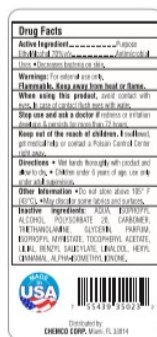


ETHYL ALCOHOL- ethyl alcohol gel

Chemco Corporation

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.

49783-700-32





ETHYL ALCOHOL

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49283-700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	0.1 mL in 100 mL
POLYSORBATE 20 (UNII: 7T1F30V5YH)	0.45 mL in 100 mL
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.05 mL in 100 mL
WATER (UNII: 059QF0KO0R)	1 mL in 100 mL
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	0.01 mL in 100 mL
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540GJV69)	0.01 mL in 100 mL
LINALOOL, (+)- (UNII: F4VNO44C09)	0.01 mL in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.35 mL in 100 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302)	0.1 mL in 100 mL

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-700-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/02/2020	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/02/2020	

Labeler - Chemco Corporation (032495954)

Registrant - Chemco Corporation (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
Chemco Corporation		032495954	manufacture(49283-700)