

PROCLEAN- benzalkonium chloride solution
Ecolab Inc.

Drug Facts

Active Ingredient

Benzalkonium chloride 0.5%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warning

For external use only

Do not use

- in eyes

When using the product

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

Stop use and ask a physician 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 hands and apply foam
- scrub hands and forearms
- rinse thoroughly and dry

Other information

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information, in USA and Canada, call 1 800 328 0026

Inactive ingredients water (aqua), cocamine oxide, hexylene glycol, PEG-180, glycerin, hydroxyethylcellulose, cocamidopropyl PG-dimonium chloride phosphate, phenoxyethanol, myristamide DIPA, myristamine oxide, methyl gluceth-20, glyceryl

caprylate/caprates, alcohol, PEG-12 dimethicone, citric acid, polyquaternium-7, fragrance, potassium hydroxide, blue 1

Questions? call **1-866-781-8787**

Representative Label and Principal Display Panel

PROCLEAN

ADVANCED

ANTIBACTERIAL

HAND SOAP

Antibacterial liquid hand soap

Active ingredient: Benzalkonium chloride 0.5%

6102017

3.78 L (1 US GAL)

For more ingredient information visit: www.shamrockfoodservice.com/proclean

This product may be patented

For questions or comments, call 1-866-444-7450.

Manufactured by

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

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774470/5400/1018

Distributed by Shamrock Foods

Phoenix, AZ

PROCLEAN

ADVANCED ANTIBACTERIAL HAND SOAP

Antibacterial liquid hand soap
Jabón antibacterial líquido para manos

Active ingredient: Benzalkonium chloride 0.5%



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Datos sobre la droga

Ingrediente activo	Propósito
Cloruro de benzalconio 0.5%	Jabón antiséptico para manos

Usos
 ■ lavado de manos para disminuir las bacterias en la piel

Advertencias
 Para uso externo solamente

No lo use
 ■ en los ojos

Cuando use este producto
 ■ si en los ojos, láveselos rápida y completamente con abundante agua
 ■ si ocurriera irritación rojura, descontinúe su uso

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

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

Instrucciones
 ■ manos húmedas y aplique el producto
 ■ refriéguese las manos y los antebrazos
 ■ enjuáguese completamente y séquese

Información adicional
 ■ para informaciones adicionales, vea la hoja de datos de seguridad
 ■ para información médica de emergencia en los EEUU o Canadá, llame al 1 800 328 0026

Ingredientes inactivos agua, óxido de cocamina, hexilenglicol, PEG-180, glicerina, hidroxietilcelulosa, cloruro de fosfato de cocamidopropil PG dimonio, fenoxietanol, miristamida DIPA, óxido de miristamina, metil glucet-20, gliceril caprilato/caprato, alcohol, PEG-12, dimeticona, ácido cítrico, policuaternio-7, fragancia, hidróxido de potasio, azul 1

¿Preguntas? llame al 1-866-781-8787

For more ingredient information visit | Para obtener más información acerca de los ingredientes, visite:
www.shamrockfoodservice.com/proclean

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

For questions or comments, call 1-866-444-7450.
 Para información sobre productos, llame al 1-866-444-7450.

Manufactured by | Fabricado por
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 774470/5400/1018

Distributed by | Distribuido por
 Shamrock Foods
 Phoenix, AZ



PROCLEAN

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-602
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYETHYL CELLULOSE (4000 MPA.S AT 1%) (UNII: ZYD53NBL45)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
GLYCERYL MONO- AND DICAPRYLOCAPRATE (UNII: U72Q2I8C85)	
ALCOHOL (UNII: 3K9958V90M)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-602-11	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/29/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/29/2018	

Labeler - Ecolab Inc. (006154611)

Revised: 8/2024

Ecolab Inc.