

CREAMOL HAND SANITIZING WIPES 80 PCS- alcohol swab
LABER KIMYA AR-GE SANAYI TICARET - LEVENT KAHRIMAN

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.

CREAMOL HAND SANITIZING WIPE 80 pcs

Alcohol: 75% (w/w) Purpose: Antimicrobial

Antimicrobial

Directions

Wipe hands, discard

Hand sanitizer to help reduce bacteria on skin. Kills 99.9% of most common germs that may cause illness.

For external use only. Flammable, keep away from fire and flame.

Do not use on open wounds, in or near the eyes and in children less than 2 months of age.

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

If irritation and redness develops and persists and in case of poisoning stop use and ask a doctor. Symptoms of Poisoning are dizziness, loss of consciousness and speech disorder.

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

isopropyl alcohol, glycerine, alpha-tocopherol, panthenol, water

Other informations

Store between 15-30 °C. Avoid freezing and excessive heat above 40°C



CREAMOL HAND SANITIZING WIPES 80 PCS
alcohol swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77892-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	45 g in 100 g
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	30 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PANTHENOL (UNII: WV9CM0O67Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL, D- (UNII: N9PR3490H9)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77892-0012-1	4 g in 1 POUCH; Type 0: Not a Combination Product	09/07/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/07/2020	

Labeler - LABER KIMYA AR-GE SANAYI TICARET - LEVENT KAHRIMAN (502952094)

Establishment

Name	Address	ID/FEI	Business Operations
LABER KIMYA AR-GE SANAYI TICARET - LEVENT KAHRIMAN		502952094	manufacture(77892-0012)

Revised: 9/2020

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