HAND RX HAND SANITIZER- alcohol solution
Blue Cross Laboratories

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 Ingredient(s)

Alcohol 62% v/v. Purpose: Antiseptic

Antiseptic

Use

- to reduce bacteria on the skin that could cause disease.
- recommended for repeat use.

Warnings

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

Do not use

avoid contact with broken skin

do not inhale or ingest

When using this product

keep out of eyes. In case of contact with eyes, flush thoroughly with water.
Stop use and ask a doctor if

skin irritation develops

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.
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105ºF
- may discolor some fabrics
- harmful to wood finishes and plastics

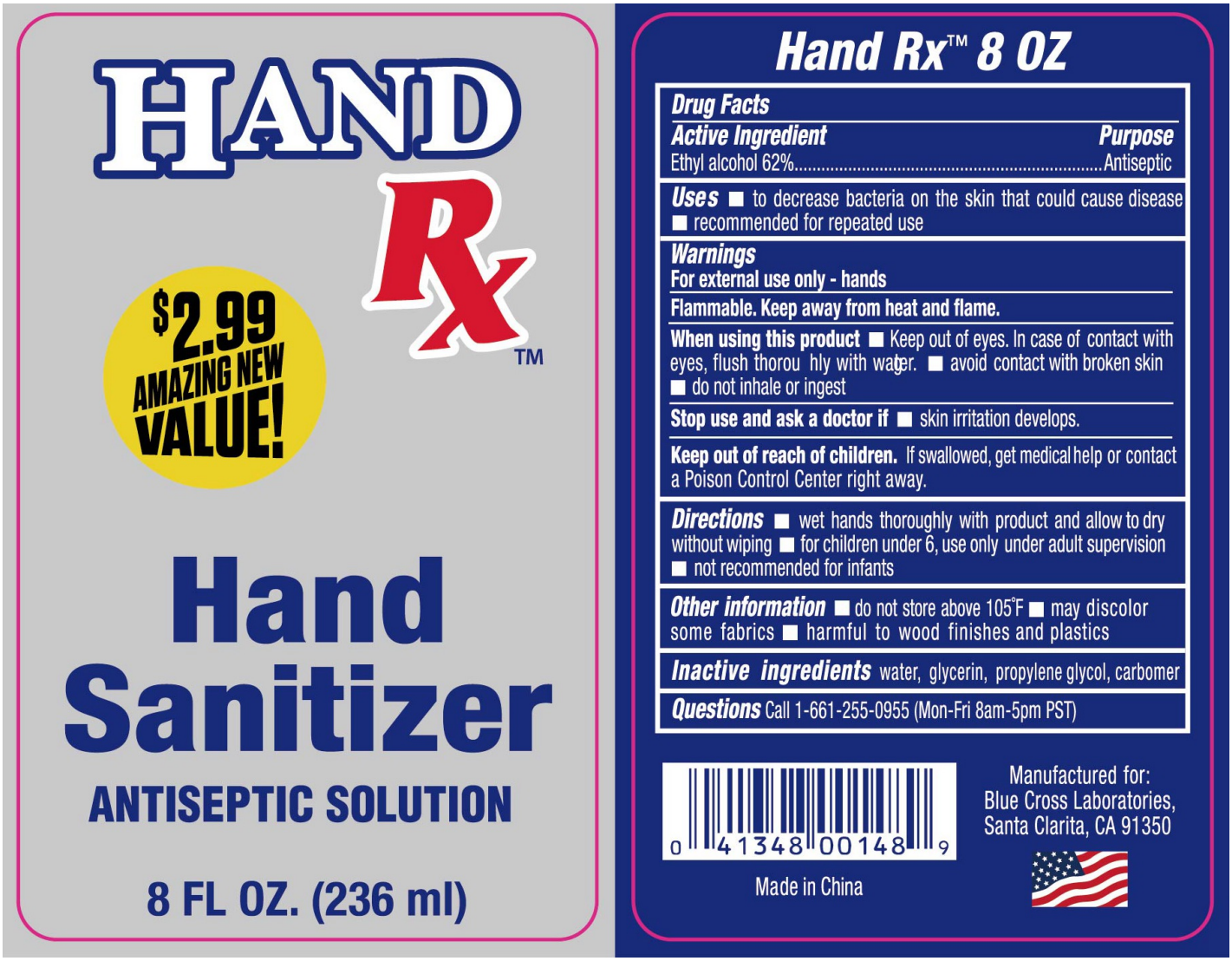
Inactive ingredients

water, glycerin, propylene glycol, carbomer

Package Label - Principal Display Panel

Hand Rx Hand Sanitizer Antiseptic Solution

8 FL OZ. (236 ml)



HAND RX HAND SANITIZER

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22431-221
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22431-221-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/11/2020	

Labeler - Blue Cross Laboratories (008298879)

Establishment

Name	Address	ID/FEI	Business Operations
Ningbo Liyuan Daily Chemical Products Co., Ltd		530766098	manufacture(22431-221)

Revised: 5/2020 Blue Cross Laboratories