

AOMI HAND SANITIZER CLEANSING PLUS- ethyl alcohol gel

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

ETHYL ALCOHOL 70.0% (w/w)

INACTIVE INGREDIENTS

Water, Carbomer 940, Triethanolamine, Jasminum Officinale (Jasmine) Extract, Prunus Mume Flower Extract, Bellis Perennis (Daisy) Flower Extract, Fragrance

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

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

Uses

To decrease bacteria on hands

Directions

- Please take out adequate amounts of gel on palm of your hand and rub your hands for 15 seconds at least.

Other Information

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

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

QUESTIONS

www.mirfeel.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



AOMI HAND SANITIZER CLEANSING PLUS

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:750 10-140
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		Alcohol	700 g in 1000 mL	
Inactive Ingredients				
Ingredient Name				Strength
Water (UNII: 059QF0KO0R)				
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
TROLAMINE (UNII: 9O3K93S3TK)				
JASMINUM OFFICINALE FLOWER (UNII: 0Q8K841432)				
PRUNUS MUME FLOWER (UNII: 2N8872050J)				
BELLIS PERENNIS FLOWER (UNII: 26I94X9A1K)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75010-140-01	1000 mL in 1 POUCH; Type 0: Not a Combination Product	09/01/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333E	09/01/2020	

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(75010-140)

Revised: 9/2020

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