CRAZY CLEANZ HAND SANITIZER EUCALYPTUS SCENTED- benzalkonium chloride liquid
Landy International

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.

51706-954 Crazy Cleanz Hand Sanitizer Eucalyptus Scented 0.1% Benzalkonium chloride

Active Ingredient(s)

Benzalkonium Chloride 0.1%

Purpose

Antiseptic

Use

helps eliminate bacteria on hands

Warnings

For external use only
Do not use
. In children less than 2 months of age. On open skin wounds
When using this product keep out of eyes, ears, and mouth.
In case of contact with eyes, rinse eyes thouroughly with water.
Stop use and ask a doctor if irritation or rash occurs.
These may be a sign of a serious condition.

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

Directions

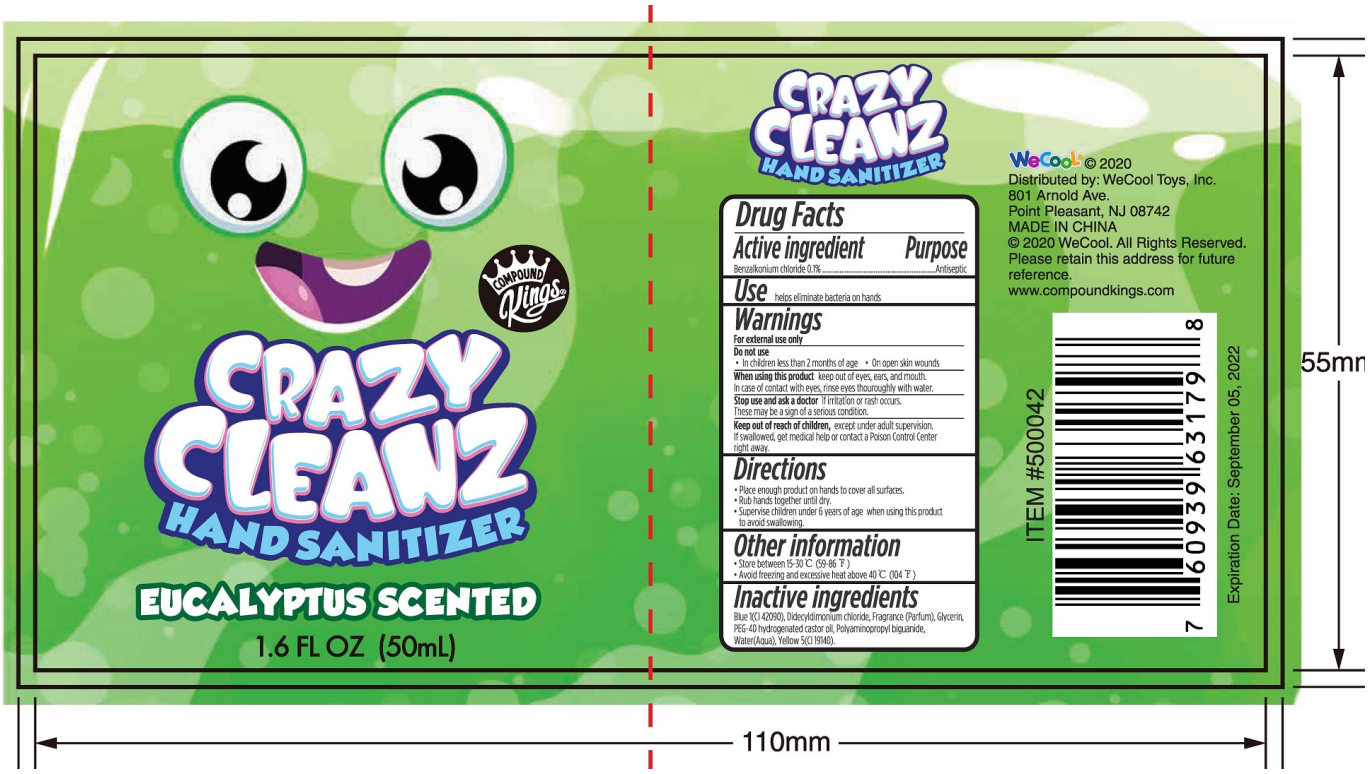
-Place enough product on hands to cover all surfaces.
· Rub hands together until dry.
.Supervise children under 6 years of age when using this product to avoid swallowing.

Inactive ingredients

Blue 1(Cl 42090), Didecyldimonium chloride,Fragrance(Parfum),Glycerin, PEG-40 hydrogenated castor oil, Polyaminopropyl biguanide,

Water(Aqua),Yellow 5(C119140).

Package Label - Principal Display Panel



CRAZY CLEANZ HAND SANITIZER EUCALYPTUS SCENTED

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51706-954
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
POLYAMINOPROPYL BIGUANIDE (UNII: DT9D8Z79ET)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51706-954-01	50 g in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/15/2020	

Labeler - Landy International (545291775)

Establishment

Name	Address	ID/FEI	Business Operations
Landy International		545291775	manufacture(51706-954)

Revised: 3/2022 Landy International