

BLOODY MARYS WICKED HAND SANITIZER- ethyl alcohol liquid
Bobbie Weiner Enterprises, L.L.c.

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.

BLOODY MARY'S® WICKED HAND SANITIZER

Drug Facts

Active Ingredient[s]

Ethyl Alcohol 80% v/v

Purpose

Antiseptic

Use[s]

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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

Do not use

- In children less than 2 months of age.
- On open skin wounds.

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Place enough product on hands to cover the surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

Inactive Ingredients Purified water USP, Glycerin, Hydrogen Peroxide

Complies with FDA standards According to WHO recommendation

Bobbie Weiner Enterprises LLC

12355 NE 13th Avenue

Suite 400 B

North Miami, FL 33161

MADE IN THE USA

WWW.DEARBLOODYMARY.COM

Packaging



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 NDC CODE: 76799-101-12
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16 FL OZ. **MADE IN USA**

0 85000 98354 6

**WICKED
HAND SANITIZER**
16 FL OZ.

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DRUG FACTS LABEL

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BLOODY MARYS WICKED HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76799-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76799-101-10	59 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/28/2020	
2	NDC:76799-101-11	118 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/28/2020	
3	NDC:76799-101-12	473 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/28/2020	

Labeler - Bobbie Weiner Enterprises, L.L.c. (014620327)

Revised: 5/2020

Bobbie Weiner Enterprises, L.L.c.