

PURE FORCE- benzalkonium chloride liquid
Ecolab Inc.

Drug Facts

Active Ingredient

Benzalkonium chloride 0.1%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin
- recommended for repeated use

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

- wash hands to remove soil
- dispense palmful
- spread to cover hands, rub in well
- air dry, do not rinse or towel dry

Other information

- for additional information, see Material Safety Data Sheet (MSDS)
- for emergency medical information in USA, call 1.800.328.0026

Inactive ingredients water (aqua), isopropyl alcohol, propylene glycol, FDC red 40,

FDC blue 1

Questions? call 1.866.444.7450

Principal Display Panel with Representative Label

Hand Sanitizing Foam

Active ingredient: Benzalkonium chloride 0.1%

25 FL OZ (750 mL)

Pure Force™

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370 Wabasha Street North Made in U.S.A.

St. Paul, MN 55102 755509/7100/0713

<p>Drug Facts (continued)</p> <p>Other information</p> <ul style="list-style-type: none"> ■ for additional information, see Material Safety Data Sheet (MSDS) ■ for emergency medical information in USA, call 1.800.328.0026 <p>Inactive ingredients water (agua), isopropyl alcohol, propylene glycol, FD&C red 40, FD&C blue 1</p> <p>Questions? call 1.866.444.7450</p> <p>For service or additional information, call 1-866-444-7450. Para obtener información adicional o servicio de la unidad, llame al 1-866-444-7450. Product No./Producto No. 8000187</p> <p>Distributed by: © 2013 Ecolab USA Inc. PureForce All Rights Reserved/Todos los derechos reservados 370 Wabasha Street North Made in U.S.A./Hecho en EE.UU. St. Paul, MN 55102 755509/7100/0713</p>	<div style="background-color: #00a0a0; color: white; padding: 5px;"> <p>Hand Sanitizing Foam <i>Sanitizante de manos en espuma</i></p> </div> <p>Active ingredient: Benzalkonium Chloride 0.1% Ingrediente activo: Cloruro de benzalkonium 0.1%</p> <p>25 FL OZ (750 mL) pure FORCE™</p> <div style="border: 1px solid black; padding: 2px;"> <p>Peel Here For Drug Facts</p>  </div>
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<p>Drug Facts</p> <table border="1" style="width: 100%;"> <tr> <th style="text-align: left;">Active ingredient</th> <th style="text-align: left;">Purpose</th> </tr> <tr> <td>Benzalkonium chloride 0.1%.....</td> <td>Antiseptic handwash</td> </tr> </table> <p>Uses</p> <ul style="list-style-type: none"> ■ for handwashing to decrease bacteria on the skin ■ recommended for repeated use <p>Warnings</p> <ul style="list-style-type: none"> ■ For external use only <p>Do not use</p> <ul style="list-style-type: none"> ■ In eyes 	Active ingredient	Purpose	Benzalkonium chloride 0.1%.....	Antiseptic handwash	<p>Drug Facts (continued)</p> <p>When using this product</p> <ul style="list-style-type: none"> ■ If in eyes, rinse promptly and thoroughly with water ■ discontinue use if irritation and redness develop <p>Stop use and ask a doctor if skin irritation or redness occurs for more than 72 hours</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p> <p>Directions</p> <ul style="list-style-type: none"> ■ wash hands to remove soil ■ dispense palmful ■ spread to cover hands, rub in well ■ air dry, do not rinse or towel dry
Active ingredient	Purpose				
Benzalkonium chloride 0.1%.....	Antiseptic handwash				

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-504
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-504-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/28/2013	
2	NDC:47593-504-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/28/2013	09/28/2017

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/28/2013	

Labeler - Ecolab Inc. (006154611)

Revised: 7/2024

Ecolab Inc.