

BLUZEN HAND SANITIZER- ethyl alcohol spray
Remcoda, Llc

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.

BLUZEN HAND SANITIZER

Drug Facts

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic

Uses

- hand sanitizer to help reduce bacteria on skin

Warnings

Flammable, keep away from fire/flame

For external use only

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water • do not inhale or ingest

Stop use and ask a doctor if

irritation and redness develop

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

Directions

- wet hands thoroughly with product and allow to dry without wiping
- supervise children under 6 years of age when using this product to avoid swallowing

Other information

- Store below 104°F (40°C)

Inactive ingredients

water, aloe barbadensis leaf juice, glycerin

Questions?

1-800-777-1603

MOISTURIZING FORMULA WITH ALOE VERA

UNSCENTED

KILLS 99.9% OF GERMS

SHOPBLUZEN.COM

Distributed by: Remcoda, LLC, New York, NY, 10018

SULFATE FREE * PARABEN FREE * VEGAN, CRUELTY FREE * MADE IN THE USA

Packaging

The image shows the front of a hand sanitizer bottle. The main text reads "HAND SANITIZER" in large letters, followed by "MOISTURIZING FORMULA WITH ALOE VERA". Below that is a circular seal with "UNSCENTED" inside. The brand name "BLUZEN" is prominently displayed, with "KILLS 99.9% OF GERMS" underneath. At the bottom, it says "2 FL. OZ (60 ML)" and "SHOPBLUZEN.COM". On the right side, there is a "Drug Facts" panel with sections for Active ingredient, Uses, Warnings, Directions, and Inactive ingredients. A vertical banner on the far right states "MADE IN THE USA" and lists product attributes: "SULFATE FREE * PARABEN FREE * VEGAN, CRUELTY FREE".

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BLUZEN HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79200-200
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: 059QF0K00R)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79200-200-02	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/30/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/30/2020		

Labeler - Remcoda, Llc (117130169)

Revised: 6/2020

Remcoda, Llc