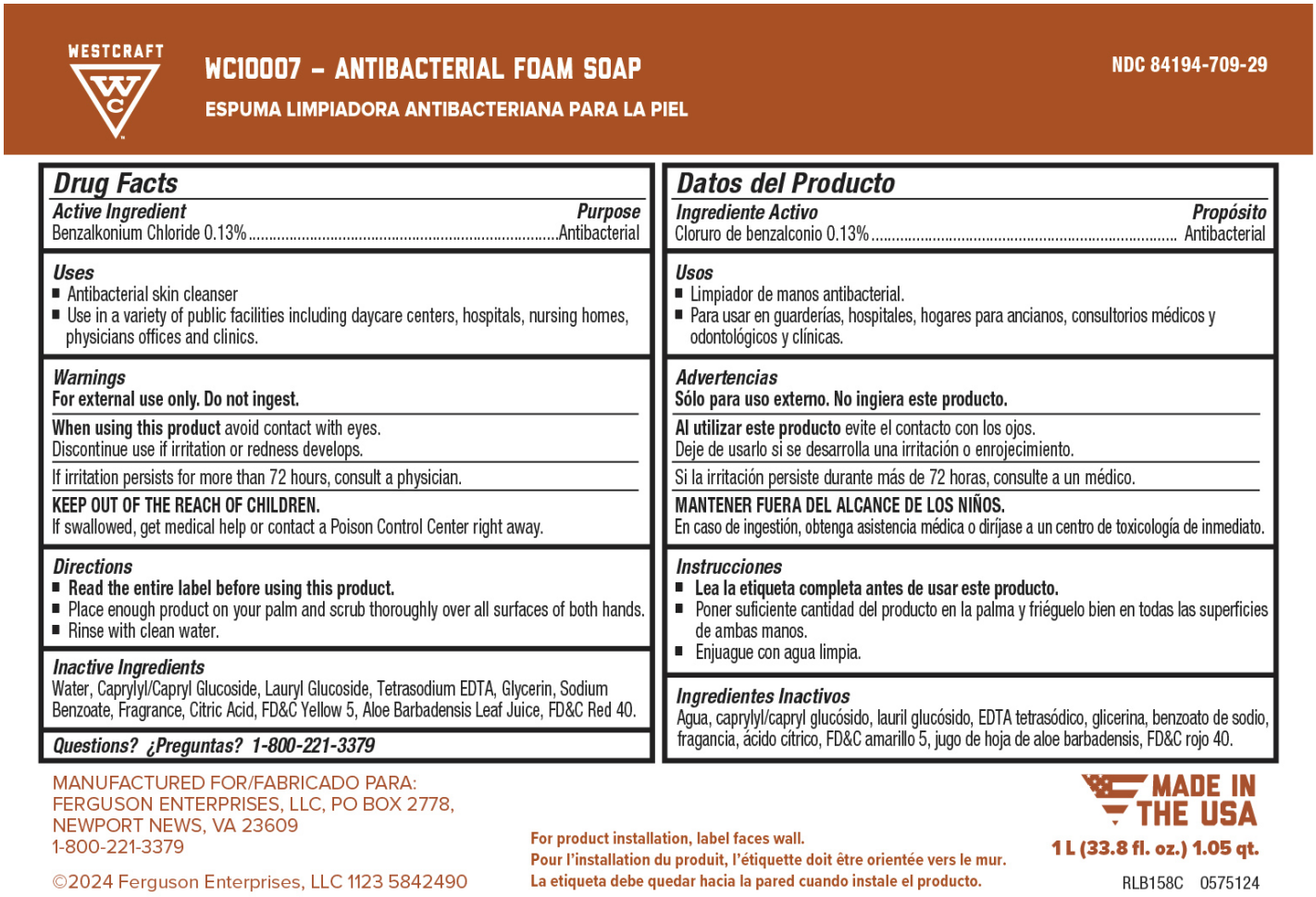
Purpose

Antibacterial

Keep out of reach of children

Keep out of the reach of children

Label



ANTIBACTERIAL FOAMSOAP

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84194-751
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

FD&C RED NO. 40 (UNII: WZB9127XOA)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84194-751-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	05/21/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/21/2024	

Labeler - Ferguson Enterprises (008955171)

Registrant - Betco Corporation (005050158)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation		005050158	manufacture(84194-751) , label(84194-751)

Revised: 6/2024 Ferguson Enterprises