

H C- h c gel
Guangzhou cimei bio technology co.ltd

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.

ALCOHOL HAND CLEAN GEL

Active Ingredient

Ethyl Alcohol 62.0%

Purpose

Antimicrobial

Uses

Hand sanitizer to help reduce bacteria on skin.

WARNINGS

Flammable, keep away from fire or flame, For external use only When using this products,do not use in or near the eyes.In case of the contact,Rinse eyes thoroughly with water, Stop use and ask doctor if irritation or rash appears and lasts, Keep out of reach for children.If swallowed, get medical help or contact a poison control center right away.

DIRECTIONS

Place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry. Children Under 6 years of age should be supervised when using this product.

Other information

Store below 106°F.(41°C)

May discolor certain fabrics or surfaces.

INACTIVE INGREDIENTS

Water(Aqua),Aloe Barbadensis Leaf Juice, Carbomer,Fragrance,Glycerin, Propylene Glycol, Tocopheryl Acetate(Vitamin E),

Triethanolamine,FD&C Blue NO.1,FD&C Yellow NO 5.

Package Label - Principal Display Panel



H C

h c gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:75233-001

Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	ALOE VERA LEAF (UNII: ZY81Z83H0X)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)			
	TROLAMINE (UNII: 9O3K93S3TK)			
	FD&C BLUE NO. 1 (UNII: HBR47K3TBD)			
	WATER (UNII: 059QF0K00R)			
	FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
	CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75233-001-01	99 mL in 1 AMPULE; Type 0: Not a Combination Product	05/02/2020	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph not final	part333A	05/02/2020	

Labeler - Guangzhou cimei bio technology co.ltd (554529699)

Registrant - Guangzhou cimei bio technology co.ltd (554529699)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou cimei bio technology co.ltd		554529699	manufacture(75233-001)