CB ADVANCED HAND SANITIZER- alcohol gel Cb Distributors, Inc.

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.

CB Advanced Hand Sanitizer

Drug Facts

Active ingredient Ethyl alcohol 75%

Purpose

Antiseptic

Use

• to help reduce bacteria on the skin.

Warnings

For external use only

Flammable. Keep away from fire or flame.

When using this product

do not use in or near the eyes. In case of contact, flush eyes with water.

Stop use and ask a doctor if

irritation or rash appears and lasts.

Keep out of reach of children.

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

Directions

• wet hands thoroughly with product and allow to dry without wiping • Children under 6, use only under adult supervision.

Other Information

• store below 105°F • may discolor certain fabrics.

Inactive Ingredients

• Purified water(Aqua), Aloe Barbadensis Leaf Extract, Acrylates/C10-30 alkyl Acrylate Crosspolymer, Tetrahydroxypropyl Ethylenediamine.

Package Labeling:

2 FL OZ (60 mL)	KILLS MOST ILLNESS CAUSING GERMS		SANTIZER				
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Production Date: 05/22/2020 Valid Time: 3 Years Expiry Date: 05/22/2023 Distributed by CB Distributors, Inc., Beloit, WI 53511 www.cbprices.com (608) 368-9909



CB ADVANCED HAND	SANITIZER								
alcohol gel									
Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:76813-014					
Route of Administration	TOPICAL								
Active Ingredient/Active Moiety									
Ingre	Basis of Strength St		rength						
ALCOHOL (UNII: 3K9958V90M) (AL		ALCOHOL 0.75 m		L in 1 mL					
Inactive Ingredients									
Ingredient Name					Strength				
WATER (UNII: 059QF0KO0R)									
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)									
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)									
EDETOL (UNII: Q4R969U9FR)									
Packaging									
# Item Code	Package Description	Ma	arketing Start Date	Marketin	g End Date				
1 NDC:76813-014-00 60 mL in 1 BO	TLE; Type 0: Not a Combination F	Product 05/	/28/2020						
Marketing Information									

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/28/2020	

Labeler - Cb Distributors, Inc. (799862735)

Revised: 6/2020

Cb Distributors, Inc.