

ALOE CHOK CHOK HAND- alcohol gel
TONYMOLY 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.

ACTIVE INGREDIENT

ALCOHOL 65% w/w

INACTIVE INGREDIENT

Water, Glycerin, Carbomer, Triethanolamine, Hydroxyacetophenone, Fragrance, Ethylhexylglycerin, Trehalose, Hamamelis Virginiana (Witch Hazel) Water, Aloe Barbadensis Leaf Water, 1,2-Hexanediol, Sodium Hyaluronate, Linalool, Hexyl Cinnamal

PURPOSE

ANTISEPTIC

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 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.

Uses

Hand gel that kills germs that can potentially cause disease.

Directions

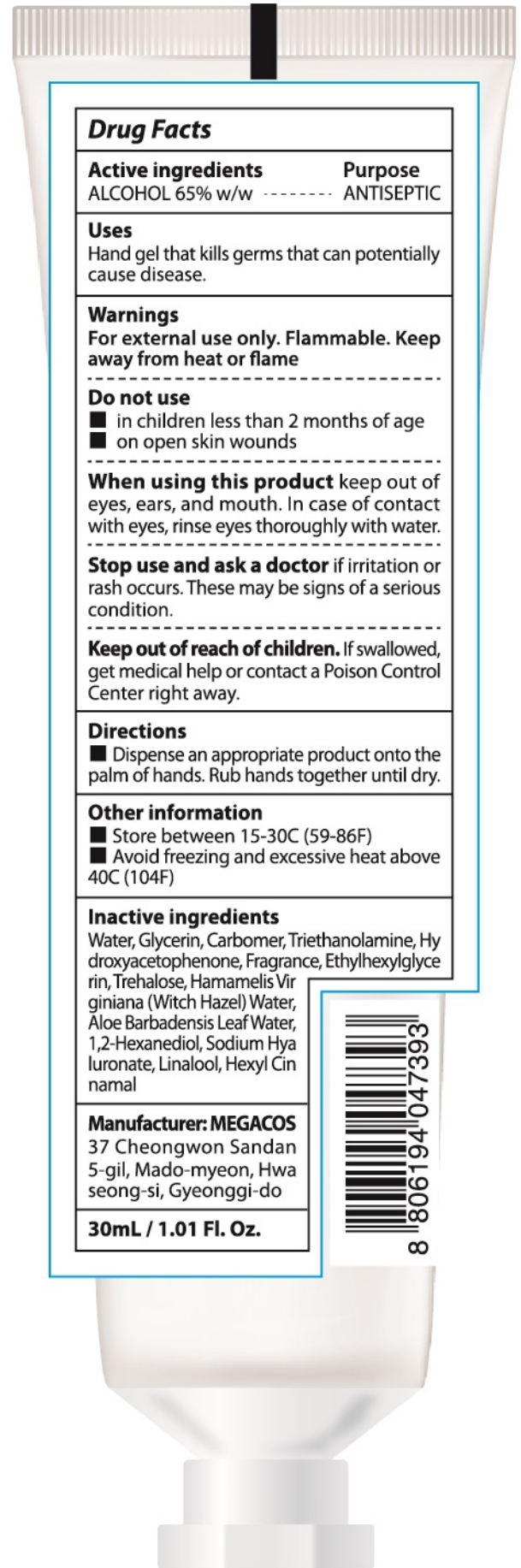
Dispense an appropriate product onto the palm of hands. Rub hands together until dry.

Other Information

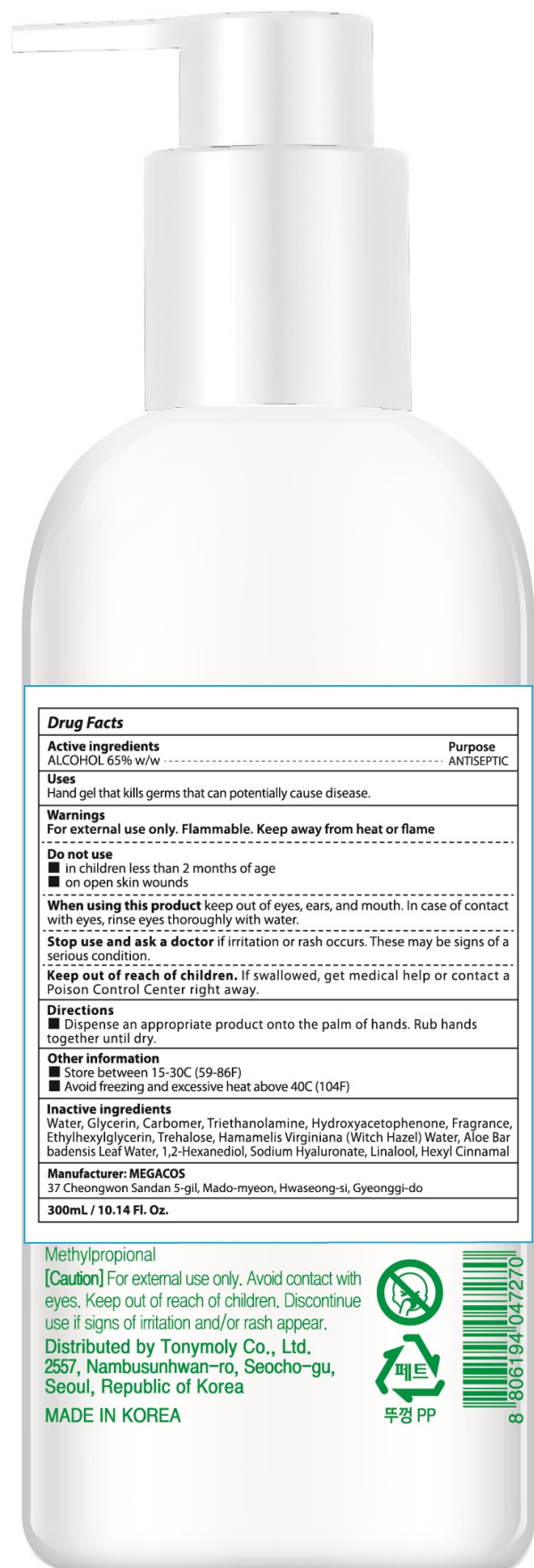
Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

PACKAGE LABEL : ALOE Chok Chok Hand Gel 30mL



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ALOE CHOK CHOK HAND

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59078-806
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Glycerin (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Trehalose (UNII: B8WCK70T7I)	
WITCH HAZEL (UNII: 101I4J0U34)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
1,2-Hexanediol (UNII: TR046Y3K1G)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59078-806-01	30 mL in 1 TUBE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:59078-806-02	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2020	

Labeler - TONYMOLY CO., LTD. (688216798)

Registrant - TONYMOLY CO., LTD. (688216798)

Establishment

Name	Address	ID/FEI	Business Operations
MEGACOS MANUFACTURING Co., Ltd.		694745986	manufacture(59078-806)

Revised: 5/2020

TONYMOLY CO., LTD.