SANITIZING WIPES- benzalkonium chloride swab
Wipes LLC

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

Do not use if

you are allergic to any of this ingredients

When using this product

avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor

If irriatation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children

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

Directions

Adults and Children 2 Years and Over: Apply to hands. Allow to dry without wiping.

Children under 2 Years: Ask a doctor before use

To dispense: Pull tab and remove large cover. Insert triangular threading loop through slits in cover. Do not push finger through the opening. Thread first sheet in center of roll through loop. Replace cover. Pull loop back through opening. Pull each sheet up and slightly to the side. Dispose of wipe in trash. **DO NOT FLUSH.**

Other information

Store below 95 ^{O}F (35 ^{O}C). keep closed tightly. May discolor certain fabric or surfaces

Inactive ingredients

Water, SD Alcohol 40, Phenoxyethanol, Decyl Glucoside, Potassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric Acid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance

Principal Display Panel

NDC 70792-1200-1

Sanitizing Wipes by WIPES.com

Kills 99.99% of most common germs that may cause illness. Ideal for offices, restaurants, health clubs and more!

Will not harm vinyl, plastic, metal and rubber

1200 Wipes

6 in. x 8 in. (15.2 cm x 20.3 cm)

SANITIZING WIPES

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70972-1200
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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70972-1200-1	1200 in 1 BAG	04/04/2017	
1		0.00273 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/04/2017	

Labeler - Wipes LLC (964227347)

Registrant - Precare Corp. (117111327)

Establishment

Name	Address	ID/FEI	Business Operations
Precare Corp.		117111327	manufacture(70972-1200)

Revised: 12/2024 Wipes LLC