DIVAPROTECT HAND SANITIZER- benzalkonium chloride liquid
Diva International 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.

Active ingredient

Benzalkonium Chloride 0.13%

Uses

Effective in destroying harmful bacteria. Do not use on DivaCup.

Warnings

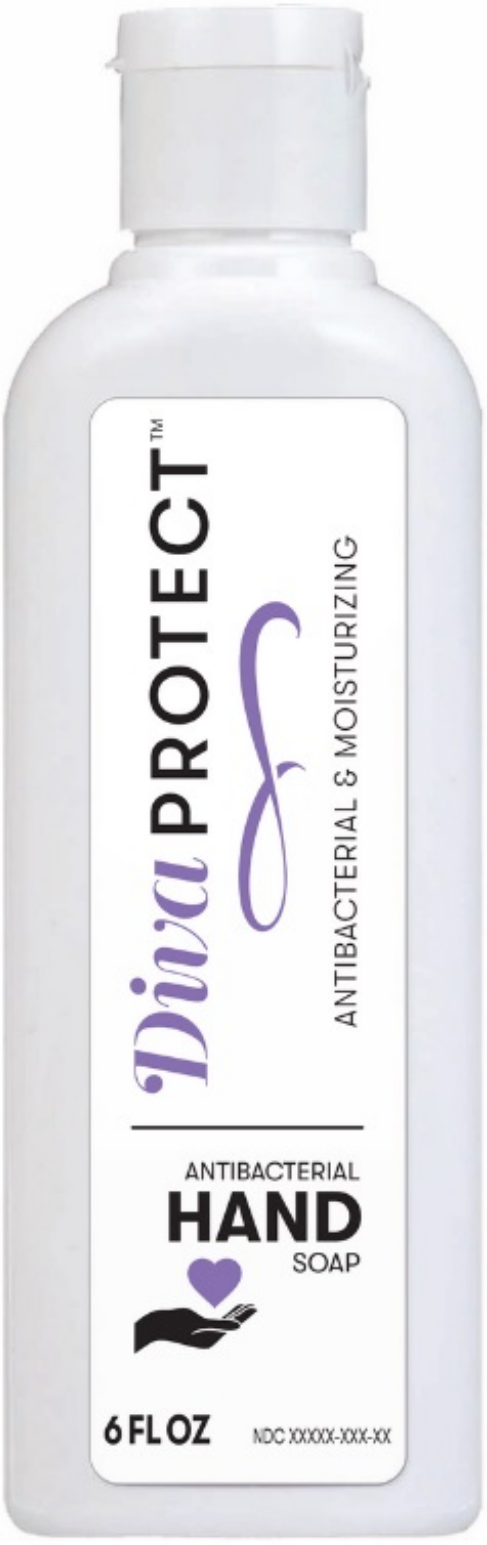
- For external use only.
- When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- Stop use and ask a doctor if irritation or rash appears and lasts.

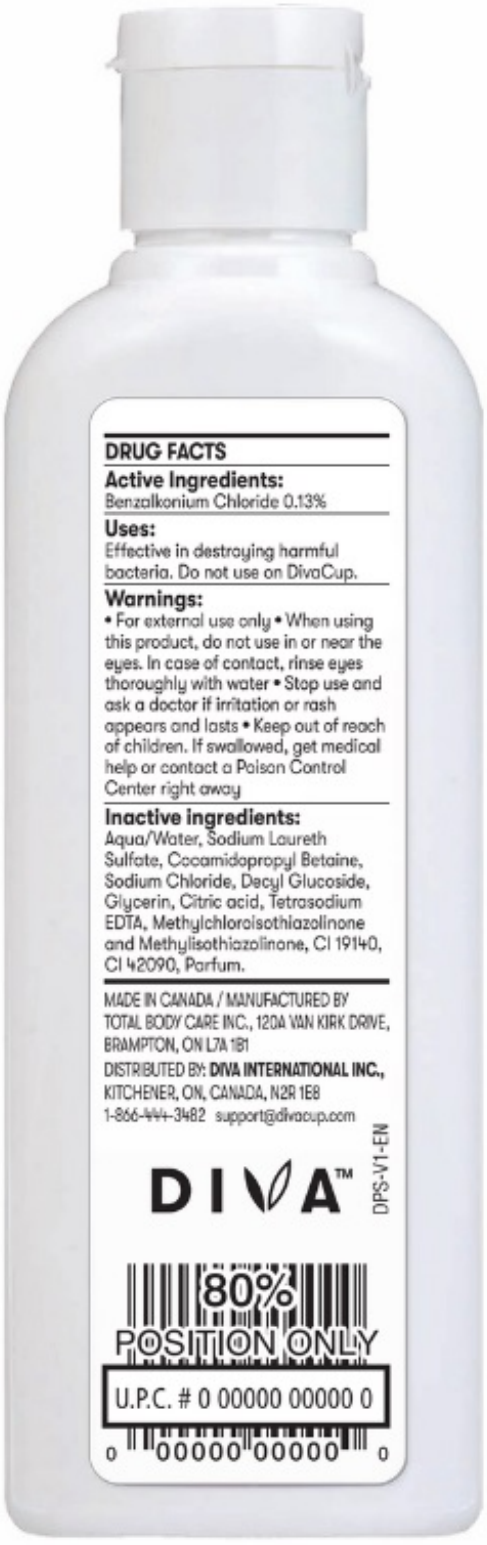
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Inactive ingredients

Aqua/Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Decyl Glucoside, Glycerin, Citric acid, Tetrasodium EDTA, Methylchloroisothiazolinone and Methylisothiazolinone, CI 19140, CI 42090, Parfum.

Package Label - Principal Display Panel


Diva PROTECT™
ANTIBACTERIAL & MOISTURIZING
ANTIBACTERIAL
HAND
SOAP
6 FL OZ
NDC XXXXX-XXX-XX


DRUG FACTS
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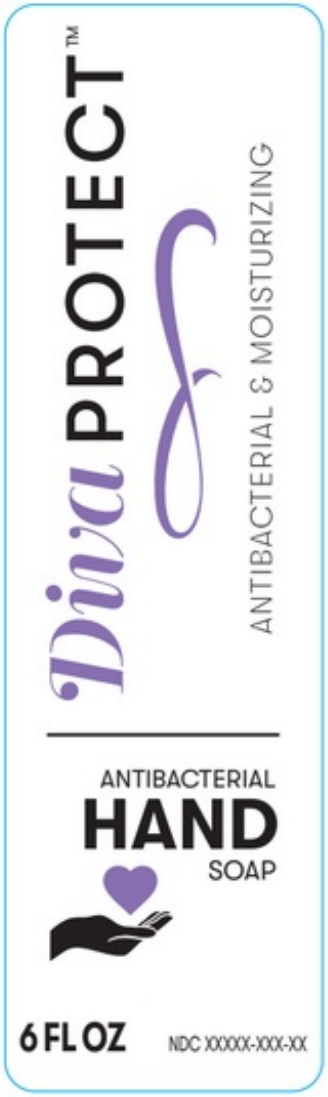
MADE IN CANADA / MANUFACTURED BY TOTAL BODY CARE INC., 120A VAN KIRK DRIVE, BRAMPTON, ON L7A 1B1
DISTRIBUTED BY: DIVA INTERNATIONAL INC., KITCHENER, ON, CANADA, N2R 1E8
1-866-444-3482 support@divacup.com
DIVA™
DPS-V1-EN
80%
POSITION ONLY
U.P.C. # 0 00000 00000 0
0 00000 00000 0



DIVAPROTECT HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77116-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	

COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE/METHYLISOTHIAZOLINONE MIXTURE (UNII: 15O9QS218W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77116-002-01	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/25/2020	

Labeler - Diva International Inc. (207651758)

Revised: 5/2020 Diva International Inc.