

ISOPROPYL ALCOHOL 50%- isopropyl alcohol 50% solution
Hydrox Laboratories

Isopropyl Rubbing Alcohol 50%

Drug Facts

Active Ingredient

Isopropyl Alcohol 50%

Purpose:

First Aid Antiseptic

Uses first aid to help prevent the risk of infection in: minor cuts, scrapes, burns.

Warnings For external use only

Flammable

- Keep away from fire or flame, heat, spark, electrical. Flash point 72°F.
- do not use with eletocautery procedures.

Ask a doctor before use if you have deep or puncture wounds, animal bites or serious burns.

When using this product

- do not get into eyes
- do not apply over large areas of the body
- do not use longer than 1 week unless directed by a doctor

Stop use and ask a doctor if condition persists or gets worse

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

<Directions clean the affected area. apply 1 to 3 times daily.

Other information does not contain, nor is intended as a substitue for grain or ethyl alcohol. will produce serious gastric disturbances if taken internally.

Inactive Ingredient purified water

Principal Display Panel

Hydrox

Isopropyl Rubbing Alcohol

A cooling and refreshing and massaging compound

50%

First Aid Antiseptic

TAMPER EVIDENT CAP FOR YOUR PROTECTION. IF RING-BAND IS DETACHED FROM CAP OR MISSING, DO NOT USE.

WARNING: FLAMMABLE!

Hydrox Laboratories

Elgin, IL 60123

The image shows the principal display panel for Hydrox Isopropyl Alcohol 50%. The top left features the Hydrox logo with a crest and the text 'OPTIMALES' and 'TRADE MARK SINCE 1913'. The brand name 'Hydrox' is prominently displayed in a large, white, serif font. Below it, 'ISOPROPYL ALCOHOL' and '50%' are written in large, bold, blue sans-serif fonts. A red 'Warning Flammable' symbol is located to the right of the brand name. The NDC number '10565-003-99' is printed below the warning. A barcode with the number '0 21599 41283 8' is positioned vertically on the right side. A tamper-evident cap warning box is located at the bottom left, and a 'FIRST AID ANTISEPTIC Made in USA' label with a small American flag is at the bottom center. The bottom of the panel features the manufacturer's name 'Mfg. by Hydrox LABORATORIES' and contact information 'Elgin, IL 60123 • www.hydroxlabs.com'. On the right side, there is a 'Drug Facts' table with sections for Active ingredient, Purpose, Uses, Warnings, Directions, and Inactive ingredient.

Drug Facts	
Active ingredient	Purpose
Isopropyl alcohol 50%	First Aid Antiseptic
Uses • helps prevent the risk of infection in: • minor cuts • scrapes • burns	
Warnings For external use only	
Flammable • keep away from fire or flame, heat, spark, electrical. Flash point 72°F. • do not use with electrocautery procedures	
Ask a doctor before use if you have deep or puncture wounds, animal bites or serious burns	
When using this product • do not get into eyes • do not apply over large areas of the body • do not use longer than 1 week unless directed by a doctor	
Stop use and ask a doctor if condition persists or gets worse	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions • clean the affected area • apply 1 to 3 times daily • may be covered with a sterile bandage • if bandaged, let dry first	
Other information • does not contain, nor is intended as a substitute for grain or ethyl alcohol • will produce serious gastric disturbances if taken internally	
Inactive ingredient	purified water

Net Content: ONE GALLON (3.8 L)

Mfg. by **Hydrox** LABORATORIES
Elgin, IL 60123 • www.hydroxlabs.com

1-07-A0033 Reorder
A0033

ISOPROPYL ALCOHOL 50%

isopropyl alcohol 50% solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10565-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	50 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10565-003-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2019	
2	NDC:10565-003-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2019	
3	NDC:10565-003-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2019	
4	NDC:10565-003-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2019	
5	NDC:10565-003-99	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/11/2019	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	manufacture(10565-003) , label(10565-003) , pack(10565-003)

Revised: 10/2023

Hydrox Laboratories