

INTELLIWRIST PERSONAL HAND SANITIZER- topical antiseptic liquid
Intelwrist, LLC

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.

Intelliwrist Personal Hand Sanitizer

Active Ingredient(s)

Active Ingredient(s)

Ethyl alcohol 70% v/v.....

Purpose

Purpose

Antiseptic

Use

Use

Hand sanitizer to help reduce bacteria on the skin that could cause disease.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

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

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

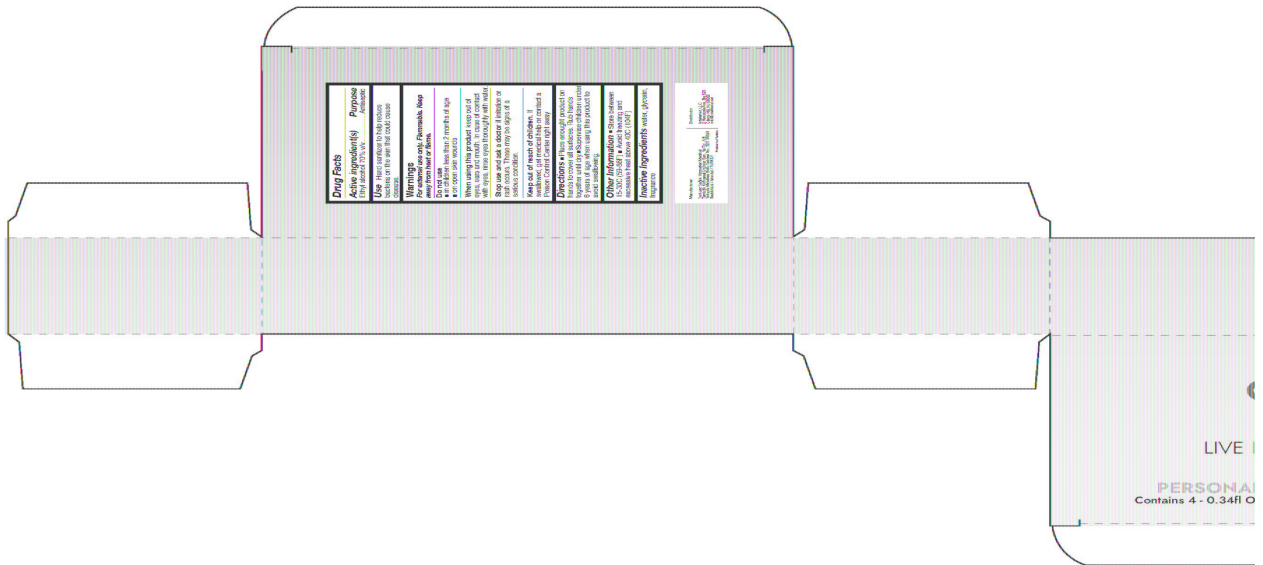
Other Information

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

Inactive Ingredients

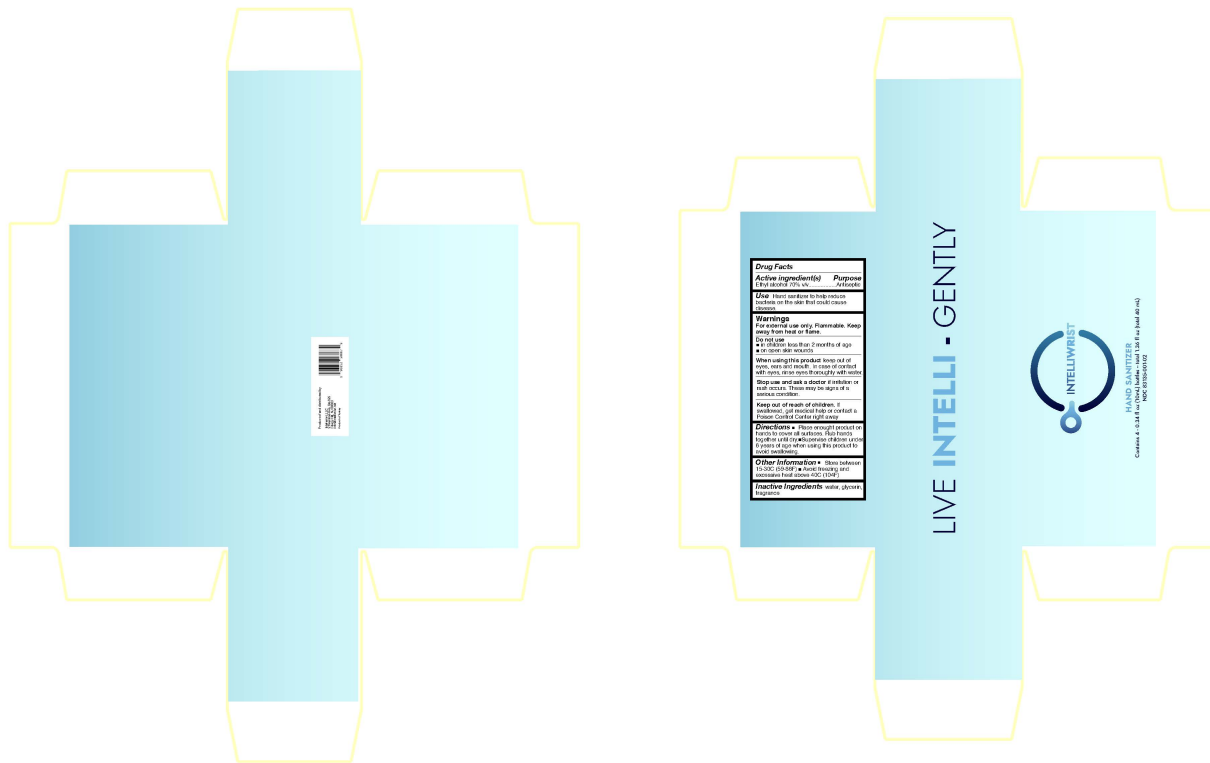
Inactive Ingredients

water, glycerin, fragrance



83135-001-01

83135-001-02



INTELLIWRIST PERSONAL HAND SANITIZER

topical antiseptic liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83135-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	28 mL in 40 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

FRAGRANCE 13576 (UNII: 5EM498GW35)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83135-001-01	40 mL in 1 KIT; Type 0: Not a Combination Product	11/30/2022	
2	NDC:83135-001-02	40 mL in 1 BOX; Type 0: Not a Combination Product	11/30/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/30/2022	

Labeler - Intelwrist, LLC (008557500)

Revised: 1/2023

Intelwrist, LLC