ALCOHOL- isopropyl alcohol liquid
Dynarex Corporation

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.

Alcohol 70%

Active Ingredient	Purpose
Isopropyl Alcohol 70% v/v	Antiseptic

PURPOSE

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.
- Not for use with electrocautinary devices or procedures
- **Ask a doctor before use if you have** deep puncture wounds, animal bites or serious burns.
- **Stop use and ask a doctor if** condition persist or gets worse.

For rubbing and massaging:

- **Caution:** Do not apply to irritated skinor if excessive irritation develops
- Avoid getting into eyes or mucous membranes

INDICATIONS & USAGE

When using this product:

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

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

IIf swallowed, get medical help or contact a Poison Control Center right away.

DOSAGE & ADMINISTRATION

DIRECTIONS:

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

OTHER INFORMATION

- Store at room temperature: 15 deg C to 30 deg C 59 deg F to 86 deg F
- avoid excessive heat
- Does not contain, nor is it intended as a substitute for grain or ethyl alcohol
- Will produce serious gastric disturbances if taken internally

INACTIVE INGREDIENT

Inactive Ingredient

- Water

PRINCIPAL DISPLAY PANEL

DYNAREX 70% ISOPROPYL ALCOHOL

70 IPA.jpg

ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-304
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-304-01	473 mL in 1 BOTTLE		
2	NDC:67777-304-02	295 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/29/2014	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Establishment

Name	Address	ID/FEI	Business Operations
Blossom Pharmaceuticals		677381470	manufacture(67777-304)

Revised: 1/2014 Dynarex Corporation