RUE21 OCEAN ANTI-BACTERIAL HAND SANITIZER - alcohol liquid ALILY (ZHANGZHOU) BATH PRODUCTS 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 Fact

Active Ingredient Purpose
Ethyl Alcohol 62% Antiseptic

Uses:

- To decrease bacteria on the skin that could cause diseases
- Recommended for repeated use.

Warnings:

For external use only-hands.

Flammable-keep away from heat and flame

When using this products

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.

Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately

Directions

- Wet hands thoroughly with product and allow to dry without wiping
- For children under 6, use only under adult supervision
- Not recommended for infants

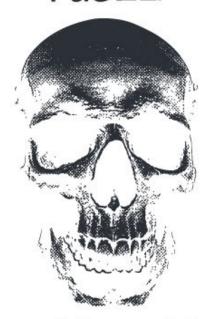
Other Information

- Do not store above 105 F
- May discolor some fabrics
- Harmful to wood finishes and plastics

Inactive Ingredients

Water (aqua), Glycerin, Triethanolamine, Fragrance (Parfume), PEG-40 Hydrogenated Castor Oil, Carbomer, DMDM Hydantoin, Hydrogenated Jojoba Oil, FD and C Yellow No. 5, FD and C Blue No. 1 pict

rue 21.



anti-bacterial hand sanitizer

Ocean 1.5 fl oz. / 45 ml Drug Facts Active Ingredients

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RUE21 OCEAN ANTI-BACTERIAL HAND SANITIZER

alcohol liquid

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:49371-043

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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	In	gredient Name		Basis of Strength	Strength	
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	A	ALCOHOL	62 g in 100 g	

DMDM HYDANTO IN (UNII: BYR0546 TOW)	0.2 g in 100 g
FD&C YELLOW NO. 5 (UNII: 1753WB2F1M)	0.0016 g in 100 g
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	0.005 g in 100 g

I	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:49371-043-01	45 g in 1 BOTTLE			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333	05/20/2010		

Labeler - ALILY (ZHANGZHOU) BATH PRODUCTS LTD. (531064269)

Registrant - ALILY (ZHANGZHOU) BATH PRODUCTS LTD. (531064269)

Establishment				
Name	Address	ID/FEI	Business Operations	
ALILY (ZHANGZHOU) BATH PRODUCTS LTD.		531064269	manufacture	

Revised: 5/2010 ALILY (ZHANGZHOU) BATH PRODUCTS LTD.