SMART HAND SANITIZER- alcohol gel Fleetwood Jamaica 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.

Smart Hand Sanitiser

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

a. Alcohol (ethanol SDA 40B 190 PF) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.

b. Glycerol (1.45% v/v).

c. REVERSE OSMOSIS water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol (Ethanol)70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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

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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

Glycerin, Acrylate copolymer, PEG-40 Hydrogenated Castor Oil, Triethanolamine, reverse osmosis water

240 mL NDC 77883-000-00



Product Informa	tion						
Product T ype		HUMAN OTC DRUG	Item Cod	ode (Source) N		NDC:77883-000	
Route of Administra	ition	TOPICAL					
Active Ingredien	t/Active Moi	ety					
Ingredient Name				Basis of Strength		strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)				AL	COHOL	$0.73\ mL$ in $1\ mL$	
GLYCERIN (UNII: PDC6A3C0OX)					0.02 mL in 1 mL		
Inactive Ingredients Ingredient Name					Strength		
WATER (UNII: 059QF					0.02 mL in 1 mL		
WATER (UNII: 059QF		Package Description			0.02 mL in 1 mL 0.245 mL in 1 mL	Marketing End Dat	
WATER (UNII: 059QF Packaging # Item Code	DKOOR)			Mar	0.02 mL in 1 mL 0.245 mL in 1 mL		
WATER (UNII: 059QF Image: Character of the system	0KO0R) 240 mL in 1 BO 3875 mL in 1 BO	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combina	on Product tion Product	Mar 05/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date		
WATER (UNII: 059QF Packaging # Item Code 1 NDC:77883-000-01 2 NDC:77883-000-01 3 NDC:77883-000-02	240 mL in 1 BO 3875 mL in 1 BO 60 mL in 1 BOT	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combina TLE; Type 0: Not a Combinatic	on Product tion Product n Product	Mar 0 5/19 0 5/19 0 5/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date /2020 /2020		
WATER (UNII: 059QF Packaging # Item Code 1 NDC:77883-000-01 3 NDC:77883-000-02 4 NDC:77883-000-03	240 mL in 1 BO 3875 mL in 1 BO 60 mL in 1 BOT 120 mL in 1 BO	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combinat TLE; Type 0: Not a Combinatic TTLE; Type 0: Not a Combinati	on Product tion Product n Product on Product	Mar 0 5/19 0 5/19 0 5/19 0 5/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date /2020 /2020 /2020		
WATER (UNII: 059QF P Kaging # Item Code 0 NDC:77883-000-01 NDC:77883-000-02 NDC:77883-000-02 NDC:77883-000-03	240 mL in 1 BO 3875 mL in 1 BO 60 mL in 1 BOT 120 mL in 1 BO	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combina TLE; Type 0: Not a Combinatic	on Product tion Product n Product on Product	Mar 0 5/19 0 5/19 0 5/19 0 5/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date /2020 /2020		
WATER (UNII: 059QF P Kaging # Item Code 1 NDC:77883-000-01 3 NDC:77883-000-02 4 NDC:77883-000-03 5 NDC:77883-000-04	240 mL in 1 BO 3875 mL in 1 BO 60 mL in 1 BOT 120 mL in 1 BO 480 mL in 1 BO	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combinat TLE; Type 0: Not a Combinatic TTLE; Type 0: Not a Combinati	on Product tion Product n Product on Product	Mar 0 5/19 0 5/19 0 5/19 0 5/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date /2020 /2020 /2020		
WATER (UNII: 059QF P Kaging # Item Code 1 NDC:77883-000-01 3 NDC:77883-000-02 4 NDC:77883-000-03	240 mL in 1 BO 3875 mL in 1 BO 60 mL in 1 BOT 120 mL in 1 BO 480 mL in 1 BO	Package Description TTLE; Type 0: Not a Combinati DTTLE; Type 0: Not a Combinat TLE; Type 0: Not a Combinatic TTLE; Type 0: Not a Combinati	on Product tion Product n Product on Product on Product	Mar 05/19 05/19 05/19 05/19 05/19	0.02 mL in 1 mL 0.245 mL in 1 mL keting Start Date /2020 /2020 /2020		

Labeler - Fleetwood Jamaica Limited (816428652)

Establishment								
Name	Address	ID/FEI	Business Operations					
Fleetwood Jamaica Limited		816428652	manufacture(77883-000)					

Revised: 6/2020

Fleetwood Jamaica Limited