


SANIPRO HAND SANITIZER 70 250ML- 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 % 250ml

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

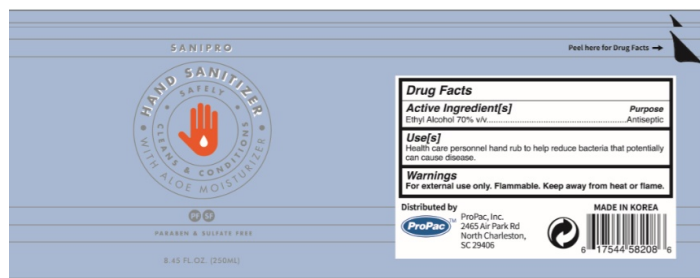


Drug Facts	
Active Ingredient(s) Ethyl Alcohol 70% v/v.	Purpose Antiseptic
Use[s] Health care personnel hand rub to help reduce bacteria that potentially can cause disease.	
Warnings For external use only. Flammable. Keep away from heat or flame.	
Distributed by ProPac, Inc. 2465 Air Park Rd North Charleston, SC 29405	MADE IN KOREA 17544-58206

Drug Facts (continued)
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.
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 Barbadensis Leaf Extract, Panthenol, Sodium Hyaluronate, Salvia Officinalis (Sage) Leaf Extract
Questions or Comments? 1-800-345-3038

Purpose

Antiseptic, Hand Sanitizer

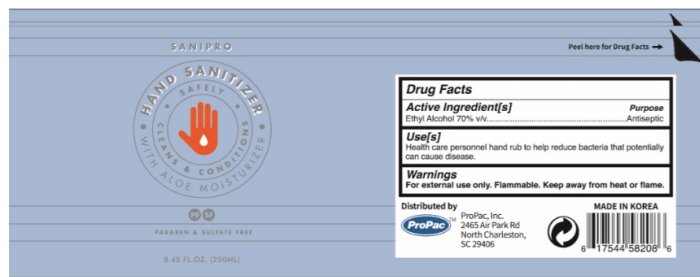


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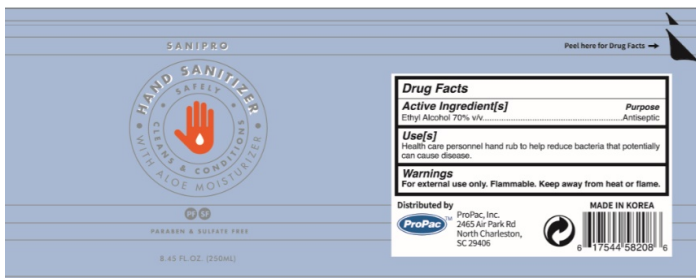
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Package Label - Principal Display Panel

SANIPRO
HAND SANITIZER
WITH ALOE MOISTURIZER

SAFELY
CLEANS & CONDITIONS
PF SF
PARABEN & SULFATE FREE
8.45 FL.OZ. (250ML)



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SANIPRO HAND SANITIZER 70 250ML

ethyle alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73862-214
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)	
PANTHENOL (UNII: WV9CM0O67Z)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SAGE (UNII: 065C5D077J)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73862-214-13	6 in 1 CASE	03/30/2020	
1	NDC:73862-214-12	48 in 1 CASE		
1	NDC:73862-214-11	12 in 1 CASE		
1	NDC:73862-214-10	250 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(73862-214)

Revised: 6/2020

E.CIS COSMETIC.Co.,Ltd