

ALCOHOL PREP PADS - benzocaine swab

Target 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.

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

Questions?

Call 1-800-910-6874

PACKAGE INFORMATION

NDC 11673-141-31

alcohol prep pads

with benzocaine
antiseptic/anesthetic
for preparation of the skin
prior to injection

up and up

80 count

80 INDIVIDUALLY WRAPPED PADS

245 10 0155 ID120478

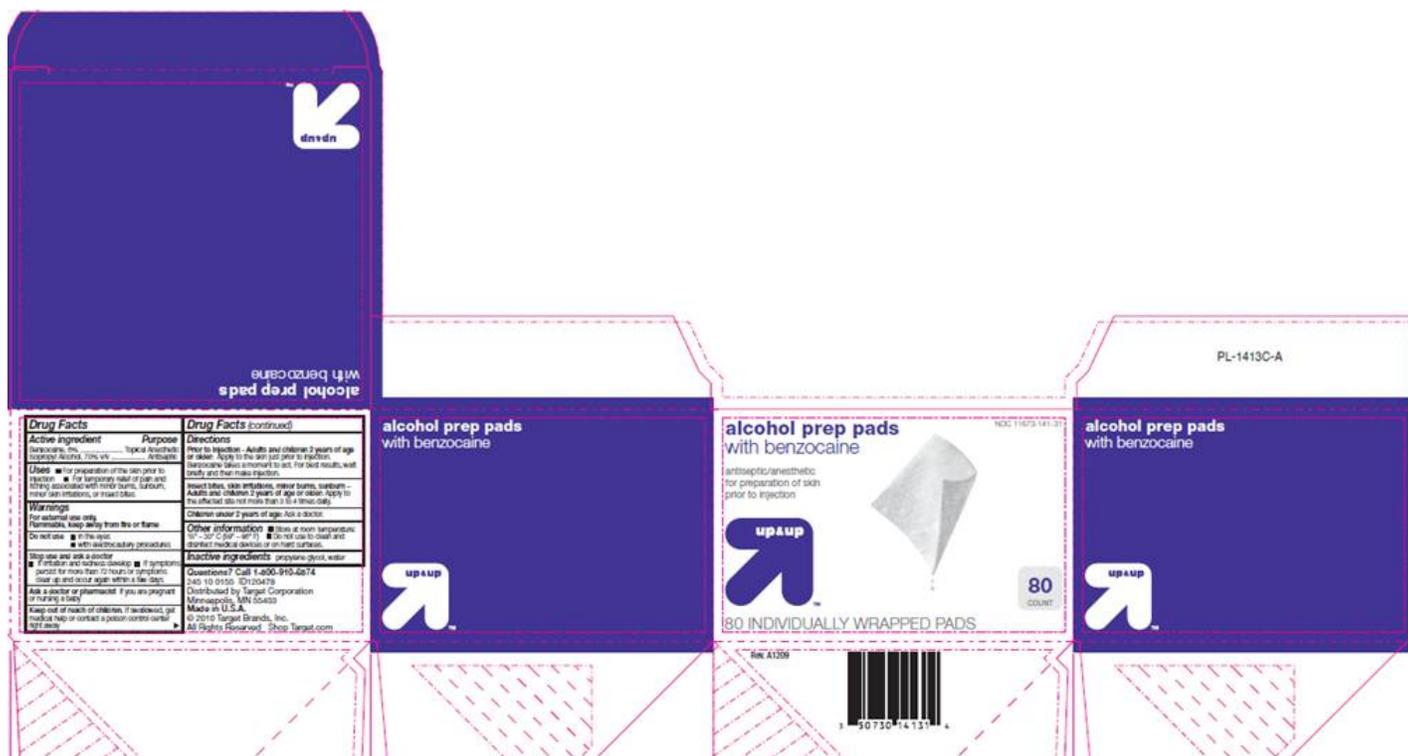
Distributed by Target Corporation

Minneapolis, MN 55403

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ALCOHOL PREP PADS

benzocaine swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-141
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: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-141-31	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 - Target Corporation (006961700)

Revised: 9/2010

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