HAND SANITIZER UNSCENTED- is opropyl alcohol spray TAICORP ENTERPRISES LIMITED

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.

Active Ingredient

Isopropyl alcohol 70% v/v

Purpose

Antiseptic

Use

To help reduce bacteria on the skin.

Warnings

- For external use only.
- Flammability warning. Keep away from heat or flame.

Do not useon children less than 2 years of age (unless directed by a doctor).

When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and consult a doctor if irritation develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Dispense enough product in your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.

Inactive ingredients

Aloe Barbandesis Leaf Extract, Glycerine, Water.

Questions?

Contact Scentuals at +1877-770-7590 Mon-Fri 9am - 4pm PST.

Package Label - Principal Display Panel

Drug Facts

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100% NATURAL



UNSCENTED SPRAY



ISOPROPYL ALCOHOL ANTISEPTIC 70%



120ml / 4fl.oz

Easy-to-use spray with no sticky residue

- Aloe & glycerine hydrates skin
- 70% alcohol keeps hands clean
- Unscented & gentle on skin

NPN: 80097896







www.scentuals.com

Manufactured by: Taicorp Enterprises Limited

Made in Canada / Fait au Canada 86 North Bend Street Unit #108, Coquitlam BC, V3K 6H1 +1877-770-7590



HAND SANITIZER UNSCENTED

isopropyl alcohol spray

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:79883-007 Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL UNII:ND2M416302) ISOPROPYL ALCOHOL ISOPROPYL ALCOHOL in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				

P	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:79883-007- 01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					
2	NDC:79883-007- 02	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					
3	NDC:79883-007- 03	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					
4	NDC:79883-007- 04	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					
5	NDC:79883-007- 05	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					
6	NDC:79883-007- 06	$3785\ mL$ in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020					

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	07/01/2020				

Labeler - TAICORP ENTERPRISES LIMITED (202875738)

Establishment							
Name	Address	ID/FEI	Business Operations				
TAICORP ENTERPRISES LIMITED		202875738	manufacture(79883-007)				

Revised: 8/2020 TAICORP ENTERPRISES LIMITED