ALOBON SANITIZERSPRAY- sanitizer spray spray Shantou Wanbang Cosmetic 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

Alcohol 75%

Purpose

Antiseptic

Use

For decreasing bacteria on the skin

Warnings

- For external use only
- Flammable, keep away from fire and flame
- Dose not contain grain alcohol; do not drink. If taken internally will produce serious gastric disturbances.

When using this product

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- Aviod the eyes and mucous membrances
- In the case of eyes and mucous membrances contact, rinse area thoroughly with water

Stop use and ask a doctor if

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- Condition worsens
- Redness or irritation develops
- If condition persists for more than 3 days

Keep out of reach of children

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If swallowed contact a doctor or Poison Control Center immediately.

Directions

- Place enough Product in your palm to cover hands and rub hands together briskyly until dry.
- Supervise children in the use of this product.
- In the case of eye contact, rinse eyes thoroughly with water.

Other information

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

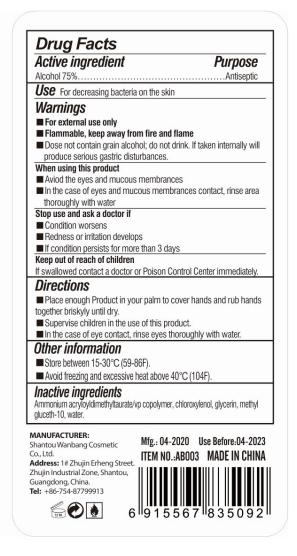
Inactive ingredients

Ammonium acryloyldimethyltaurate/vp copolymer, chloroxylenol, glycerin, methyl gluceth-10, water.

Package Label/Principal Display Panel

NDC 78692-613-34; 100 mL





ALOBON SANITIZERSPRAY

sanitizer spray spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78692-613
Route of Administration	TOPICAL		

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

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
WATER (UNII: 059QF0KO0R)			
METHYL GLUCETH-10 (UNII: N0 MWT4C7WH)			
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)			
CHLOROXYLENOL (UNII: 0 F32U78 V2Q)			

	Pa	ckaging			
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 N	NDC:78692-613-34	$100\ mL$ in 1 BOTTLE; 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	part333A	09/01/2020		

Labeler - Shantou Wanbang Cosmetic Co., Ltd. (552659581)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shantou Wanbang Cosmetic Co., Ltd.		552659581	manufacture (78692-613)	

Revised: 8/2020 Shantou Wanbang Cosmetic Co., Ltd.