CAAT KIT (COVID-19 ANTIBODY ACCELERATED TEST KIT)- is opropyl alcohol Asclemed USA, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

CAAT Kit TM

(COVID-19 Antibody Accelerated Test Kit)

Active Ingredient

Purpose

Isopropyl Alcohol 70% v/v

Antiseptic

Purpose:

Antiseptic

Uses

For first aid to decrease germs in

- minor cuts
- scrapes
- burns

For preparation of the skin prior to injection

Warnings

For external use only

Flammable - keep away from fire or flame

Do not use

with electrocautery procedures

When using this product do not

- get into eyes
- apply over large areas of the body
- in case of deep or puncture wounds, animal bites or serious burns consult a doctor

Stop use and ask a doctor if

- condition persists or gets worse or lasts for more than 72 hours
- do not use longer than 1 week unless directed by a doctor

Keep out of reach of children.

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

Directions

- apply to skin as needed
- discard after single use

Other information

Protect from freezing and avoid excessive heat

Inactive ingredient

Water

NDC: 76420-704-10

CAAT Kit™ COVID-19 Antibody Accelerated Test Kit

Kit Contains

- 10 Test Cassettes with Buffer and Package Insert
- 10 Droppers
- 10 Isopropyl Alcohol 70% Prep Pads
- 10 Single Use Lancets
- 10 Biohazard Specimen Bags
- 10 Adhesive Bandages

Package Contains 10 Single Use Tests

For Professional Use Only

For In Vitro Diagnostic Use

Distributed by:

EnovachemTM
PHARMACEUTICALS
Torrance, CA 90501

NDC: 76420-704-10

CAAT Kit

COVID-19 Antibody Accelerated Test

| K | it | Co | n | ta | ins |
|---|----|----|---|----|-----|
|---|----|----|---|----|-----|

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CAAT KIT (COVID-19 ANTIBODY ACCELERATED TEST KIT)

isopropyl alcohol kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76420-704

Packaging

| l | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|------------------|---|-----------------------------|---------------------------|
| ı | 1 | NDC:76420-704-10 | 1 in 1 POUCH: Type 1: Convenience Kit of Co-Package | 04/13/2020 | |

Quantity of Parts

| Part # | Package Quantity | Total Product Quantity |
|--------|------------------|------------------------|
| Part 1 | 10 POUCH | 50 mL |

Part 1 of 1

ISOPROPYL ALCOHOL

isopropyl alcohol swab

Product Information

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| Active higherinactive wholety | | | |
|--|----------------------|--------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302) | ISOPROPYL ALCOHOL | 70 mL in 100 mL | |

Inactive Ingredients

| 0 | | |
|----------------------|-----------------|----------|
| | Ingredient Name | Strength |
| D (LINII, OFOOEOROD) | | |

WATER (UNII: 059QF0KO0R)

Packaging

| ı | 8 8 | | | |
|---|-------------|--|-----------------------------|--------------------|
| l | # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| l | 1 | 5 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 | 0 1/0 1/20 0 7 | |

Marketing Information

| 8 | | | | | |
|-----|-----------------------|--|----------------------|--------------------|--|
| | Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| | unapproved drug other | | 04/13/2020 | | |
| - 1 | | | | | |

Labeler - Asclemed USA, Inc. (059888437)

Revised: 4/2020 Asclemed USA, Inc.