HAND SANITIZING GEL BY CN PHARMA- alcohol gel
Canadian National Pharma Group 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 Ingredient

Ethanol 75%

Purpose

Antiseptic

Uses

- Hand Sanitizer to help reduce bacteria that potentially can cause disease.
- For use when soap and water are not available.

For external use only.
Flammable. Keep away from heat or Flame.
Do not use:

- In children less than 2 months of age
- On open skin wounds

When using this product keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

Store between 15-30^{o}C (59-86^{o}F)
Avoid freezing and excessive heat above 40^{o}C (104^{o}F)

Inactive ingredients

Glycerin, hydrogen peroxide, carbomer, triethanolamine, denatonium benzoate, fragrance, distilled water.

Package Label - Principal Display Panel

HAND SANITIZING GEL BY CN PHARMA

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77536-001
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**
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
WATER (UNII: 059QF0KO0R)	
MENTHA SPICATA OIL (UNII: C3M81465G5)	

EAST INDIAN LEMONGRASS OIL (UNII: UP0M8M3VZW)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
TEA TREE OIL (UNII: VIF565UC2G)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
COCONUT OIL (UNII: Q9L0O73W7L)	
ORANGE OIL (UNII: AKN3KSD11B)	
ALOE (UNII: V5VD430YW9)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77536-001-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
2	NDC:77536-001-02	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
3	NDC:77536-001-03	275 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
4	NDC:77536-001-04	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
5	NDC:77536-001-05	945 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
6	NDC:77536-001-06	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
7	NDC:77536-001-07	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2020	

Labeler - Canadian National Pharma Group Inc (204126080)

Establishment

Name	Address	ID/FEI	Business Operations
Canadian National Pharma Group Inc		204126080	manufacture(77536-001)

Revised: 9/2020 Canadian National Pharma Group Inc