DALAN THERAPY ANTIBACTERIAL HAND WASH, ACTIVE PROTECTION-
benzalkonium chloride liquid
DALAN KIMYA ENDUSTRI ANONIM SIRKETI

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.

Dalan Therapy Antibacterial Hand Wash, Active Protection

Drug Facts

Active ingredients

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

For handwashing to decrease bacteria on the skin.

Warnings

For external use only:

When using this product.

- Do not use in the eyes.

Stop use and ask a doctor if

irritation and redness develop. if condition persists for more than 72 hours consult a doctor

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wet hands and forearms. Apply 5 milliliters (teaspoonful) or palmful to hands and forearms. Scrub thoroughly for 60 seconds. Rinse and Repeat

Inactive ingredients:

Water, Cocamidopropyl Betaine, Lauryl Amine Oxide, Cocamine Oxide, Cocamidopropylamine Oxide, Sodium Chloride, Glycerine, Lactic Acid, Sodium Lactate, PEG-120 Methyl Glucose Dioleate, Fragrance, Laureth-7 Citrate, Tetrasodium EDTA, Benzophenone-3, Triethylene Glycol, Propylene Glycol, Benzyl Alcohol, Methylchloroisothiazolinone, Methylsothiazolinone, Limonene, Yellow 5, Red 40.

Package Labeling:

DALAN THERAPY ANTIBACTERIAL HAND WASH, ACTIVE PROTECTION

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51209-014
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)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	

COCAMINE OXIDE (UNII: QWA2IZI6FI)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID (UNII: 33X04XA5AT)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
EDETATE SODIUM (UNII: MP1J8420LU)	
OXYBENZONE (UNII: 95OOS7VE0Y)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51209-014-00	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/03/2020	

Labeler - DALAN KIMYA ENDUSTRI ANONIM SIRKETI (566219285)

Revised: 4/2020 DALAN KIMYA ENDUSTRI ANONIM SIRKETI