

ISOPROPYL ALCOHOL- isopropyl alcohol solution
Hydrox Laboratories

Isopropyl Rubbing Alcohol 70% USP

Drug Facts

Active Ingredient

Isopropyl Alcohol 70%

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, USP

A cooling and refreshing and massaging compound

70%

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

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0 21599 12344 4
1-07-D0022B

Hydrox
TRADEMARK SINCE 1913

NDC 10565-002-32

ISOPROPYL RUBBING ALCOHOL, USP

A cooling and refreshing rubbing and massaging compound.

70%

FIRST AID ANTISEPTIC

WARNING: FLAMMABLE!

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32 FL. OZ. (946 mL)

Hydrox Laboratories
Elgin, IL 60123

1-07-D0024F

ISOPROPYL ALCOHOL

isopropyl alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10565-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	1 mL in 100 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10565-002-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2017	
2	NDC:10565-002-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/05/2019	
3	NDC:10565-002-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2017	
4	NDC:10565-002-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2017	
5	NDC:10565-002-99	3800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/05/2017	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	manufacture(10565-002) , label(10565-002) , pack(10565-002)