BEAUTAIME HAND DOCTOR- alcohol gel ASAN C&S

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

Active ingredients: ETHANOL 62.0% w/w

INACTIVE INGREDIENT

Inactive ingredients

Water, Glycerin, Propylene Glycol, Triethanolamine, Carbomer, Aloe Barbadensis Leaf Powder, Aloe Extract

PURPOSE

Purpose: ANTISEPTIC

WARNINGS

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

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

Uses

Uses

Hand sanitizer to help reduce bacteria that potentially can cause disease.

Directions

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

Other information

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

PACKAGE LABEL - BEAUTAIME HAND DOCTOR GEL 70mL





Beau T'aime Hand Doctor Gel

Sterilizing hand sanitizer



Everyday Hand Doctor Gel

Washing your hands is the best habit for protecting your health

Ethanol62%



500ml

Drug Facts

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Distributor: ASAN C&S / 28, Hyuam-ro 79beon-gil, Wollong-myeon, Paju-si, Gyeonggi-do, Korea

Manufacturer: ASAN C&S / 28, Hyuam-ro 79beon-gil, Wollong-myeon, Paju-si, Gyeonggi-do, Korea

Net Wt:

500ml / 16.90 Fl. Oz.

BEAUTAIME HAND DOCTOR

alcohol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:73697-110

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ALOE VERA LEAF (UNII: ZY81Z83H0X)

Ingredient NameBasis of StrengthStrengthALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)ALCOHOL62 g in 100 mL

Inactive Ingredients

Ingredient Name

Strength

Water (UNII: 059QF0KO0R)

Glycerin (UNII: PDC6A3C0OX)

Propylene Glycol (UNII: 6DC9Q167V3)

Trolamine (UNII: 9O3K93S3TK)

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:73697-110- 01	70 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/01/2020				
2 NDC:73697-110- 02	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020				

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333E	04/01/2020					

Labeler - ASAN C&S (631139649)

Registrant - ASAN C&S (631139649)

Establishment						
Name	Address	ID/FEI	Business Operations			
ASAN C&S		631139649	manufacture(73697-110)			

Revised: 4/2020 ASAN C&S