

COLOR ELEMENT ALCOHOL FREE HAND SANITIZER- benzalkonium chloride liquid
Zhejiang iColor Biotech Co., Ltd

Color Element Alcohol Free Hand Sanitizer

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose:

Antiseptic

Uses

For hand washing to help reduce bacteria that potentially can cause disease. Recommended for repeated use. For use when soap and water are not available.

Warnings

For external use only

When using this product

do not use in or near eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash appears on the skin.

Keep out of reach of children.

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

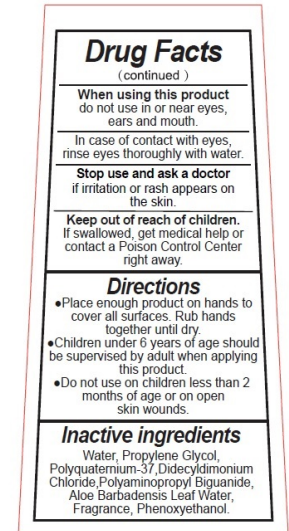
Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Children under 6 years of age should be supervised by adult when applying this product.
- Do not use on children less than 2 months of age or on open skin wounds.

Inactive ingredients

Water, Propylene Glycol, Polyquaternium-37, Didecylidimonium Chloride, Polyaminopropyl Biguanide, Aloe Barbadensis Leaf Water, Fragrance, Phenoxyethanol.

Package Labeling:30ml



Package Labeling:2ml



COLOR ELEMENT ALCOHOL FREE HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74934-048
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIDCYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
POLYAMINOPROPYL BIGUANIDE (UNII: DT9D8Z79ET)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74934-048-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/07/2020	01/31/2026
2	NDC:74934-048-02	2 mL in 1 BAG; Type 0: Not a Combination Product	09/07/2020	01/31/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/07/2020	01/31/2026

Labeler - Zhejiang iColor Biotech Co., Ltd (554528308)

Revised: 1/2024

Zhejiang iColor Biotech Co., Ltd