PANROSA INSTANT HAND SANITIZER CLEAR- alcohol gel PANROSA ENTERPRISES, INC.

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.

PANROSA Instant Hand Sanitizer Clear

Drug Facts

Active ingredients

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

- hand sanitizer to help decrease bacteria on the skin. when water, soap & towel are not available.
- recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

Do not apply around eyes.

Do not use

in ears & mouth

When using this product.

avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor if

redness or irritation develop and persist for more than 72 hours.

Keep out of reach of children.

Children must be supervised in use of this product.

Directions

- pump as need into your palms and thoroughly spread on both hands.
- rub into skin until dry.

Other information

- store at 20°C (68°F to 77°F).
- may discolor fabrics.

Inactive ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrace, aloe barbadensis leaf juice, tocopheryl acetate

Package Labeling:





Package Labeling: 60ml







product.

Drug Facts (continued) **Directions** • place enough product into your palms and thoroughly spread on both hands. • rub into skin until drv. **Other information ■** store below 110 °F (43°C). ■ may discolor certain fabrics of surfaces *Inactive ingredients* water, carbomer, triethanolamine,

glycerin, propylene glycol,

frágrance, aloé barbadénsis

leaf juice, tocopheryl acetate.

Package Labeling: 1L





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PANROSA ENTERPRISES, INC. 550 Monica Circle Corona, CA 92880 www.panrosa.com Formulated and Developed in the U.S.A. Responsibly sourced and manufactured in China.

PANROSA INSTANT HAND SANITIZER CLEAR

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50302-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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

Inactive Ingredients Strength **Ingredient Name** WATER (UNII: 059QF0KO0R) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) TROLAMINE (UNII: 903K93S3TK) **GLYCERIN** (UNII: PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:50302- 003-00	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/21/2020			
2	NDC:50302- 003-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2020			
3	NDC:50302- 003-11	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	02/21/2020				
final	i'					

Labeler - PANROSA ENTERPRISES, INC. (859957578)

Revised: 10/2021 PANROSA ENTERPRISES, INC.