ISOPROPYL RUBBING ALCOHOL 50% - isopropyl alcohol liquid North & South Wholesalers LLC

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.

Isopropyl Rubbing Alcohol 50%

Active Ingredients (by volume)

Isopropyl alcohol (50% conc.)

Purpose

First aid antiseptic

Uses

• first aid to help prevent the risk of infection in minor cuts, scrapes and burns

Warnings

For external use only; flammable, keep away from fire or flame, heat, spark, electrical

Ask a doctor before use if you have

• deep punctured 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 one week unless directed by a doctor

Stop using this product if

• condition persists or gets worse

Keep this and all drugs out of the reach of children

In case of accidental ingestion, seek professional assistance or contact a Poison control center (1-800-222-1212) immediately

Directions

- clean effected area
- apply small amount of this product on the area 1-3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first

Other information

- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol, will produce serious gastric disturbance if taken internally

Inactive Ingredient

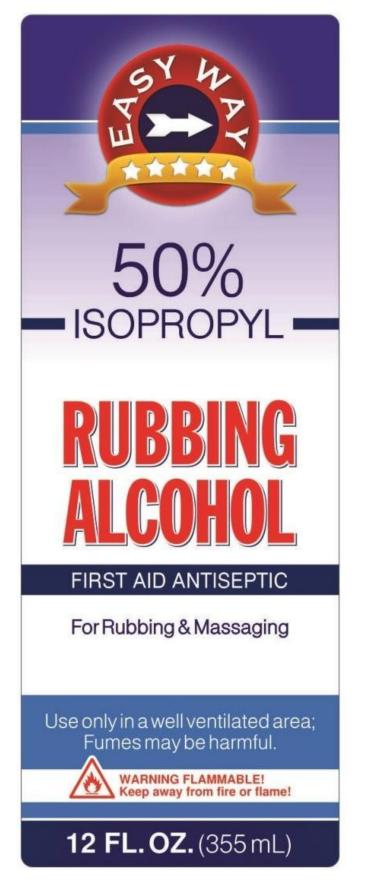
purified water

PRINCIPAL DISPLAY PANEL

ISOPROPYL RUBBING ALCOHOL 50%

FIRST AID ANTISEPTIC

12 FL.OZ (355 mL)



ISOPROPYL RUBBING ALCOHOL 50%

isopropyl alcohol liquid

| 50% by | rl rubbing alcohol volume | | |
|--|--|-----------------------|-------------------------------------|
| Drug | Facts | | |
| and the second division of the second divisio | gredients (by volun Icohol | | Purpos first aid antisept |
| Uses ■ fi scrapes ar | st aid to help prevent the d burns | risk of infec | tion in minor cut |
| Warnin For externa spark, elec | use only; flammable, keep | away from fire | e or flame, heat, |
| | or before use if you have nctured wounds, animal bite | s or serious bu | rns |
| do not g | g this product et into eyes oply over large areas of the b se longer than one week unl | ody ess directed b | y a doctor |
| Stop using | this product if on persists or gets worse | | |
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| apply si May be | fected area fected area nall amount of this product covered with a sterile band ged, let dry first | | 1-3 times daily |
| store at does no ethyl alc | nformation room temperature t contain, nor is intendec shol. will produce serious gas e Ingredient purifie | tric disturbanc | |
| | R EVIDENT: DO NOT ED SAFETY FOIL IS 390 | BROKENO | |

| Product Inform | ation | | | | | | | |
|--|---|----------------|--------------------------|--------------------|----------------------|-----------------------|-------|-----------------------|
| Product Type | | HUMAN OTC DRUG | Item Coo | Item Code (Source) | | NDC:70201-001 | | |
| Route of Administration | | | TOPICAL | | | | | |
| Active Ingredie | nt/A | ctive Moie | ety | | | | | |
| | | Ing | redient Name | | | Basis of Stre | ength | Strength |
| ISOPROPYL ALCOHOL (UNII: ND2M4 UNII:ND2M416302) | | | | | ISOPROPYL ALCOHOL | | | |
| Inactive Ingred | ients | | | | | | | |
| | | Iı | igredient Name | | | | Stre | ngth |
| Packaging | | | | | | | | |
| # Item Code | | | Package Description | | Ma | rketing Start Date | Μ | larketing End Date |
| 1 NDC:70201-001- 01 | 118 m Produ | | LE, PLASTIC; Type 0: Not | a Combination | | | | |
| 2 NDC:70201-001- 02 | 177 n Produ | | LE, PLASTIC; Type 0: Not | a Combination | | | | |
| 3 NDC:70201-001- 03 | 237 n Produ | | LE, PLASTIC; Type 0: Not | a Combination | | | | |
| 4 NDC:70201-001- 04 | -001- 296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | | | | | | |
| 5 NDC:70201-001- 05 | 01- 355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | | | | | | |
| 6 NDC:70201-001- 06 | 414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | | | | | | |
| 7 NDC:70201-001- 07 | 473 n Produ | | LE, PLASTIC; Type 0: Not | a Combination | | | | |
| | | | | | | | | |
| Marketing In | | | | | | | | |
| Marketing Categ | - | | on Number or Monogra | aph Citation | Marketi | ing Start Date | Mark | keting End Da |
| OTC monograph not | final | part333A | | | 11/03/201 | 5 | | |

Labeler - North & South Wholesalers LLC (004948495)

Registrant - Anicare Pharmaceuticals Pvt. Ltd (916837425)

| Establishment | | | | | | | | | |
|----------------------------------|---------|-----------|----------------------------|--|--|--|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | | | | |
| Anicare Pharmaceuticals Pvt. Ltd | | 916837425 | manufacture(70201-001) | | | | | | |