

**BLUE SANDALWOOD- 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.*

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**Touchland Blue Sandalwood 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

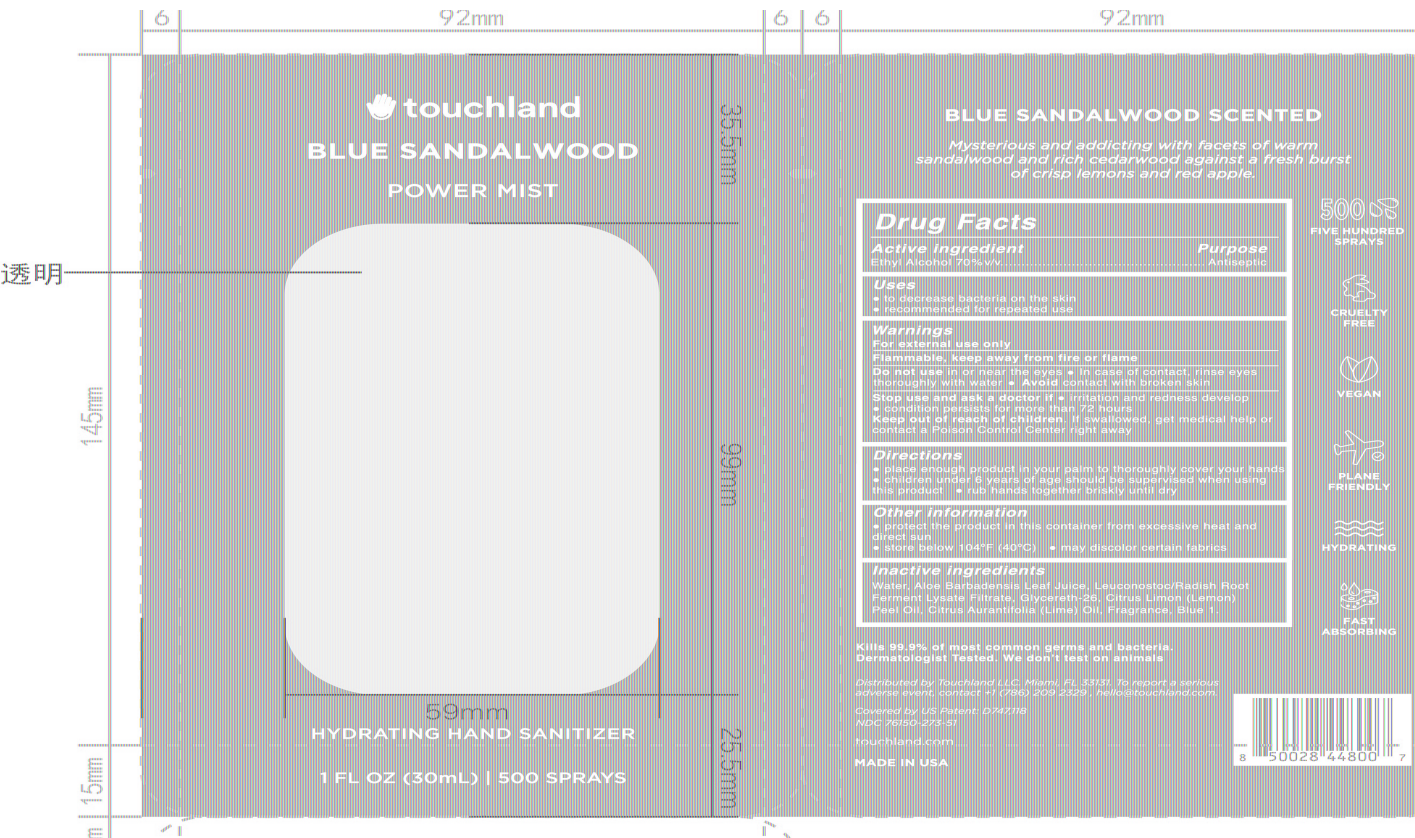
Store below 104F (40C)

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, Fragrance, blue 1

pouch label



back label



## BLUE SANDALWOOD

hand sanitizer spray

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76150-273
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
<b>GLYCERETH-26</b> (UNII: NNE56F2N14)	
<b>LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE</b> (UNII: D2QHA03458)	
<b>LEMON OIL</b> (UNII: I9GRO824LL)	
<b>LIME OIL</b> (UNII: UZH29XGA8G)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76150-273-51	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