BLUZEN HAND SANITIZER WINTERMINT- ethyl alcohol gel 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 GEL-WINTERMINT

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

- place enough product in your palm to thoroughly cover your hands
- rub hands together briskly until dry
- supervise children under 6 years of age when using this product to avoid swallowing.

Other information

■ Store below 110°F (43°C)

Inactive ingredients

glycerin, hydrogen peroxide, purified water, polyethylene glycol, carbomer copolymer, fragrance *Questions?* **1-800-777-1603**

CLEANSING FORMULA

WINTERMINT

KILLS 99.9% OF GERMS
MADE IN THE USA
SULFATE FREE PARABEN FREE
VEGAN, CRUELTY FREE
SHOPBLUZEN.COM

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

Packaging



BLUZEN HAND SANITIZER WINTERMINT ethyl alcohol gel **Product Information** Product Type HUMAN OTC DRUG Item Code (Source) NDC:79200-507 **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		
GLYCERIN (UNII: PDC6A3C0OX)			
HYDROGEN PERO XIDE (UNII: BBX060AN9V)			
WATER (UNII: 059QF0KO0R)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79200-507- 16	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/09/2020	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	09/09/2020				

Labeler - Remcoda, Llc (117130169)

Revised: 9/2020 Remcoda, Llc