

FRESHORIZE HAND SANITIZER- alcohol lotion

Freshorize, 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.

Drug Facts

Keep out of reach of children

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Purpose

Kills 99.9% Bacteria

Requires no water or towels.

Apply small amount to hands and rub lightly until dry.

Alcohol denat, Aqua, acrylates/C10-30 alkyl, acrylate cross polymer, triethanolamine (TEA), parfum

Active Ingredients

62.5% Ethyl Alcohol.

Use Information

Requires no water or towels. Apply small amount to hands and rub lightly until dry.

Warnings

Safety: For External Use Only. Use on Hands only.

Flammable. Keep away from fire or flame.

Avoid contact with Eyes, should this occur rinse immediately with clean warm water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Display of Package Label



Hand Sanitizer

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www.freshorize.com
www.avidairlineproducts.com

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Active ingredient: 62.5% Ethyl Alcohol
Ingredients: Alcohol denat, Aqua, acrylates/C10-30 alkyl acrylate cross polymer, triethanolamine (TEA), parfum
Manufactured under licence for Freshorize Ltd, Freshorize Ltd, Knowledge dock business centre, University way, London, E16 2HD, UK.
Tel: +44 7960 821632 Fax: +44 208223 7502
Made in China
Batch number:



NET 9.975 FL. OZ. 295ml

FRESHORIZE HAND SANITIZER

alcohol lotion

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		TOPICAL	NDC:52305-200	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	
Alcohol (UNII: 3K9958V90M) (alcohol - UNII:3K9958V90M)			Alcohol	
			185 mL in 295 mL	
Inactive Ingredients				
Ingredient Name			Strength	
Water (UNII: 059QF0KO0R)				
ETHANOLAMINE (UNII: 5KV86114PT)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52305-200-50	295 mL in 1 BOTTLE, PUMP		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333A	06/01/2010	

Labeler - Freshorize, Ltd. (424168503)

Registrant - Freshorize, Ltd. (424168503)

Establishment			
Name	Address	ID/FEI	Business Operations
Freshorize, Ltd.		424168503	manufacture

Revised: 11/2010

Freshorize, Ltd.