AOMI HAND SANITIZER- alcohol spray Mirfeel 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.

ACTIVE INGREDIENT

ALCOHOL 70.0% (w/w)

INACTIVE INGREDIENT

Water, Glycerin, Fragrance, Cymbopogon Citratus Extract, Anthemis Nobilis Flower Extract, Aspalathus Linearis Extract, Melaleuca Alternifolia(Tea Tree)Leaf Extract

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 sanitizer to help reduce bacteria that potentially can cause disease.

Directions

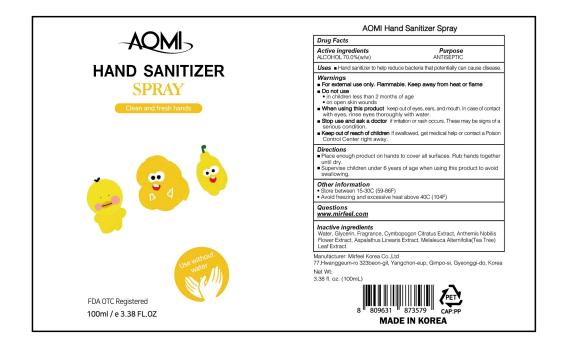
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

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

QUESTIONS

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





AOMI HAND SANITIZER

alcohol spray

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Prod	luct	Intori	mation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:75010-040

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	70 g in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Glycerin (UNII: PDC6A3C0OX)			
CHAMAEMELUM NO BILE FLO WER (UNII: O2T154T6OG)			
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)			

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75010-040-	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination	0 = /0 1/20 20	

0 1 P	roduct	03/01/2020		
Marketing Information				
Marketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fin	al part333E	05/01/2020		

Labeler - Mirfeel Korea Co., Ltd (695004258)

Registrant - Mirfeel Korea Co., Ltd (695004258)

Establishment				
Name	Address	ID/FEI	Business Operations	
Mirfeel Korea Co., Ltd		695004258	manufacture(75010-040)	

Revised: 6/2020 Mirfeel Korea Co., Ltd