

4270 FIRST AID KIT- 4270 first aid kit

4401 FIRST AID KIT- 4401 first aid kit

Honeywell Safety Products 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](#).

0498-4270, 4401: First Aid Kit (Amm Inh, PVP wipes, sting relief-SF00001090, SF00001103)

Ammonia Inhalent

Active ingredient (in each ampule)

Ammonia 15%

Ammonia Inhalent

Purpose

Respiratory stimulant

Ammonia Inhalent

Uses

- to prevent or treat fainting

Ammonia Inhalent

Warnings

For external use only

Do not use

- if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

- condition persists

Keep out of reach of children

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

Ammonia Inhalent

Directions

- hold inhalant away from face and crush ampoule between thumb and forefinger at position indicated on sleeve.
- hold near nostrils for inhalation of volatile vapor

Ammonia Inhalent

Other information

- store at room temperature away from light

Ammonia Inhalent

Inactive ingredients

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water

Ammonia Inhalent

Questions

1-800-430-5490

PVP

Active Ingredient

Povidone-iodine solution USP, 10% (equivalent to 1% titratable iodine)

PVP

Purpose

First aid antiseptic

PVP

Uses

- first aid to help prevent the risk of infection in minor cuts, scrapes, and burns

PVP

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body
- on individuals who are allergic or sensitive to iodine

Ask a doctor before use if you have

- deep or puncture wounds,
- animal bites
- serious burns

When using this product

- do not use longer than one week unless directed by a doctor

Stop use and ask a doctor if

- conditions persists or gets worse
- irritation and redness develops

Keep out of reach of children.

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

PVP***Directions***

Reverse cardboard sleeve, then crush at dot between thumb and forefinger. Allow solution to saturate tip and apply solution to injury.

- clean affected area
- apply to affected area 1 to 3 times daily
- may be covered with a sterile bandage
- discard swab after single use

PVP***Other information***

- store at room temperature away from light
- keep from freezing or excessive heat
- do not use if package is torn or open

PVP***Inactive ingredient***

citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

PVP***Questions***

1-800-430-5490

Sting Relief***Active ingredient (in each wipe)***

Ethyl alcohol 50.0% Lidocaine HCl 2.0%

Sting Relief**Purpose**

Antiseptic

Topical pain relief

Sting Relief

Uses

- prevent infection in minor scrapes, and temporary relief of itching of insect bites

Sting Relief

Warnings

For external use only

Flammable, keep away from open fire or flame

Do not use

- over large areas of the body
- in eyes
- over raw or blistered areas

Stop use and ask a doctor

- if conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children

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

Stig Relief

Directions

adults and children 2 years and older: Apply to cleaned affected area not more than 3 times daily.

children under 2 years of age: consult a doctor.

Sting Relief

Inactive ingredients

benzalkonium chloride, menthol, and purified water

Questions or Comments?

1-800-430-5490

4270

SF00001090 Kit Contents

1 1 X 3 PLASTIC 50/BOX

1 WOVEN 2" X 3" 25/BOX

1 KNUCKLE BAND 8 PER
1 AMMONIA INHALANTS 10 PER
1 EYE DRESS PKT W/4 ADH STRIPS
2 INSTANT COLD PACK 4" X 6"
1 FINGERTIP BANDAGE, 10 PER
1 PVP IODINE WIPES 10 PER
2 ADHES TAPE W/P 1"X 2 1/2 YD
2 GAUZE BANDAGE 2"X2 YDS STRETCH GZ
1 FIRST AID GUIDE ASHI
1 GZE PADS STERILE 3"X 3" 10'S
1 CPR FILTERSHIELD 77-100
1 SCISSOR BDGE 4" RED PLS HDL
1 KIT TWEEZER 3 1/2" SLANTED
1 180 EMPTY BLANK NO LOGO
1 POCKET INSERT RED #180 KIT 4R
2 BANDAGE COMP 4" W/TELFAPAD 1
1 LBL STOCK 6-3/8"X4"
1 LBL STOCK 4"X2-7/8"
3 PR LRG NITRILE GLVES ZIP BAG
1 WATER-JEL BURN DRESSING 4 X 4
1 NOX A STING WIPES 10

4401

SF00001103 kit contents

1 1 X 3 PLASTIC 50/BOX
1 WOVEN 2" X 3" 25/BOX
1 KNUCKLE BAND 8 PER
1 AMMONIA INHALANTS 10 PER
1 EYE DRESS PKT W/4 ADH STRIPS
2 INSTANT COLD PACK 4" X 6"
1 FINGERTIP BANDAGE, 10 PER
1 PVP IODINE WIPES 10 PER
2 ADHES TAPE W/P 1"X 2 1/2 YD

2 GAUZE BANDAGE 2"X2 YDS STRETCH GZ

1 FIRST AID GUIDE ASHI

1 GZE PADS STERILE 3"X 3" 10'S

1 CPR FILTERSHIELD 77-100

1 SCISSOR BDGE 4" RED PLS HDL

1 KIT TWEEZER 3 1/2" SLANTED

2 BANDAGE COMP 4" W/TEFPA PAD 1

1 LBL STOCK 6-3/8"X4"

1 LBL STOCK 4"X2-7/8"

3 PR LRG NITRILE GLVES ZIP BAG

1 WATER-JEL BURN DRESSING 4 X 4

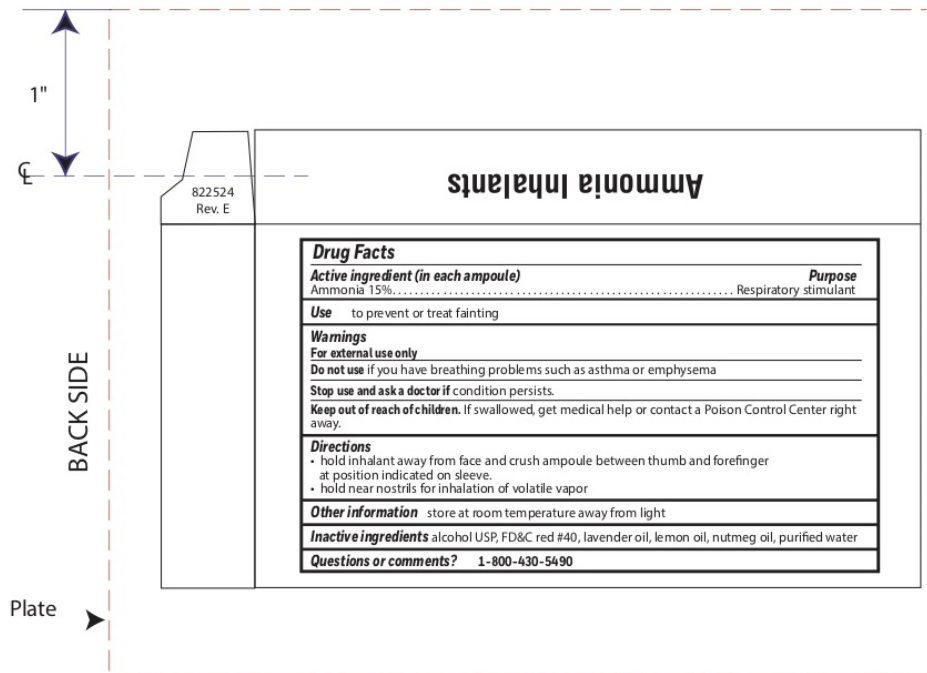
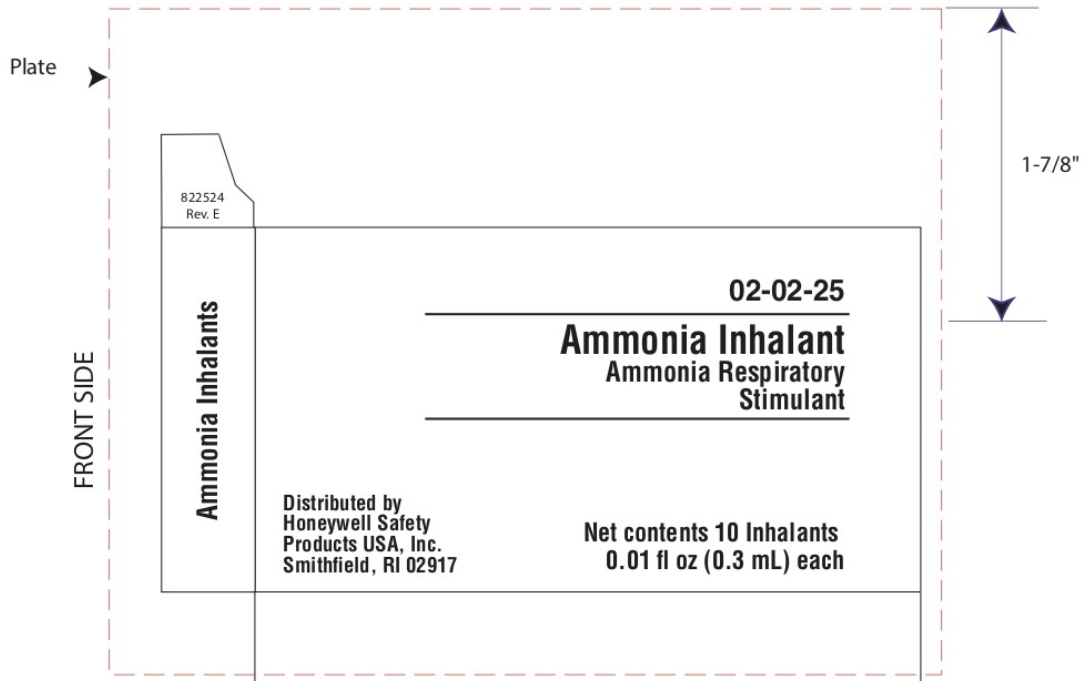
1 POLYBAG 14" X 20" '3 MIL'

1 STING relief WIPES 10

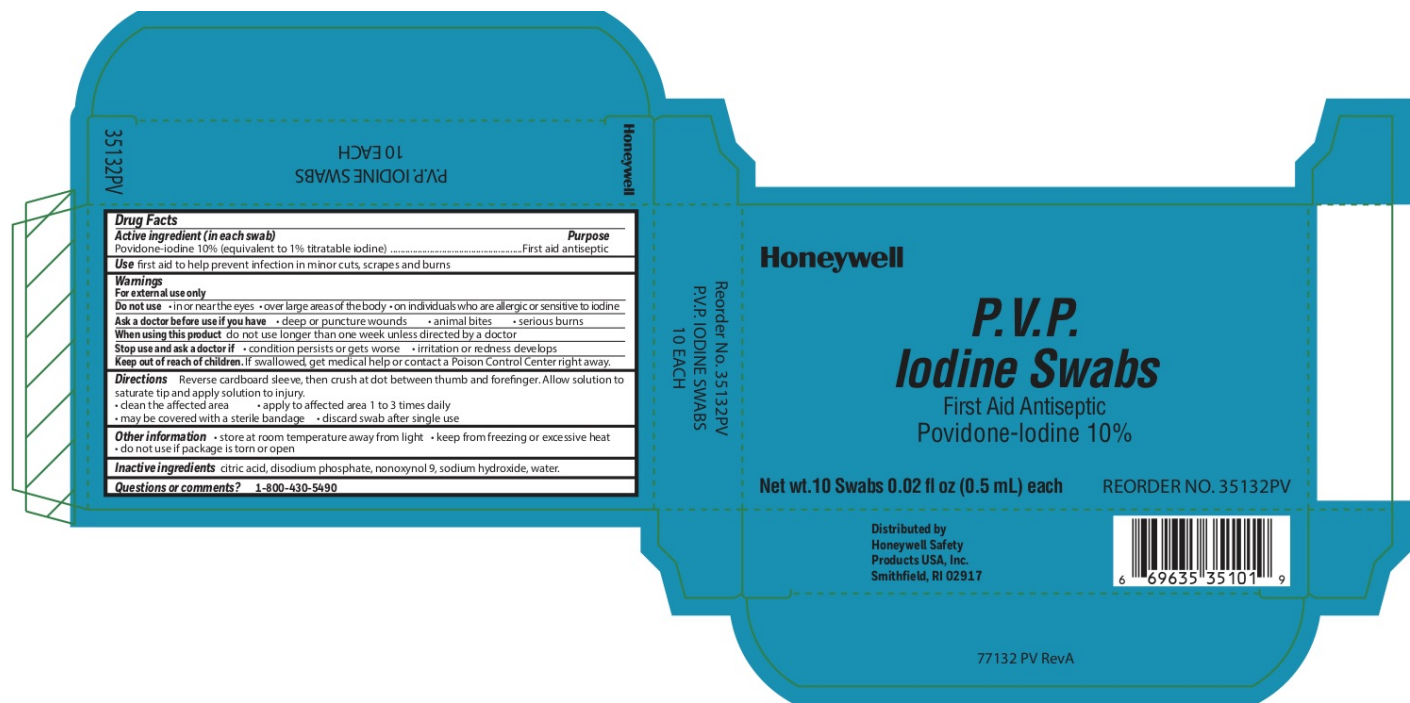
Ammonia Inhalant

Principal Display Panel

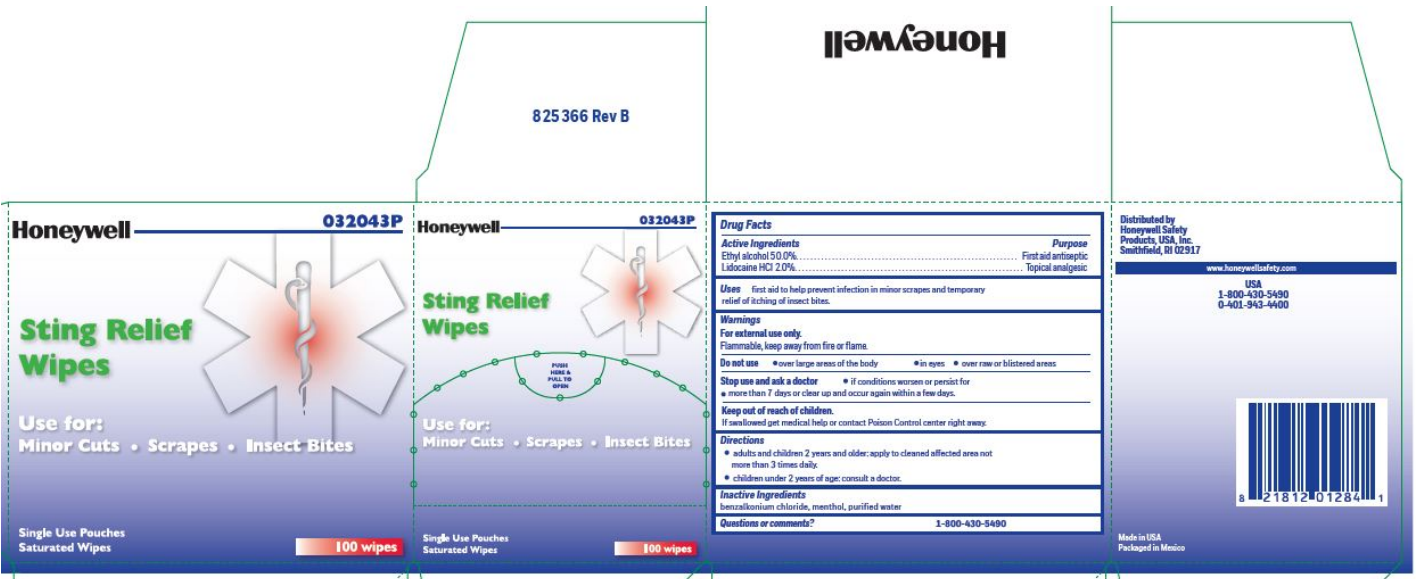
796006 Rev. E Unit Carton Printing Plate for "A" size carton.



Principal Display Panel



Sting Relief
Principal Display Panel



4270 Kit Label
SF00001090



Distributed by Honeywell Safety Products USA, Inc Smithfield, RI 02917

4401 Kit Label
SF00001103

46001363 Rev.C

Prints 3 colors

Black, Red (PMS 186) and Blue (PMS 072)

Refill Information

US Poison Control 1-800-222-1222



Contact your authorized Honeywell Safety Products
Distributor with your refill orders.

Honeywell

www.honeywellsafety.com

| USA | CANADA | EUROPE |
|----------------|----------------|--------------------|
| 1-401-943-4400 | 1-514-351-7233 | +31 (0) 118 656400 |
| 1-800-430-4110 | 1-888-212-7233 | |

46001363 Rev. C

4270 first aid kit kit

Product Information

| | | | |
|--------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0498-4270 |
|--------------|----------------|--------------------|---------------|

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0498-4270-01 | 1 in 1 KIT; Type 0: Not a Combination Product | 03/14/2019 | |

Quantity of Parts

| Part # | Package Quantity | Total Product Quantity |
|--------|------------------|------------------------|
| Part 1 | 10 AMPULE | 3 mL |
| Part 2 | 10 POUCH | 3 mL |
| Part 3 | 10 POUCH | 4 mL |

Part 1 of 3

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

| | |
|-------------------------|--------------------------|
| Item Code (Source) | NDC:0498-3334 |
| Route of Administration | RESPIRATORY (INHALATION) |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-------------------|
| AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X) | AMMONIA | 0.045 g in 0.3 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|----------------------------|----------|
| ALCOHOL (UNII: 3K9958V90M) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0498-3334-00 | 0.3 mL in 1 AMPULE; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 09/18/2018 | |

Part 2 of 3

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

| | |
|-------------------------|---------------|
| Item Code (Source) | NDC:0498-0121 |
| Route of Administration | TOPICAL |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|---------------|
| POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4) | IODINE | 10 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---------------------------------------|----------|
| NONOXYNOL-9 (UNII: 48Q180SH9T) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0121-00 | 0.3 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 |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 09/18/2018 | |

Part 3 of 3

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information

| | |
|--------------------------------|---------------|
| Item Code (Source) | NDC:0498-0733 |
| Route of Administration | TOPICAL |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------------------|-------------------|
| LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE HYDROCHLORIDE ANHYDROUS | 20 mg in 1 mL |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 0.5 mL in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|-----------------|
| MENTHOL (UNII: L7T10EIP3A) | |
| WATER (UNII: 059QF0KO0R) | |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|----------|------------------|--|-----------------------------|---------------------------|
| 1 | NDC:0498-0733-00 | 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 |
|---------------------------|---|-----------------------------|---------------------------|
| unapproved drug other | | 03/14/2019 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------------|---|-----------------------------|---------------------------|
| unapproved drug other | | 03/14/2019 | 01/17/2020 |

4401 FIRST AID KIT

4401 first aid kit kit

Product Information

| | | | |
|---------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0498-4401 |
|---------------------|----------------|---------------------------|---------------|

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0498-4401-01 | 1 in 1 KIT; Type 0: Not a Combination Product | 03/14/2019 | |

Quantity of Parts

| Part # | Package Quantity | Total Product Quantity |
|--------|------------------|------------------------|
| Part 1 | 10 AMPULE | 3 mL |
| Part 2 | 10 POUCH | 3 mL |
| Part 3 | 10 POUCH | 4 mL |

Part 1 of 3**AMMONIA INHALENT**

ammonia inhalent inhalant

Product Information

| | |
|--------------------------------|--------------------------|
| Item Code (Source) | NDC:0498-3334 |
| Route of Administration | RESPIRATORY (INHALATION) |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-------------------|
| AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X) | AMMONIA | 0.045 g in 0.3 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|----------------------------|----------|
| ALCOHOL (UNII: 3K9958V90M) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0498-3334-00 | 0.3 mL in 1 AMPULE; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 09/18/2018 | 12/01/2022 |

Part 2 of 3

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

| | |
|-------------------------|---------------|
| Item Code (Source) | NDC:0498-0121 |
| Route of Administration | TOPICAL |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|---------------|
| POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4) | IODINE | 10 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---------------------------------------|----------|
| NONOXYNOL-9 (UNII: 48Q180SH9T) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0121-00 | 0.3 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 |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 09/18/2018 | |

Part 3 of 3

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information

| | |
|-------------------------|---------------|
| Item Code (Source) | NDC:0498-0733 |
| Route of Administration | TOPICAL |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------------------|----------------|
| LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE HYDROCHLORIDE ANHYDROUS | 20 mg in 1 mL |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 0.5 mL in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| MENTHOL (UNII: L7T10EIP3A) | |
| WATER (UNII: 059QF0KO0R) | |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0733-00 | 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 |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 03/14/2019 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|--|----------------------|--------------------|
| unapproved drug other | | 03/14/2019 | |

Labeler - Honeywell Safety Products USA, INC (118768815)