# UP AND UP MOISTURIZING BODY WASH- benzalkonium chloride liquid TARGET CORPORATION

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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## **DRUG FACTS**

## ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE 0.13%

#### **PURPOSE**

ANTIBACTERIAL

#### **USES**

FOR WASHING TO DECREASE BACTERIA ON THE SKIN

## WARNINGS

FOR EXTERNAL USE ONLY

## WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USING THIS PRODUCT AND ASK DOCTOR IF

IRRITATION AND REDNESS DEVELOPS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

## **DIRECTIONS**

SQUEEZE A SMALL AMOUNT ONTO A WET WASHCLOTH, SPONGE, POUF OR HANDS AND APPLY TO BODY. WORK INTO A RICH LATHER AND RINSE OFF

## OTHER INFORMATION

STORE AT ROOM TEMPERATURE

## INACTIVE INGREDIENTS

WATER (AQUA), COCAMIDOPROPYL BETAINE, DECYL GLUCOSIDE, HYDROXYETHYLCELLULOSE, GLYCERIN, CITRIC ACID, POLOXAMER 124, FRAGRANCE (PARFUM), TETRASODIUM EDTA, POLYQUATERNIUM-7, SODIUM CITRATE, ALOE BARBADENSIS LEAF JUICE, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, BLUE 1 (CI 42090), EXT. VIOLET 2 (CI 60730)

## **QUESTIONS OR COMMENTS?**

1-800-910-6874

## LABEL COPY

Compare to the performance of Dial® Spring Water® Body Wash\* moisturizing body wash antibacterial new formula sulfate free triclosan free fresh scent up&up 21 FL OZ (621 mL)

up & up® antibacterial moisturizing body wash, with its invigorating scent, will moisturize and refresh your skin. Its antibacterial formula will make sure you're clean.

## Drug Facts

Active ingredient Benzalkonium Chloride 0.13%

Purpose Antibacterial

Uses ■ For washing to decrease bacteria on the skin.

## Warnings

For external use only.

When using this product ■ avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop using this product and ask doctor if irritation and redness develops.

Keep out of reach of children. ■ In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions ■ Squeeze a small amount onto a wet washcloth, sponge, pouf or hands and apply to body. Work into a rich lather and rinse off.

Other information 
Store at room temperature.

Inactive ingredients: Water (Aqua),
Cocamidopropyl Betaine, Decyl Glucoside,
Hydroxyethylcellulose, Glycerin, Citric Acid,
Poloxamer 124, Fragrance (Parfum), Tetrasodium
EDTA, Polyquaternium-7, Sodium Citrate, Aloe
Barbadensis Leaf Juice, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1 (CI 42090),
Ext. Violet 2 (CI 60730).

Questions or comments? 1-800-910-6874

This product is cruelty free, gluten free and manufactured in an environmentally sustainable facility.

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Distributed by Target Corporation
Minneapolis, MN 55403
Made in Canada

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06-19529

06-19528

benzalkonium chloride liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-815	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>BENZALKO NIUM CHLO RIDE</b> (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL		

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3011KX)				
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)				
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70 SE62ZAR)				
GLYCERIN (UNII: PDC6A3C0OX)				
CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP)				
POLOXAMER 124 (UNII: 1S66E28KXA)				
EDETATE SO DIUM (UNII: MP1J8420LU)				
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)				
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)				

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	<b>Marketing End Date</b>
1	NDC:11673-815-21	621 mL in 1 BOTTLE, PLASTIC		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	03/05/2014		

## Labeler - TARGET CORPORATION (006961700)

## Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(11673-815)	

Revised: 3/2014 TARGET CORPORATION