

SWISHER- 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 0.5%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings

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 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 hands and apply foam
- scrub hands and forearms
- rinse thoroughly and dry

Other information

- READ SAFETY DATA SHEET (SDS) BEFORE USING THIS PRODUCT
- 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 (aqua), cocamine oxide, hexylene glycol, PEG-180, glycerin, cocamidopropyl PG-dimonium chloride phosphate, phenoxyethanol, polyquaternium-7, myristamide DIPA, myristamine oxide, citric acid, methyl gluceth-20, glyceryl caprylate/caprato, alcohol, PEG-12 dimethicone, potassium citrate, fragrance, potassium hydroxide, blue 1, methylparaben, propylparaben.

Questions? call **1.866.444.7450**

Principal display panel & representative package

SWISHER

HIGH PERFORMANCE ANTIBACTERIAL HAND SOAP

6101901

750 mL (25 US FL OZ)

Active ingredient:

Benzalkonium chloride 0.5%

772409/5400/0618

For more ingredient information visit: www.ecolab.com/sds

This product may be patented: www.ecolab.com/patents

For more information visit: www.swsh.com

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

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Datos sobre la droga (continuado)

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 la espuma
- refriéguese las manos y los antebrazos
- enjuáguese completamente y seque

Información adicional

- LEA LA FICHA DE DATOS DE SEGURIDAD ANTES DE USAR EL PRODUCTO

Datos sobre la droga (continuado)

■ **INFORMACIÓN DE EMERGENCIA SOBRE SALUD:**
1 800 328 0026. Si está fuera de los Estados Unidos y Canadá, llame a cubo revertido al 1 651 222 5352 (número en los EE.UU.).

Ingredientes inactivos agua, óxido de cocamina, hexilenglicol, PEG-180, glicerina, cocamidopropil PG-dimonio cloruro fosfato, fenoxietanol, policuaternio 7, DIPA miristamida, óxido de miristamina, ácido cítrico, metilgluceth-20, glicéridos caprílico / capríco, alcohol, PEG-12 dimeticona, citrato de potasio, fragancia, azul 1.

¿Preguntas? llame al 1.866.444.7450

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

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

For more information visit: www.swsh.com

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Made in U.S.A. | Hecho en EE.UU.



HIGH PERFORMANCE ANTIBACTERIAL HAND SOAP

Jabón antibacterial avanzada espuma mano

6101901

750 mL (25 US FL OZ)

Active ingredient | Ingrediente activo: Benzalkonium chloride 0.5%



772409/5400/0618

Drug Facts

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Drug Facts (continued)

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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

Advertencia
Para uso externo solamente

No lo use

- en los ojos

Cuando use este producto

- si ocurriera irritación roja, descontinúe su uso
- si en los ojos, láveselos rápida y completamente con abundante agua

SWISHER
benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-591
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)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERYL CAPRYLOCAPRATE (UNII: U72Q2I8C85)	
ALCOHOL (UNII: 3K9958V90M)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-591-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/20/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/20/2018	

