

QUICKTOUCH ALCOHOL PREP PAD- alcohol prep pads swab
SLV Pharmaceuticals LLC

QuickTouch Alcohol Prep Pad

Active Ingredient

Isopropyl Alcohol, 70% v/v

Purpose

Antiseptic, Sterile Solution

Uses

Antiseptic cleanser

Kills harmful bacteria and germs

First aid to help prevent infection

Warnings

For External Use Only

Avoid contact with the eyes

If contact occurs, flush eyes with water

Flammable, keep away from fire or flame.

Do Not Use

With electrocautery procedures

In the eyes

Stop Use and ask a doctor if

Irritation and redness develops

If condition persists for more than 72 hours, consult a physician

Discontinue use and consult a healthcare practitioner if

Irritation develops

Keep out of reach of children

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

Directions

Use as part of your daily cleansing routine

May be covered with a sterile bandage

Other Information

Store at room temperature 15°-30°C (59°-86°F)

Avoid excessive heat

Inactive Ingredients

Water

Quick Touch Alcohol Prep Pad



QUICKTOUCH ALCOHOL PREP PAD

alcohol prep pads swab

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:73317-4417 |
| 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 |

Inactive Ingredients

| Ingredient Name | Strength |
|--------------------------|----------|
| WATER (UNII: 059QF0KO0R) | |

Product Characteristics

| Color | Score |
|-------|-------|
| Shape | Size |
| white | |
| | |

| Flavor | | Imprint Code | | |
|------------------------------|--|--|----------------------|--------------------|
| Contains | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:73317-4417-1 | 1 in 1 PACKET; Type 0: Not a Combination Product | 09/16/2024 | |
| Marketing Information | | | | |
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M003 | 09/16/2024 | | |

Labeler - SLV Pharmaceuticals LLC (081225162)

Registrant - SLV Pharmaceuticals LLC (081225162)

Revised: 9/2024

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