HAND SANITIZER HAND RX- ethyl alcohol liquid
Blue Cross 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.

Hand Rx Hand Sanitizer

Active ingredients purpose

Ethyl Alcohol 62% Antimicrobial

Kills over 99% of most common germs

Keep out of reach of children

Keep out of eyes.In case of eye contact immediately flush eyes with water then contact physician.

Warning: flammable. Keep away from flame. If swallowed, contact physician. For external use only. Do not store above 110 degrees farenheit.

directions:

put a thumbnail size amount in your palm and rub your hands together briskly until dry

Children under 6 years of age should be supervised when using Hand Rx.

Inactive Ingredients: Water, Glycerin, Propylene Glycol, Carbomer

Hand Rx

Hand Sanitizer

Kills 99.99% of germs

8 Fl Oz. (236 ml)

8 OZ Hand Rx™ Hand Sanitizer

Kills Over 99% of Most Common Germs within seconds

Hand Rx allows you to wash your hands whenever conventional soap and water use is not available. Use as an aid to reduce exposure to infectious germs and to supplement daily handwashing. Helps meet U.S. Government standards by protecting against exposure to bloodborne diseases. Nontoxic-For External Use Only.

Keep bottle of *Hand Rx ™* in

- **Kitchen**
- **Medical Offices**
- **Nurseries**
- **Automobile**
- **Bathrooms**
- **Workplace**

Drug Facts

Active Ingredients*.....*Purpose

Ethyl Alcohol 62%.....Antimicrobial

Directions*:** • Put a thumbnail size amount in your palm and rub your hands together briskly until dry. • Children under 6 years of age should be supervised when using ***Hand Rx ™

***Inactive Ingredients*:** Water, Glycerin, Propylene Glycol, Carbomer.

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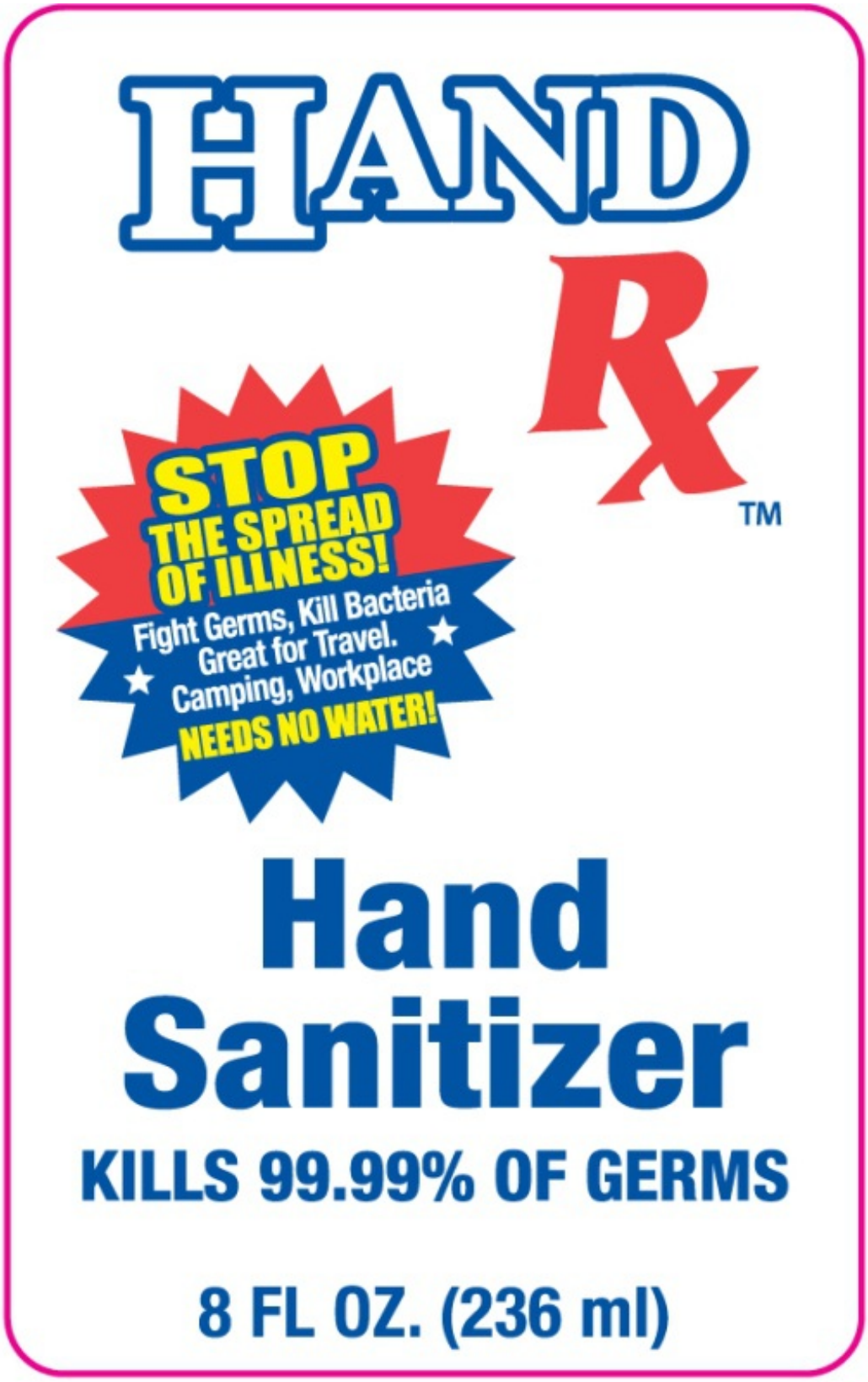

HAND SANITIZER HAND RX

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22431-581

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22431-581-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/12/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/12/2010	

Labeler - Blue Cross Laboratories, Inc. (008298879)

Registrant - Blue Cross Laboratories, Inc. (008298879)

Establishment

Name	Address	ID/FEI	Business Operations
Blue Cross Laboratories, Inc.		008298879	manufacture(22431-581)

Revised: 1/2017 Blue Cross Laboratories, Inc.