

UNSCENTED- hand sanitizer spray
Bell International Laboratories, 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.

Touchland Unscented Hand Sanitizer Spray

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

To decrease bacteria on the skin

Recommended for repeated use

Warnings

For external use only

Flammable, keep away from fire or flame

Do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Avoid contact with broken skin

Stop use and ask a doctor

- if irritation and redness develop
- Condition persists for more than 72 hours

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

Children under 6 years of age should be supervised when using this product

Rub hands together briskly until dry

Other Information

Protect the product in this container from excessive heat and direct sun

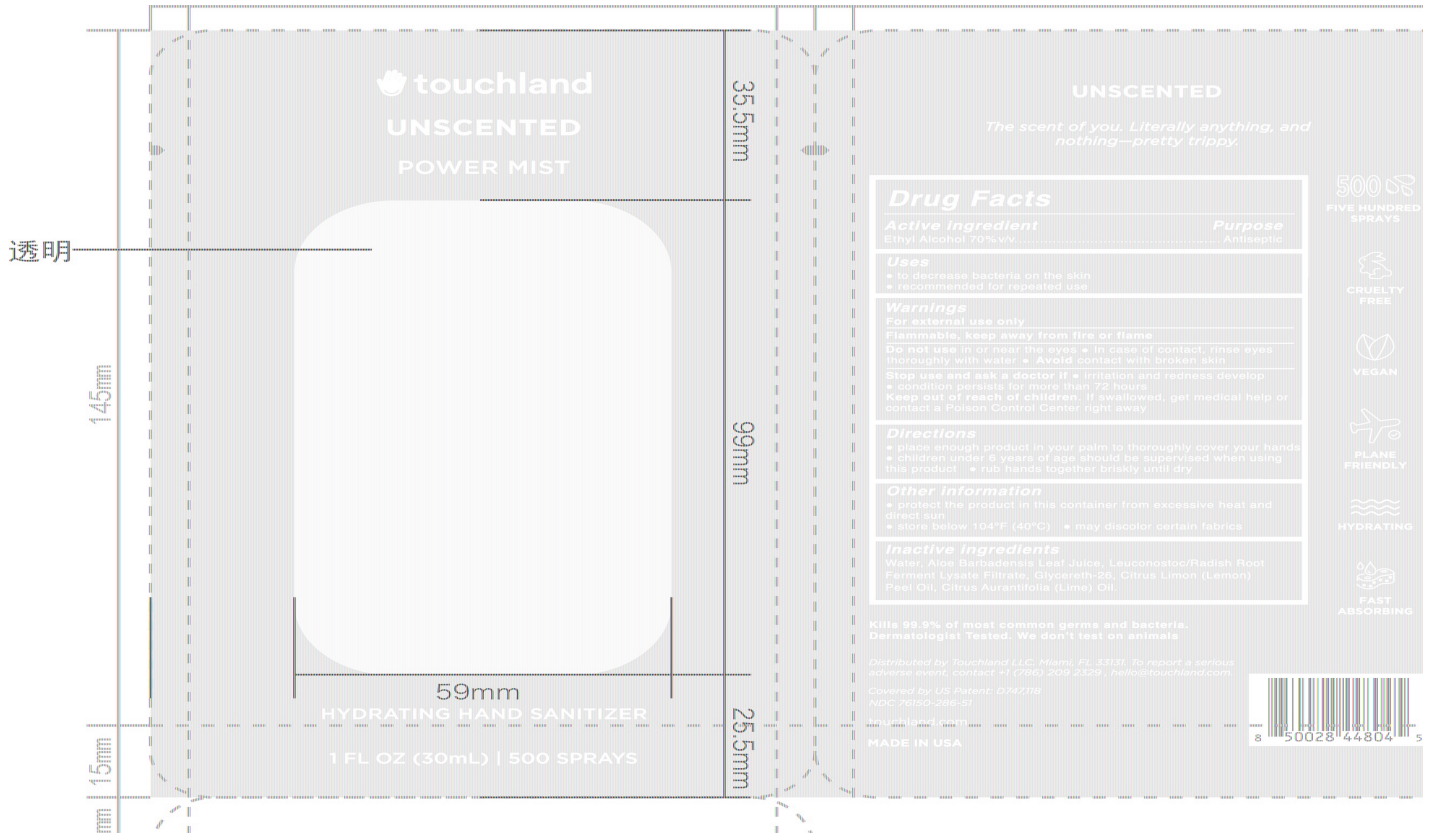
Store below 104F (40C)

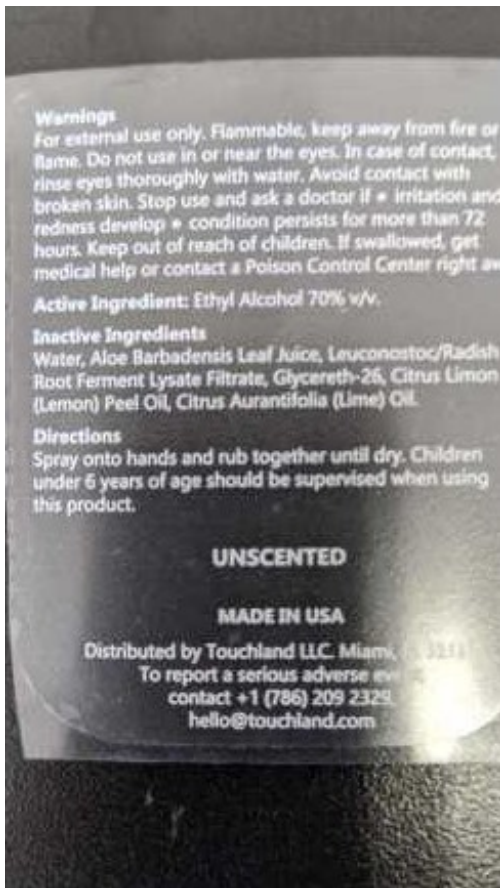
May discolor certain fabrics

Inactive Ingredients

Water, Aloe Barbadensis Leaf Juice, Leuconostoc/Radish Root Ferment Lysate Filtrate, Glycereth-26, Citrus Limon (Lemon) Peel Oil, Citrus Aurantifolia (Lime) Oil

label





UNSCENTED

hand sanitizer spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76150-286
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
GLYCERETH-26 (UNII: NNE56F2N14)	
WATER (UNII: 059QF0KO0R)	
LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458)	
LEMON OIL (UNII: I9GRO824LL)	
LIME OIL (UNII: UZH29XGA8G)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76150-286-51	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2021	

Labeler - Bell International Laboratories, Inc. (967781555)

Revised: 8/2021

Bell International Laboratories, Inc.