

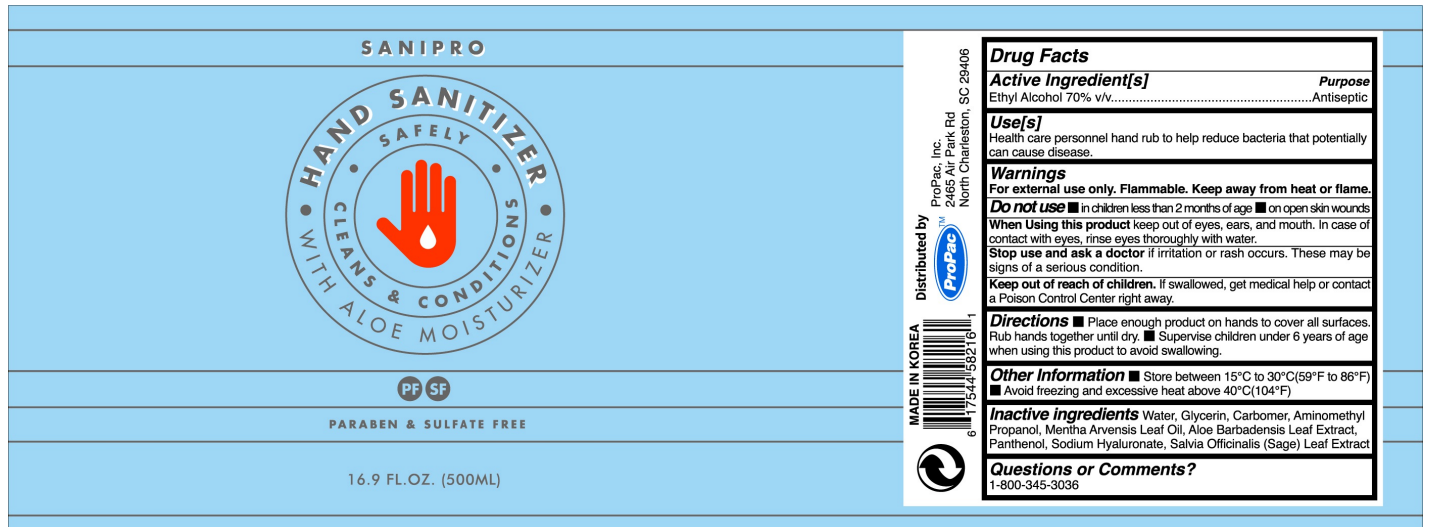
SANIPRO HAND SANITIZER 70 500ML- ethyle alcohol gel
E.CIS COSMETIC.Co.,Ltd

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.

SANIPRO Hand Sanitizer 70% 500ml

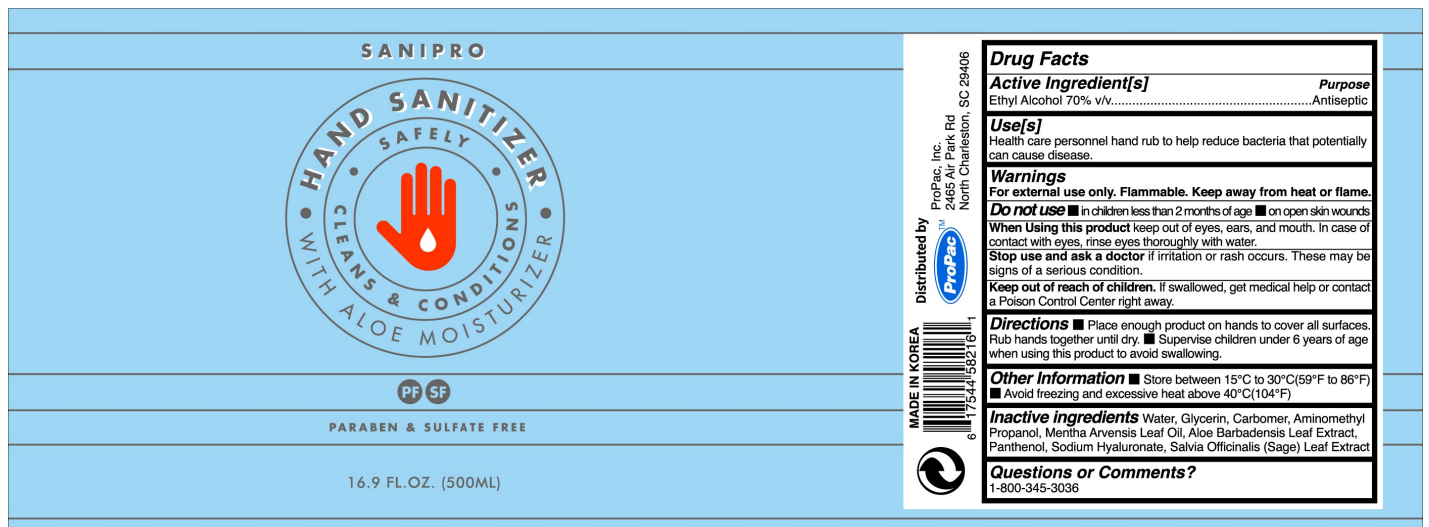
Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic



Purpose

Antiseptic, Hand Sanitizer



Use

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

SANIPRO

PF SF

PARABEN & SULFATE FREE

16.9 FL.OZ. (500ML)

MADE IN KOREA

Distributed by ProPac, Inc. 2465 Air Park Rd North Charleston, SC 29406

6 17544 58216 1

Drug Facts

Active Ingredient[s]	Purpose
Ethyl Alcohol 70% v/v.....	Antiseptic

Use[s]
Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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 hands together until dry. ■ Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information ■ Store between 15°C to 30°C(59°F to 86°F)
■ Avoid freezing and excessive heat above 40°C(104°F)

Inactive ingredients Water, Glycerin, Carbomer, Aminomethyl Propanol, Mentha Arvensis Leaf Oil, Aloe Barbadosis Leaf Extract, Panthenol, Sodium Hyaluronate, Salvia Officinalis (Sage) Leaf Extract

Questions or Comments?
1-800-345-3036

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SAFELY
CLEANS & CONDITIONS
WITH ALOE MOISTURIZER

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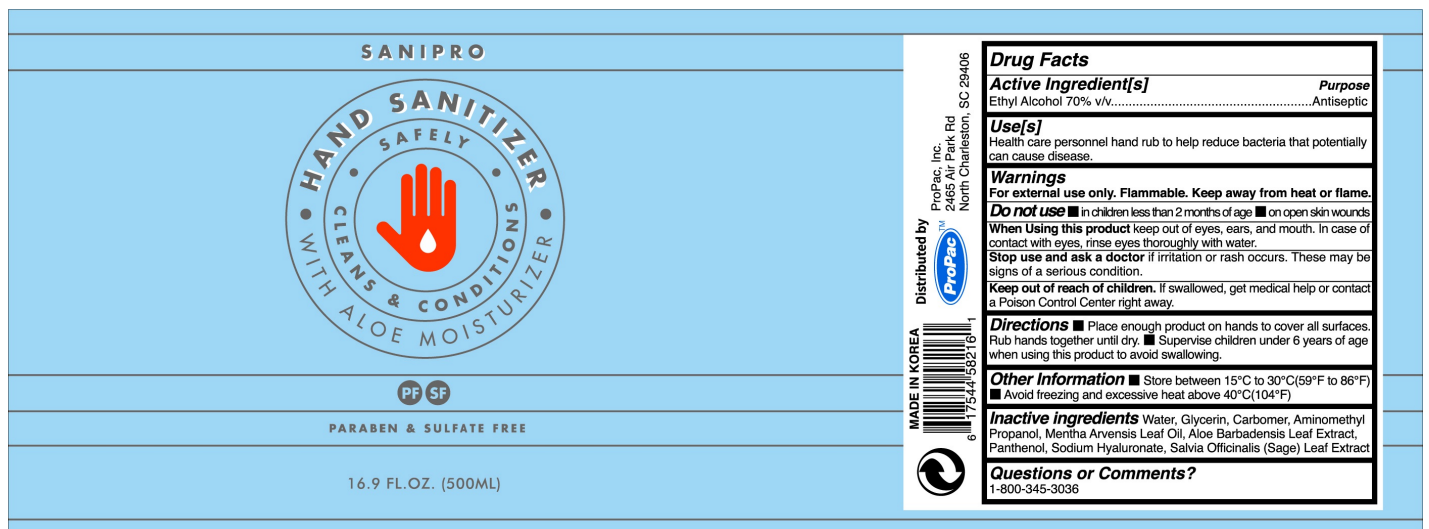
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Package Label - Principal Display Panel

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SANIPRO HAND SANITIZER 70 500ML

ethyle alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73862-215
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
PANTHENOL (UNII: WV9CM0O67Z)	
SAGE (UNII: 065C5D077J)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73862-215-11	24 in 1 CASE	03/30/2020	
1	NDC:73862-215-10	500 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - E.CIS COSMETIC.Co.,Ltd (689846270)

Establishment

Name	Address	ID/FEI	Business Operations
E.CIS COSMETIC.Co.,Ltd		689846270	manufacture(73862-215)

Revised: 6/2020

E.CIS COSMETIC.Co.,Ltd