BEAUTAIME HAND DOCTOR A- 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 70.0% 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

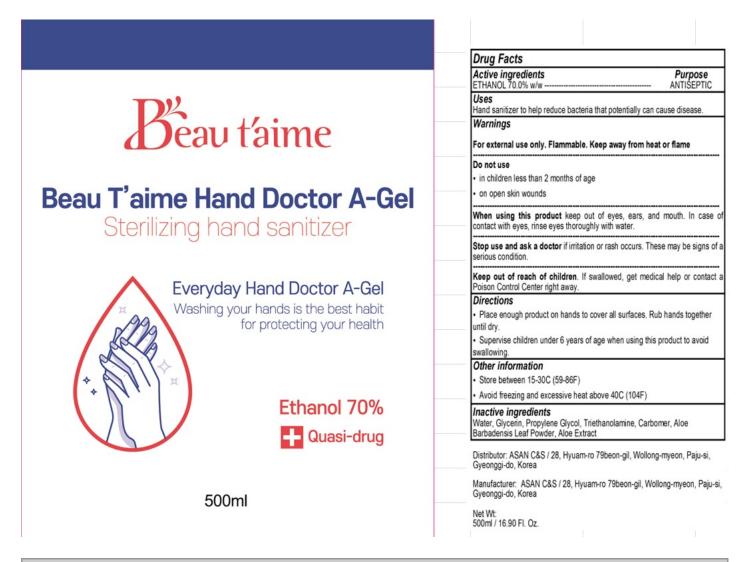
Other information

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

PACKAGE LABEL - BEAUTAIME HAND DOCTOR A GEL 70mL



PACKAGE LABEL - BEAUTAIME HAND DOCTOR A GEL 500mL



BEAUTAIME HAND DOCTOR A

alcohol gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:7		3697-120	
Route of Administration	TOPICAL				
Active Ingredient/Active Mo	iety				
Ingr		Basis of Strength		Strength	
ALCOHOL (UNII: 3K9958V90M) (AL		ALCOHOL 7		'0 g in 100 mL	
					0
					0
					0
Inactive Ingredients					
Inactive Ingredients	Ingredient Name				Strength
Inactive Ingredients Water (UNII: 059QF0K00R)	Ingredient Name				
	Ingredient Name				
Water (UNII: 059QF0KO0R)					
Water (UNII: 059QF0K00R) Glycerin (UNII: PDC6A3C0OX)					

ALOE VERA LEAF (UNII: ZY81Z83H0X)

ALOE (UNII: V5VD430YW9)

Packaging								
#	Item Code		Package Description	Marketing Start Date	Marketing End Date			
1	NDC:73697-120- 01	70 m	L in 1 CONTAINER; Type 0: Not a Combination Product	04/01/2020				
2	NDC:73697-120- 02	0- 500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		04/01/2020				
Marketing Information								
	Marketing Categ	orv	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	Marketing Catego	ury			-			

Labeler - ASANC&S (631139649)

Registrant - ASANC&S (631139649)

Establishment

Name	Address	ID/FEI	Business Operations
ASAN C&S		631139649	manufacture(73697-120)

Revised: 4/2020

ASAN C&S