CROSSCO- is opropyl alcohol solution Max Chemical, 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.

CROSSCO® ANTISEPTIC Topical Solution

Drug Facts

Active ingredient

Isopropyl alcohol 70%

Purpose

Antiseptic

Uses

- hand sanitizer to decrease bacteria on the skin
- recommended for repeated use
- for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame For external use only

Do not use

- in children less than 2 months of age
- on open skin wounds

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

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

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 between 15-30°C (59-86°F)
- avoid freezing and excessive heat above 40°C (104°F)
- do not use if the cap is broken

Inactive ingredients

water

Questions?

+1-787-765-6100

You may also report serious side effects to this phone number. Mon-Fri 9:00 AM - 5:00 PM

Antiseptic Hand Rub Non-sterile Solution

Distributed by: MAX CHEMICAL, INC.

La Brisa#6, Urb.Sabana Llana, San Juan, PR, 00924

Packaging



70% **ISOPROPYL** ALCOHOL ANTISEPTIC

Topical Solution



Antiseptic Hand Rub

Non-sterile Solution

16 fl oz (1 pt) (473 mL)

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Lot No:

Expiry:

BLANCO GROUP

DE210-16

CROSSCO

isopropyl alcohol solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74213-382

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
ISOPROPYL A	ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -	ISOPROPYL	70 mL

UNII:ND2M416302)	ALCOHOL	in 100 mL
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Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74213-382-08	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	
2	NDC:74213-382-10	296 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	
3	NDC:74213-382-17	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	
4	NDC:74213-382-04	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/17/2020	

Labeler - Max Chemical, Inc. (118159136)

Establishment			
Name	Address	ID/FEI	Business Operations
Max Chemical, Inc.		118 159 136	manufacture(74213-382)

Revised: 4/2020 Max Chemical, Inc.