DOCTOR HAND FRESH GEL- alcohol gel C-MAX KOREA 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.

Drug Facts

ethanol

water, carbopol, polysorbate 60, triethanolamine, peppermint oil

Antiseptic

instant healthcare personnel hand antiseptic to reduce bacteria that potentially can cause disease instant hand antiseptic to decrease bacteria on the skin recommended for repeated use

KEEP OUT OF REACH OF THE CHILDREN

Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.

Supervise children in the use of this product.

For external use only.

Flammable, keep away from fire or flame.

When using this product keep out of eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.

Stop use and ask a doctor if significant irritation or sensitization develops.

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

for external use only

Active Ingredient	Purpose Antiseptic
Uses Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use not available.	
Warnings 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 e with water. Stop use and ask a doctor if initation or rash occurs. These may be signs of a serio Keep out of reach of children. If swallowed, get medical help or contact a Poison C	us condition.
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 swallowin	
Other Information Store between 1-30C (33.8-88F) Avoid freezing and excessive heat above 40C (104F)	
Inactive Ingredients Water, Carborner, Triethanolamine, Polysorbate 20, Lemon Flavor	

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70949-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

	Ingredient Name	Basis of Strength	Strength
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	310 mL in 500 mL

Inactive Ingredients	
Ingredient Name	Strength
PEPPERMINT (UNII: V95R5KMY2B)	
TROLAMINE (UNII: 903K93S3TK)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:70949-002-	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/02/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/21/2020	

Labeler - C-MAX KOREA CO.,LTD. (557821673)

Registrant - C-MAX KOREA CO.,LTD. (557821673)

Establishment				
Name	Address	ID/FEI	Business Operations	
C-MAX KOREA CO.,LTD.		557821673	manufacture(70949-002)	

Revised: 4/2020 C-MAX KOREA CO.,LTD.