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.

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

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

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

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

Inactive ingredients

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

Package Label - Principal Display Panel

SANIPRO
HAND SANITIZER
WITH ALOE MOISTURIZER
SAFELY
CLEANS & CONDITIONS
PF SF
PARABEN & SULFATE FREE
16.9 FL.OZ. (500ML)

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