HAND SANITIZER- alcohol gel
Mr Knox LLP.

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.

Hand Sanitizer Gel

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antimicrobial

Purpose

Antimicrobial, Hand Sanitizer

Use

For handwashing to decrease bacteria on the skin.

Recommended for repeated use.

Warnings

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

When using this product keep away from eyes. In case of eye contact, rinse eyes with water.

Stop use and ask a doctor if redness and irritation develops. If issues persist, seek medical attention.

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

Stop use and ask a doctor if redness and irritation develops. If issues persist, seek medical attention.

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

Directions

- Place enough product in your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Other information

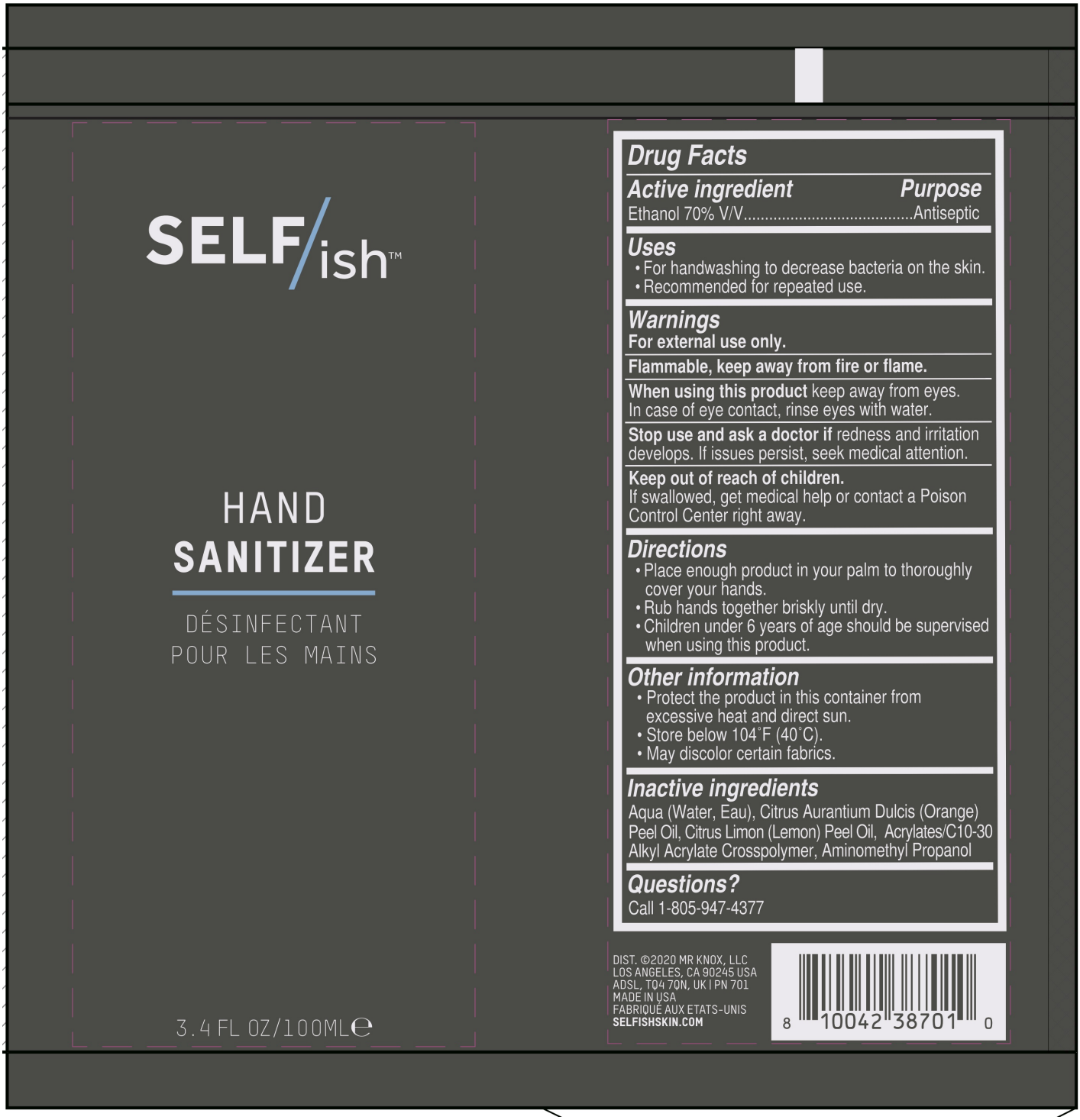
- Protect the product in this container from excessive hear and direct sun.
- Store below 104F (40C).
- May discolor certain fabrics.

Inactive ingredients

Aqua (Water, Eau), Citrus Aurantium Dulcis (Orange) Peel Oil, Citrus Limon (Lemon) Peel Oil, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol

Package Label - Principal Display Panel

100 mL NDC: 78831-701-01



HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78831-701
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**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78831-701-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Mr Knox LLP. (117547167)

Revised: 6/2020 Mr Knox LLP.