

AKOIA HAND SANITIZER- alcohol gel
NG BEAUTY WORLD FZCO

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.

AKOIA HAND SANITIZING & Moisturizing Gel

Drug Facts

Active Ingredient[s]

Denatured Alcohol (Ethanol)

Purpose

Antiseptic

Use[s]

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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.

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

Inactive ingredients Aqua, Glycerin, Acrylates /C10-30 Alkyl Acrylate Crosspolymer, Parfum (Fragrance),

Aminomethyl Propanol, Aloe Barbadensis (Aloe Vera) Extract.

• INFINITE GLOW •

HAND SANITIZING & Moisturizing Gel

MOISTURIZING FORMULA

AKOIA: A hand sanitizing and moisturizing gel that sanitizes your hands while keeping them soft and smooth. The formula contains 70% alcohol that helps guarantees maximum possible protection against bacteria and germs.

Produced by:

NG Beauty World FZCO Dubai, UAE

For: AXA Beauty World S.R.L.

Piazza Borromeo, 12 - 20123 Milano, Italy

AXA Beauty World Limited

1164 High Road, Whetstone

London - UK

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Email: info@axabw.com

Tel: +4407720670777

Made in UAE

Packaging

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1000 ml

AKOIA HAND SANITIZER

alcohol gel

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		TOPICAL	NDC:79919-001	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	70 mL in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79919-001-01	5 mL in 1 PACKET; Type 0: Not a Combination Product	08/05/2020	
2	NDC:79919-001-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/05/2020	
3	NDC:79919-001-03	100 mL in 1 TUBE; Type 0: Not a Combination Product	08/05/2020	
4	NDC:79919-001-04	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/05/2020	
5	NDC:79919-001-05	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/05/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333A	08/05/2020	

Labeler - NG BEAUTY WORLD FZCO (557947924)

Establishment			
Name	Address	ID/FEI	Business Operations
NG BEAUTY WORLD FZCO		557947924	manufacture(79919-001)