

**BRIGHTON PROFESSIONAL PREMIUM ANTIBACTERIAL FOAM SO AP-
chloroxylenol liquid
Staples**

BRIGHTON PROFESSIONAL™ Premium Antibacterial Foam Soap

Active ingredient

Chloroxylenol 0.3%

Purpose

Antimicrobial

Uses

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

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

Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

Inactive ingredients

Water (Aqua), Alcohol, Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Xylenesulfonate, Cocamide MEA, Glycerin, Isopropyl Alcohol, Lactic Acid, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Ammonium Sulfate, Fragrance (Parfum), Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)

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Premium Antibacterial
Foam Soap
Jabón espumoso
antibacterial de alta calidad

Manufactured for / Fabricado para:
Staples Contract & Commercial, Inc.
500 Staples Drive, Frammingham, MA 01702 USA, 1-888-322-0912
Made in USA / Hecho en los E.E.U.U. ©2017 Staples, Inc.

Not for Individual Sale

18503

1200 mL (40.5 US/ÉU FL OZ)

1790-L00-HG6-F

Drug Facts

Active ingredient	Purpose
Chloroxylenol 0.3%	Antimicrobial

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In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

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

Directions

- Wet Hands
- Apply product and thoroughly cover hand with lather
- Rinse well and dry hands completely

Inactive ingredients Water (Aqua), Alcohol, Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Xylenesulfonate, Cocamide MEA, Glycerin, Isopropyl Alcohol, Lactic Acid, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Ammonium Sulfate, Fragrance (Parfum), Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)

Datos Farmacológicos

Ingrediente activo	Propósito
Cloroxileno 0,3%	Antimicrobiano

Usos

- Lavado de manos empleado para disminuir la cantidad de bacterias en la piel
- Recomendado para uso reiterado

Advertencias

Sólo para uso externo

Al utilizar este producto, evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.

Dejar de usar el producto y consultar a un médico si aparece y persiste una irritación o erupción cutánea

Mantener fuera del alcance de los niños. En caso de ingestión, de inmediato acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.

Modo de uso

- Moje las manos
- Aplique el producto hasta cubrir completamente sus manos con la espuma
- Enjuague bien y seque las manos completamente

Ingredientes inactivos Agua, Alcohol etílico, Lauret sulfato de amonio, Lauril sulfato de amonio, Propilenglicol, Xilenosulfonato de amonio, MEA cocamida, Glicerina, Alcohol isopropílico, Acido láctico, Palmitato de retinilo, Aceite de Simmondsia Chinensis, EDTA tetrasódico, Acetato de tocoferilo, Aceite de maíz (Zea Mays), Sulfato de amonio, Fragancia, Metilcloroisotiazolinona, Metilisotiazolinona, Rojo 4, Amarillo 6

BRIGHTON PROFESSIONAL PREMIUM ANTIBACTERIAL FOAM SOAP
chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49514-081
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.003 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	
AMMONIUM LAURETH-2 SULFATE (UNII: 698O4Z48G6)	
AMMONIUM LAURYL SULFATE (UNII: Q7A02R1M0B)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMMONIUM XYLENESULFONATE (UNII: 4FZY6L6XCM)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LACTIC ACID (UNII: 33X04XA5AT)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
JOJOBA OIL (UNII: 724GKU717M)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CORN OIL (UNII: 8470G57WFM)	
AMMONIUM SULFATE (UNII: SU46BAM238)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49514-081-40	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/30/2007	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/30/2007	

Labeler - Staples (151064821)