

**CENTROLL NAP ADVANCED HAND SANITIZER GEL- alcohol gel
SUNGIN PHARMA**

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.

[82037-510-01] [Gel] [Bottle+Pump] [500 mL] Centroll NAP ADVANCED Hand Sanitizer Gel

Active ingredient

Alcohol 70 %

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin.

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

keep out of eyes, ears, and mouth. 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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough product on hands to cover all surfaces. Rub hand together until dry.
- Apply it in a well-ventilated area.
- Supervise children under 6 years of age when using this product to avoid swallowing.
- 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 contents from drying out and contaminants from entering, close lid completely after use.
- Keep product in its original container as storing the product in anything other than the original container may result in accidents or cause the integrity of the product to diminish.
- Do not use over bandages, castings, etc. as irritation may occur.

Other information

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

Inactive ingredients

Water, Citric acid, Aminomethylpropanol, Urea, Carbomer homopolymer, Tocopherol acetate, Isopropyl myristate, Glycerin, Dexpanthenol, Glycyrrhiza Glabra (Licorice) Root Extract



Drug Facts	
Active ingredient	Purpose
Alcohol 70 %	Antiseptic
Uses Hand sanitizer to help reduce bacteria on the skin. 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 keep out of eyes, ears, and mouth. 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 swallowed, get medical help or contact a Poison Control Center right away.	
Directions ■ Place enough product on hands to cover all surfaces. Rub hand together until dry. ■ Apply it in a well-ventilated area. ■ Supervise children under 6 years of age when using this product to avoid swallowing. ■ 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 contents from drying out and contaminants from entering, close lid completely after use. ■ Keep product in its original container as storing the product in anything other than the original container may result in accidents or cause the integrity of the product to diminish. ■ Do not use over bandages, castings, etc. as irritation may occur.	
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Manufacturer: SUNGIN PHARMA B, 317, Damsun-ro, Geumseong-myeon, Damyang-gun, Jeollanam-do, 57353, Korea

Lot No. : 1002

Mfg. Date : 2021. 05. 25

Made in Korea



CENTROLL NAP ADVANCED HAND SANITIZER GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82037-510
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
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
UREA (UNII: 8W8T17847W)	
DEXPANTHENOL (UNII: 1O6C93RI7Z)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82037-510-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/22/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/22/2021	

Labeler - SUNGIN PHARMA (688719819)

Establishment

Name	Address	ID/FEI	Business Operations
SUNGIN PHARMA		688719819	manufacture(82037-510)

