CAMPHOR- camphor 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.

49283-331-04

CAMPHOR

camphor gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49283-331
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	94.8 mL in 99 mL

Inactive Ingredients	
Ingredient Name	**Strength**
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
WATER (UNII: 059QF0KO0R)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
OLETH-20 (UNII: YTH167I2AG)	
MENTHOL (UNII: L7T10EIP3A)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DIPROPYLENE GLYCOL MONOCAPRATE MONOCAPRYLATE (UNII: 5980W979EL)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-331-28	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/11/2020	
2	NDC:49283-331-05	18927 mL in 1 PAIL; Type 0: Not a Combination Product	06/23/2020	
3	NDC:49283-331-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/11/2020	

Labeler - Chemco Corporation (032495954)

Registrant - Chemco Corporation (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
Chemco Corporation		032495954	manufacture(49283-331)