

**POVIDONE IODINE PREP PAD- antiseptic swab**  
**Dynarex Corporation**

-----  
**1108 Povidone Iodine Prep Pad NDC 67777-002-01**  
**1108UB-10 Povidone Iodine Prep Pad NDC 67777-002-03**

***Active Ingredient***

Povidone-Iodine USP 10% w/v (9.85% w/w)

***Purpose***

First Aid Antiseptic

***Use(s)***

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

***Warnings***

**For External Use Only**

***Do not use***

- In the eyes or over large areas of the body
- Longer than one week unless directed by a doctor

***Ask a doctor before use if you have***

- Deep or puncture wounds
- Animal bites
- Serious burns

***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 (1-800-222-1222) right away.

***Directions***

- Tear at notch
- Remove prep pad
- 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
- Use only once

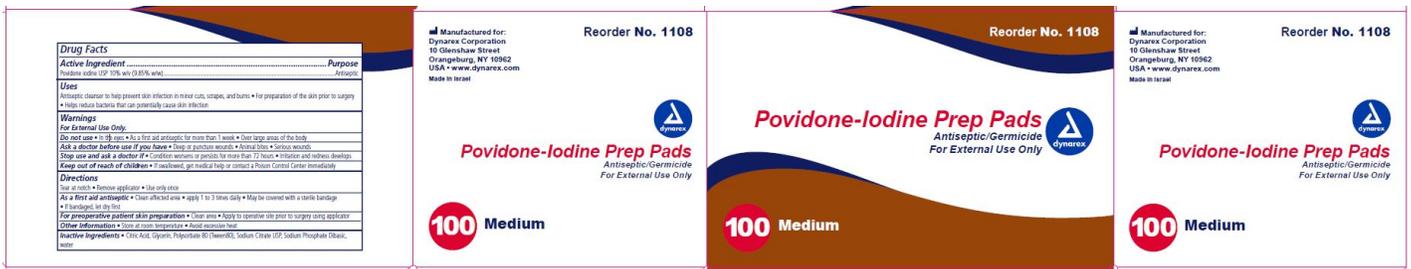
### Other Information

- Store at room temperature 15°-30°C (59°-86°F)
- Flammable. Keep away from fire or flame

### Inactive Ingredients

Anhydrous Citric Acid, Glycerin, Polysorbate 80, Sodium Citrate USP, Sodium Phosphate Dibasic, Water

### Label

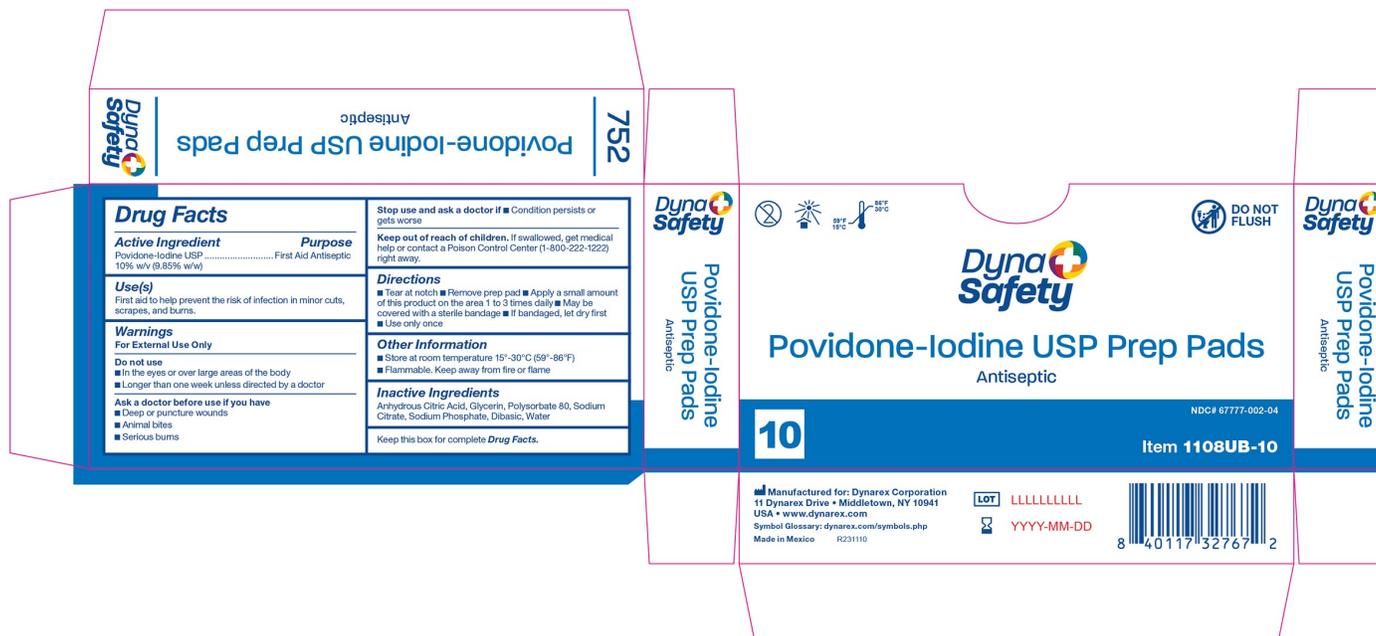


### Label



Povidon Iodine Prep Pad

## Label 1108UB-10



1108UB-10

### POVIDONE IODINE PREP PAD

antiseptic swab

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67777-002
<b>Route of Administration</b>	TOPICAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 g in 100 g

#### Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

#### Product Characteristics

<b>Color</b>		<b>Score</b>	
--------------	--	--------------	--

<b>Shape</b>	RECTANGLE	<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-002-01	1000 in 1 CASE	12/17/2018	
1	NDC:67777-002-02	100 in 1 BOX		
1		0.3 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-002-03	1000 in 1 CASE	12/17/2018	
2	NDC:67777-002-04	10 in 1 BOX		
2		0.3 g in 1 PACKET; 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	12/17/2018	

**Labeler** - Dynarex Corporation (008124539)

Revised: 7/2024

Dynarex Corporation