CB ADVANCED HAND SANITIZER- ethyl alcohol gel gel
Nutritech Pharmaceuticals, 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.

Active Ingredients

Ethyl alcohol 75%

Purpose

Antiseptic

Uses

Hand sanitizer

Warnings

For external use only.

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

Discontinue if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

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

Directions

Place enough product in palm to cover hands and rub hands together briskly until dry.

Inactive Ingredients

Purified Water, Glycerin, Triethanolamine, Carbomer fragrance, Vitamin E, Aloe Vera.

Made in China

Drug Facts

Active ingredient	Purpose
Ethyl alcohol 75%	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 • water, alcohol, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, acrylates/C10-30 alkyl acrylate, fragrance, crosspolymer

Distributed by
CB Distributors, Inc.,
Beloit, WI 53511
www.cbprices.com

3 65860 21003 8

CB ADVANCED HAND SANITIZER

ethyl alcohol gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73529-353
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73529-353-01	237 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/16/2020	

Labeler - Nutritech Pharmaceuticals, Inc (117342868)

Revised: 5/2020 Nutritech Pharmaceuticals, Inc