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.

49283-007-01

ETHYL ALCOHOL
ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49283-007
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**
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540GJV69)	
ISOMETHYL-.ALPHA.-IONONE (UNII: 9XP4LC555B)	
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
TROLAMINE (UNII: 9O3K93S3TK)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-007-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/07/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/07/2020	

Labeler - Chemco Corporation (032495954)

Registrant - Chemco Corporation (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
Chemco Corporation		032495954	manufacture(49283-007)

Revised: 8/2020 Chemco Corporation