ALCOHOL PREP- isopropyl alcohol swab
Dynarex Corporation

1113, 1114, 1116, 1113UB-24, 1113UB-10, 1113-50 Sterile Alcohol Prep Pads

Active Ingredient

Isopropyl Alcohol 70%

Purpose

Antiseptic

Use(s)

- First aid to help prevent infection in minor cuts, scrapes, and burns
- For preparation of the skin prior to an injection

Warnings

For External Use only

Flammable. Keep away from fire or flame.

Do not use

- In the eyes or over large areas of the body
- Longer than 1 week unless directed by a doctor
- With electrocautery procedures

Ask a doctor before use if you have

Deep or puncture wounds, animal bites, or serious burns.

Stop use and ask a doctor if

Condition persists for more than 72 hours or gets worse.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

• **When used for first aid:** Clean the affected area. Apply a small amount of this product on the area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.

• **When used to prepare an injection site: A**pply product to the injection site prior to injection.

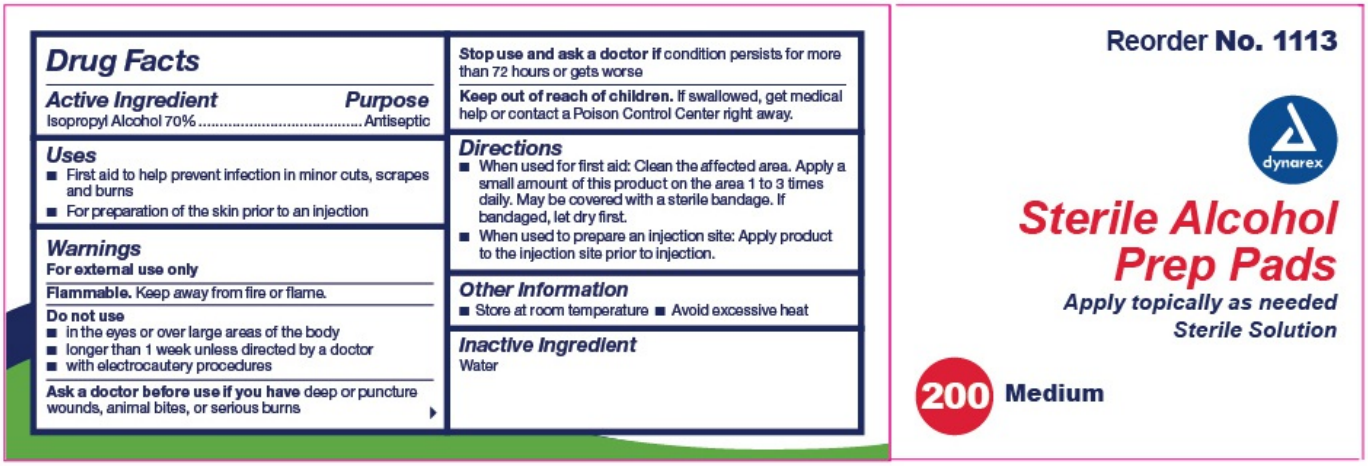
Other Information

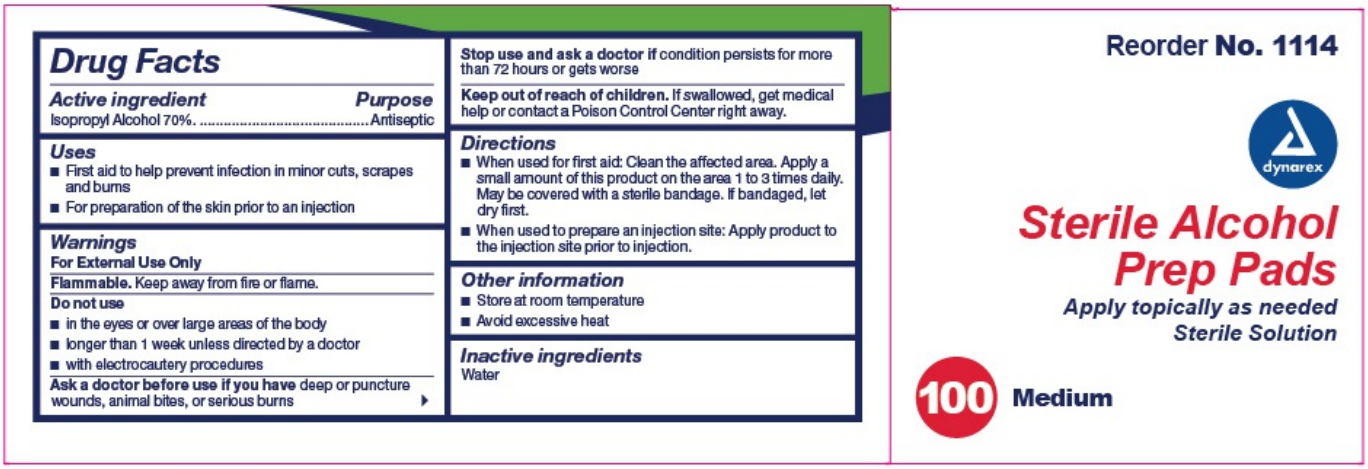
- Store at room temperature
- Avoid excessive heat

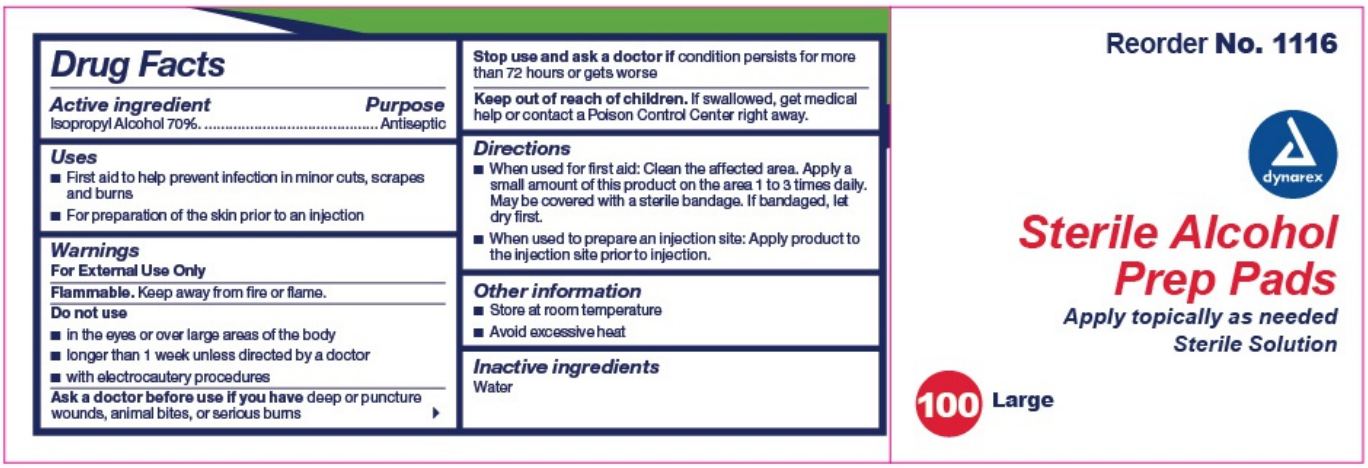
Inactive Ingredient

Water

Label







Label 1113-50

1113-50

Label 1113UB-24

1113UB-24

Label 1113UB-10

1113UB-10

Label 1113-50

1113-50 PHFL010

ALCOHOL PREP

isopropyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-121
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)	

Product Characteristics

Color		**Score**	
Shape	RECTANGLE	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-121-20	20 in 1 CASE	07/01/2010	
1	NDC:67777-121-19	100 in 1 BOX		
1	NDC:67777-121-14	0.55 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:67777-121-18	10 in 1 CASE	07/01/2010	
2	NDC:67777-121-17	100 in 1 BOX		
2	NDC:67777-121-16	0.7 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:67777-121-12	10 in 1 CASE	07/01/2010	
3	NDC:67777-121-11	200 in 1 BOX		
3	NDC:67777-121-13	0.55 mL in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:67777-121-26	40 in 1 CASE	07/01/2010	
4	NDC:67777-121-27	50 in 1 BOX		
4		0.55 mL in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:67777-121-22	100 in 1 CASE	07/01/2010	
5	NDC:67777-121-21	24 in 1 BOX		
5		0.55 mL in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:67777-121-24	100 in 1 CASE	07/01/2010	

6	NDC:67777-121-23	10 in 1 BOX		
6		0.55 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/01/2010	

Labeler - Dynarex Corporation (008124539)

Revised: 12/2025 Dynarex Corporation