

**COVEX ANTIBACTERIAL BAR COOL PROTECTION- benzalkonium chloride soap
DALAN KIMYA ENDUSTRI ANONIM SIRKETI**

Covex Antibacterial Bar Soap Cool 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 to hands and forearms. Scrub thoroughly for 60 seconds. Rinse and repeat

Inactive ingredients:

Sodium Tallowate, Sodium Palmate, Water, Sodium Palm Kernelate, Fragrance, Sodium Chloride, Glycerine, Zinc Oxide, C12-15 Alkyl Lactate, Titanium Dioxide, Tetrasodium EDTA, Etidronic Acid, Disodium Distyrylbiphenyl Disulfonate, Linalool, Limonene, Geraniol,

Citral.

Package Labeling:51209-022-90



Package Labeling:51209-022-27



COVEX ANTIBACTERIAL BAR COOL PROTECTION
 benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51209-022
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 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ETIDRONIC ACID (UNII: M2F465ROXU)	
DISODIUM DISTYRYLBIPHENYL DISULFONATE (UNII: 81G7704MYR)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
GERANIOL (UNII: L837108USY)	
CITRAL (UNII: T7EU009VPP)	
SODIUM TALLOWATE, BEEF (UNII: 07RIK6QMEW)	
SODIUM PALMATE (UNII: S0A6004K3Z)	
WATER (UNII: 059QF0KO0R)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51209-022-90	90 g in 1 BOX; Type 0: Not a Combination Product	02/24/2021	
2	NDC:51209-022-27	270 g in 1 PACKAGE; Type 0: Not a Combination Product	02/24/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/24/2021	

Revised: 1/2026

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