ALCOHOL PREP PADS WITH PAIN RELIEF - benzocaine swab
McKesson

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.

DRUG FACTS

ACTIVE INGREDIENTS
Benzocaine 6%
Isopropyl alcohol 70% v/v

PURPOSE

Topical Anesthetic

Antiseptic

USES
- For preparation of the skin prior to injection
- For temporary relief of pain and itching associated with minor burns, sunburn, minor skin irritations, or insect bites

WARNINGS

For external use only.

Flammable, keep away from fire or flame.

Do not use
- in the eyes
- with electrocautery procedures

Stop use and ask a doctor
- if irritation and redness develop
- if symptoms persist for more than 72 hours or symptoms clear up and occur within a few days.

Ask a doctor or pharmacist

if you are pregnant or nursing a baby.

Keep out of reach of children

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

DIRECTIONS

Prior to Injection - Adults and children 2 years of age or older:
Apply to the skin just prior to injection. Benzocaine takes a minute to act. for best results, wait briefly and then make injection.

Insect bites, skin irritations, minor burns, sunburn -
Adults and children 2 years of age or older:
Apply to the affected site not more than 3 to 4 times daily.

Children under 2 years of age: Ask a doctor.

OTHER INFORMATION

- Store at room temperature: 15° - 30° C (59° - 86° F)
- Do not use to clean and disinfect medical devices or on hard surfaces

INACTIVE INGREDIENTS
propylene glycol, water

CARTON INFORMATION

sunmark®

NDC 49348-897-39

alcohol prep pads

Antiseptic / Anesthetic
Safe and easy to use
Isopropyl Alcohol 70%

WITH BENZOCAINE FOR PAIN RELIEF
80 INDIVIDUALLY WRAPPED FOIL PACKETS

Another Quality Product Distributed by McKesson
One Post Street, San Francisco, CA 94104
Money Back Guarantee

Visit us at www.sunmarkbrand.com

ALCOHOL PREP PADS WITH PAIN RELIEF

benzocaine swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49348-897
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
benzocaine (UNII: U3RSY48JW5) (benzocaine - UNII:U3RSY48JW5)	benzocaine	0.06 mL
isopropyl alcohol (UNII: ND2M416302) (isopropyl alcohol - UNII:ND2M416302)	isopropyl alcohol	0.70 mL

Inactive Ingredients	
Ingredient Name	**Strength**
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49348-897-39	80 in 1 CARTON		
1		1 in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/19/2010	

Labeler - McKesson (177667227)

Revised: 9/2010 McKesson