TOPSANI- benzalkonium chloride liquid
United Promotions 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.

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topsani
HAND SANITIZER + MOISTURIZER

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.125%

Purpose

Antimicrobial

Indications

For handwashing to decrease bacteria on the skin.

Warning

For external use only.

When using this product avoid contact with the eyes. Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

Keep out of reach of children. If swallowed get medical help or contact a poison control center right away.

Directions

Wet hands thoroughly with product and allow to dry without wiping.

Inactive Ingredients

Blue 1, Fragrance, Hydroxypropyl Methylcellulose, Polysorbate-20, SD Alcohol 40-B, Urea, Water

Questions or Comments?

Call 1-877-541-6055

Dist. by UPI, Atlanta, GA 30326 USA

PRINCIPAL DISPLAY PANEL - 56 mL Bottle Label

non-alcohol based

PronTech™
Technology

topsani

HAND SANITIZER + MOISTURIZER

Lasting Anti-Bacterial Protection

KILLS 99.999%
of microorganisms

all you
need is a
dimesize
drop

2 FL OZ (56 mL)

non-alcohol based

PronTech™ Technology

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TOPSANI

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24439-321
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.25 mL in 1000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
UREA (UNII: 8W8T17847W)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	BLUE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24439-321-10	56 mL in 1 BOTTLE, PLASTIC		
2	NDC:24439-321-11	236 mL in 1 BOTTLE, PUMP		
3	NDC:24439-321-12	946 mL in 1 BOTTLE, PLASTIC		
4	NDC:24439-321-13	946 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	11/01/2011	

Labeler - United Promotions Inc. (796252054)

Revised: 12/2011 United Promotions Inc.