

ANTIBACTERIAL FOAM SOAP- benzalkonium chloride soap
Ferguson Enterprises

WC10007 - Antibacterial Foam Soap

Active ingredient

Benzalkonium Chloride 0.13% w/w

Uses

Antibacterial skin cleanser

Effective in destroying harmful 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, Lauramine Oxide, Methylchloroisothiazolinone, Methylisothiazolinone, Tetrasodium Glutamate Diacetate, Water

Purpose

Antibacterial

Keep out of reach of children

Keep out of the reach of children

Label



WC10007 – ANTIBACTERIAL FOAM SOAP
ESPUMA LIMPIADORA ANTIBACTERIANA PARA LA PIEL

NDC 84194-709-29

Drug Facts		Datos del Producto	
Active Ingredient	Purpose	Ingrediente Activo	Propósito
Benzalkonium Chloride 0.13%	Antibacterial	Cloruro de benzalconio 0.13%	Antibacterial
Uses		Usos	
<ul style="list-style-type: none"> Antibacterial skin cleanser Use in a variety of public facilities including daycare centers, hospitals, nursing homes, physicians offices and clinics. 		<ul style="list-style-type: none"> Limpiador de manos antibacterial. Para usar en guarderías, hospitales, hogares para ancianos, consultorios médicos y odontológicos y clínicas. 	
Warnings		Advertencias	
<p>For external use only. Do not ingest.</p> <p>When using this product avoid contact with eyes. Discontinue use if irritation or redness develops. If irritation persists for more than 72 hours, consult a physician.</p> <p>KEEP OUT OF THE REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control Center right away.</p>		<p>Sólo para uso externo. No ingiera este producto.</p> <p>Al utilizar este producto evite el contacto con los ojos. Deje de usarlo si se desarrolla una irritación o enrojecimiento. Si la irritación persiste durante más de 72 horas, consulte a un médico.</p> <p>MANTENER FUERA DEL ALCANCE DE LOS NIÑOS. En caso de ingestión, obtenga asistencia médica o diríjase a un centro de toxicología de inmediato.</p>	
Directions		Instrucciones	
<ul style="list-style-type: none"> Read the entire label before using this product. Place enough product on your palm and scrub thoroughly over all surfaces of both hands. Rinse with clean water. 		<ul style="list-style-type: none"> Lea la etiqueta completa antes de usar este producto. Poner suficiente cantidad del producto en la palma y fríguelo bien en todas las superficies de ambas manos. Enjuague con agua limpia. 	
Inactive Ingredients		Ingredientes Inactivos	
Water, Caprylyl/Capryl Glucoside, Lauryl Glucoside, Tetrasodium EDTA, Glycerin, Sodium Benzoate, Fragrance, Citric Acid, FD&C Yellow 5, Aloe Barbadensis Leaf Juice, FD&C Red 40.		Agua, caprylyl/capryl glucósido, lauril glucósido, EDTA tetrasódico, glicerina, benzoato de sodio, fragancia, ácido cítrico, FD&C amarillo 5, jugo de hoja de aloe barbadensis, FD&C rojo 40.	
Questions? ¿Preguntas? 1-800-221-3379			

MANUFACTURED FOR/FABRICADO PARA:
 FERGUSON ENTERPRISES, LLC, PO BOX 2778,
 NEWPORT NEWS, VA 23609
 1-800-221-3379

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For product installation, label faces wall.
 Pour l'installation du produit, l'étiquette doit être orientée vers le mur.
 La etiqueta debe quedar hacia la pared cuando instale el producto.



1 L (33.8 fl. oz.) 1.05 qt.

RLB158C 0575124

ANTIBACTERIAL FOAM SOAP

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