FRAGRANCE FREE SANITIZING WIPES- benzalkonium chloride cloth GoodEarth Distribution Llc

Fragrance Free Sanitizing Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13 %

Purpose

Antimicrobal

Use

Hand sanitizer to help reduce bacteria on the skin.

Recommended for repeated use.

Warnings

For external use only.

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly in water.

Discontinue use if irritation and redness develop. If conditions persist for more than 72 hours, consult a physician.

If swallowed get medical help or contact a poison control center immediately.

Keep out of reach of children.

Directions

- Wet hands thoroughly with product and allow to dry.
- Be sure to use entire wipe.
- Discard after single use.
- Children under 6 years of age should be supervised when using this product.

Inactive ingredients

Benzoic Acid, Caprylyl/Capryl Oligoglucoside, Dehydroacetic Acid, Phenoxyethanol, Poly(Laurylglucoside)-7, Propylene Glycol, Water

1800 Wipe Pouch Label

GoodEarth

DISTRIBUTION

FRAGRANCE FREE

Kills 99.99%

of most common germs

that may cause illness

Sanitizing

wipes

1800

count

Premium fabric quickly

removes dirt and soil

MICRODOT •



FRAGRANCE FREE SANITIZING WIPES

benzalkonium chloride cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71745-001	
Route of Administration	CUTANEOUS			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg	

Inactive Ingredients		
Ingredient Name	Strength	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)		
WATER (UNII: 059QF0KO0R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)		
POLY(LAURYLGLUCOSIDE)-7 (UNII: VB00RDE21R)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
BENZOIC ACID (UNII: 85KN0B0MIM)		

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71745-001- 02	2 in 1 BOX	09/25/2017		
1	NDC:71745-001- 01	1800 in 1 POUCH; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	09/25/2017		

Labeler - GoodEarth Distribution Llc (079808558)

Registrant - Innocore Sales & Marketing Inc (201152597)

Revised: 12/2023 GoodEarth Distribution Llc