ALCOHOL PREP PAD-LARGE - isopropyl alcohol swab
Global Biomedical Technologies, LLC

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 PREP PAD-LARGE:
Drug Facts

ACTIVE INGREDIENT:

70% isopropyl alcohol, saturated prep pads.

PURPOSE:

Bandage releasing agent.

USE:

To release bandage from the skin.

WARNINGS:

For external use only: Flammable, keep away from Fire or flame.

DO NOT USE with electrocautery procedures or in the eyes. If contact occurs, flush eyes with water.

STOP USE if irritation and redness develops. If condition persists consult your health care practitioner.

KEEP OUT OF REACH OF CHILDREN. If swallowed get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Wipe vigorously over bandage for several seconds (to release bandage from the skin) and then discard. Store at room temperature (59° - 86°F).

INACTIVE INGREDIENT:

purified water.

Principal Display Panel – Carton Label

Comfort
Release®

Global Biomedical
Technology

FOR
SENSITIVE SKIN

- **Water resistant**
- **Breathable**
- **Long lasting**
- **Painless & trauma-free release***

DERMATOLOGIST
RECOMMENDED

*Apply rubbing alcohol with a cotton ball or an alcohol prep pad to the bandage for painless, trauma-free removal.

10 Comfort Release® Bandages

2” x 4" [5.1 cm x 10.2 cm]

Includes **20 Alcohol Prep Pads** for Removal

ALCOHOL PREP PAD-LARGE

isopropyl alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72727-0001
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:72727-0001-2	20 in 1 BOX	03/20/2019	
1		0.4 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 not final	part333A	03/20/2019	

Labeler - Global Biomedical Technologies, LLC (080609342)

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