3 PACK HAND SANITIZER- alcohol gel Double Body Biotech Limited

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 75%

Purpose

Antiseptic

Use

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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

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

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 Posion Control Center right away.

Directions

Wet hands thoroughly with gel, rub vigorously and allow to dry without wiping. Children under 6 years of age should be supervised when using this product.

Other information

Store below 106 F (41C).

Inactive ingredients

water,
glycereth-26,
aloe barbadensis extract
carbomer,
sodium hydroxide,
may contain: red 33, red 4, yellow 5, blue 1, red 40

Label

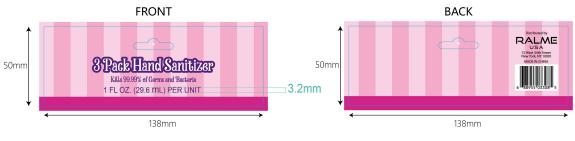
PRODUCT NAME: 3 PACK HAND SANITIZER

RANGE: 29.6ML



防水胶贴(water proof label): 22 x 44mm

560EA-0675PRODUCT NAME: 3 PACK HAND SANITIZER RANGE: 29.6ML



双面印刷卡纸:138 x 50mm

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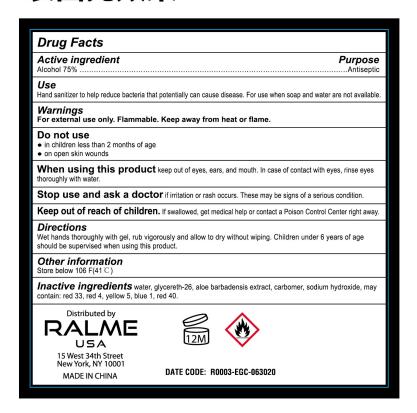
防水胶贴(water proof label): 22 x 44mm

PRODUCT NAME: HAND SANITIZER

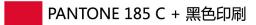
RANGE: 3X30ML

GLOSSY FINISH

表面光效果



防水胶贴(water proof label): 100 x 100mm



PRODUCT NAME: 3 PACK HAND SANITIZER

RANGE: 29.6ML



防水胶贴(water proof label): 22 x 44mm

3 PACK HAND SANITIZER alcohol gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:74657-003

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Panta	Λŧ	Δ	minic	tration

TOPICAL

Active	Ingredient/Active	Moiety
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Ingredient Name		Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
GLYCERETH-26 (UNII: NNE56F2N14)			
ALOE (UNII: V5VD430 YW9)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)			
WATER (UNII: 059QF0KO0R)			
D&C RED NO.33 (UNI: 9DBA0SBB0L)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			

Packaging

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ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 N	DC:74657-003-01	29.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/06/2020	

Marketing Information

Mar Keting Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/06/2020	

Labeler - Double Body Biotech Limited (560246954)

Establishment

Listablishment				
Name	Address	ID/FEI	Business Operations	
Double Body Biotech Limited		560246954	manufacture(74657-003)	

Double Body Biotech Limited Revised: 7/2020