

FACILIPRO- benzalkonium chloride solution
Ecolab Inc.

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

Benzalkonium Chloride 0.55%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings

For external use only

Do not use

- In eyes

When using this product

- if in eyes, rinse promptly and thoroughly with water.
- discontinue use if irritation and redness develop.

Stop use and ask a doctor if skin irritation or redness occurs for more than 72 hours

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

Directions

- wet skin and spread a small amount on hands and forearms
- scrub well, rinse thoroughly and dry

Other information

- for additional information, see Safety Data Sheet (SDS)
- **EMERGENCY HEALTH INFORMATION:** 1 800 328 0026. If located outside the United States and Canada, call collect 1 651 222 5352 (number is in the US).

Inactive ingredients water, hexylene glycol, cocamine oxide, PEG-180, glycerin, cocamidopropyl PG-dimonium chloride phosphate, methyl gluceth-20, phenoxyethanol, myristamide DIPA, caprylic/capric glycerides, citric acid, hydroxypropyl guar hydroxypypropyltrimonium chloride, PEG-12 dimethicone, ethanol, tocopheryl acetate, fragrance, potassium hydroxide

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Questions? call **1-800-35-CLEAN (352-5326)**

Principal Display Panel and representative label

ECOLAB

6100960

FACILIPRO

Premium Antibacterial

Foam Hand Soap

1250 mL (42.3 US FL OZ)

Active ingredient: Benzalkonium Chloride 0.55%

761097/5402/0321

Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA ·

tel: 1 800 35 CLEAN (352 5326)

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www.gofacilipro.com

Datos sobre la droga (continuado)

Cuando use este producto

- en los ojos, láveselos rápida y completamente con abundante agua
- descontinúe su uso si ocurre irritación o enrojecimiento

Deje de usarlo y vea a un doctor si la irritación en la piel y rosado persiste dura más de 72 horas

Manténgase fuera del alcance de los niños. Si se impiere, obtenga ayuda médica o póngase en contacto con un centro de envenenamiento inmediatamente.

Instrucciones

- moje la piel y extienda pequeña cantidad en sus manos y antebrazos
- fróteselos bien, enjuáguelos completamente y luego séquese los

Datos sobre la droga (continuado)

Información adicional

- para obtener información adicional, consulte la de Ficha de Seguridad
- INFORMACIÓN DE EMERGENCIA SOBRE SALUD: 1 800 328 0026. Si está fuera de los Estados Unidos y Canadá, llame a cobro revertido al 1 651 222 5352 (número en los EE.UU.).

Ingredientes inactivos agua, glicol de hexileno, óxido de cocamina, PEG-180, glicerina, cocamidopropil PG-dimonio cloruro fosfato, metilgluceth-20, fenoxietanol, DIPA miristamida, glicéridos caprílico / capríco, ácido cítrico, cloruro de guar hidroxipropil hidroxipropiltrimonio, PEG-12 dimeticona, etanol, acetato de tocoferol, fragancia, hidróxido de potasio

Datos sobre la droga (continuado)

¿Preguntas? llamada 1-800-35-CLEAN (352-5326)

This product may be patented | Este producto puede ser patentado: www.ecolab.com/patents

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tel: 1 800 35 CLEAN (352 5326)
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ECOLAB 6100960

FACILIPRO™

Premium Antibacterial Foam Hand Soap
Premium jabón espuma antibacterial para manos

1250 mL (42.3 US FL OZ)

Active ingredient: Benzalkonium Chloride 0.55%

761097/5402/0321 0 25469 02500 5

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Datos sobre la droga

Ingrediente activo	Propósito
Cloruro de benzalconio 0.55%....	Lavado de manos antiséptico

Usos

- lavado de manos para disminuir las bacterias en la piel

Advertencia
Para uso externo solamente

No lo use

- en los ojos

FACILIPRO

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-565
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	5.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
glycerin (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HYDROXYPROPYL GUAR (2500-4500 MPA.S AT 1%) (UNII: 3A1I7376TC)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
ALCOHOL (UNII: 3K9958V90M)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-565-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/29/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/29/2016	

