ADVANCED 2 IN 1 HAND SANITIZER- alcohol gel THE SUN 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 70.0%

INACTIVE INGREDIENTS

Water, Carbomer, Aminomethyl Propanol, Aloe Vera Gel, Grapefruit Seed Extract. Green Tea Extract, Butylene Glycol, Disodium EDTA, Phenoxyethanol

PURPOSE

Antiseptic

WARNINGS

For external use only. Flammable. Keep away from heat and flame.

Do not use ■ In children less than 2 months of age. ■ On open skin wounds

When using this product \blacksquare Keep out of eyes, ears, and mouth \blacksquare 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 medical advice is needed, have the product container or label at hand.

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

Uses

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- For children under 6, use only under adult supervision
- Avoid direct inhalation of vapors during application (Headaches and irritation to mucous membranes may occur when directly inhaled.)
- This product is intended only for hand sanitizing.
- To prevent drying out and contamination, close the lid completely after use.
- Keep in their original container not to cause accidents nor diminish the integrity of the product.

■ Do not use over bandages, castings, etc. ad irritation may occur.

Other Information

Other Information

- Store below 43°C (110°F)
- May discolor certain fabrics or surfaces

QUESTIONS

www.onnaturalusa.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





ADVANCED 2 IN 1 HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74923-100
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength	
	Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	350 mL in 500 mL	

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
AminomethylPropanol (UNII: LU49E6626Q)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
CITRUS PARADISI SEED (UNII: 12F08874Y7)			
GREEN TEA LEAF (UNII: W2ZU1RY8B0)			
Butylene Glycol (UNII: 3XUS85K0RA)			
EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM)			
Phenoxyethanol (UNII: HIE492ZZ3T)			

Packaging				
7	# Item Code Package Description		Marketing Start Date	Marketing End Date
	NDC:74923-100- 01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	05/01/2020		

Labeler - THE SUN CO.LTD (694433104)

Registrant - THE SUN CO.LTD (694433104)

Establishment				
Name	Address	ID/FEI	Business Operations	
THE SUN CO.LTD		694433104	manufacture(74923-100)	

Revised: 6/2020 THE SUN CO.LTD