**4232 FIRST AID KIT- 4232 first aid kit**
**4221 FIRST AID KIT- 4221 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.*

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**0498-4232: First Aid Kit (1st aid Sp, EW, ASA, triple, burn spray WS, alcohol wipe- 340460F, 68180LBR)**
**0498-4221: First Aid Kit**

**Eyewash**
***Active ingredient***

Sterile Water 99%

**Eyewash**
***Purpose***

Eyewash

***Eyewash***
***Uses***

- for flushing the eye to remove loose foreign material, air pollutants or chlorinated water

**Eyewash**
***Warnings***

**For external use only** Obtain immediate medical treatment for all open wounds in or near eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

**Do not use**

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away.

**Stop use and ask a doctor if**

- you experience eye pain
- changes in vision
- continued redness or irritation of the eye
- condition worsens or persists

**Keep out of reach of children**

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

**Eyewash**
***Directions***

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

**Eyewash**
***Inactive ingredients***

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

**Eyewash**
***Questions***

1-800-430-5490

**Aspirin**
***Active ingredient (in each tablet)***

Aspirin 325 mg (NSAID)* *nonsteroidal anti-inflammatory drug

**Aspirin**
***Purpose***

Pain reliever/fever reducer

**Aspirin**
***Uses***

temporarily reduces fever and relieves minor aches and pains associated with:

- a cold
- headache
- toothache
- muscular aches
- backache
- minor pain of arthritis
- premenstrual and menstrual periods

**Aspirin**
***Warnings***

**Reye's syndrome**: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these

symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

**Allergy alert:** Aspirin may cause a severe allergic reaction which may include:
- hives
- facial swelling
- asthma (wheezing)
- shock

**Stomach bleeding warning:** This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:are:
- age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

**Do not use**
- if you are allergic to aspirin or any other pain reliever/fever reducer

**Ask a doctor before use if**
- stomach bleeding warning applies to you
- you have a history of stomach problems such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis or kidney disease
- you are taking a diuretic
- you have asthma

**Ask a doctor or pharmacist before use if you are**
- taking a prescription drug for diabetes, gout or arthritis

**Stop use and ask a doctor if**
- you experience any of the following signs of stomach bleeding:
- feel faint
- vomit blood
- have bloody or black stools
- have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- ringing in the ears or loss of hearing occurs
- any new symptoms appear

**If pregnant or breast-feeding,**

**If pregnant or breat-feeding,** ask a health professional before use. It is especially important not to use aspirin during the last three months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

**Keep out of reach of children.**

- In case of overdose, get medical help or contact Poison Control Center right away.

**Aspirin**
***Directions***

- drink a full glass of water with each dose
- adults and children 12 years of age and older: take 1 or 2 tablets every 4 hours while symptoms last, not more than 12 tablets in 24 hours
- children under 12 years of age: consult a doctor

**Aspirin**
***Other information***

- store at room temperature 15° - 30°C (59° - 86°F)
- TAMPER EVIDIENT PACKETS
- DO NOT USE IF OPEN OR TORN

**Aspirin**
***Inactive ingredients***

corn starch, croscarmellose sodium*, hypromellose*, microcrystalline cellulose*, mineral oil*, polyethylene glycol*, povidone, propylene glycol, silicon dioxide, stearic acid*, titanium dioxide*

*may contain these ingredients

**Aspirin**
***Questions or Comments***

1-800-430-5490

**Ammonia**
***Active ingredient***

Ammonia 15%

**Ammonia**
***Purpose***

Respiratory stimulant

**Ammonia**
***Uses***

- to prevent or treat fainting

**Ammonia**
***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**
***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**
***Other information***

- store at room temperature away from light

**Ammonia**
***Inactive ingredient***

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

**Ammonia**
***Questions or Comments?***

1-800-430-5490

**Water Soluble 1st Aid Spray**
***Active ingredient***

Benzethonium chloride 0.2% w/w - Benzocaine 10% w/w

**Water Soluble 1st Aid Spray**
***Purpose***

Topical antiseptic

Topical anesthetic

**Water Soluble 1st Aid Spray**
***Uses***

for temporary relief of pain and itching and helps protect against infection in

- minor cuts and scrapes
- insect bites
- minor skin irritations

**Water Soluble 1st Aid Spray**
***Warnings***

**For external use only**

**Flammable**

- keep away from fire or flame
- contents under pressure
- do not puncture or incinerate container
- do not expose to temperature above 120 0 F

**Do not use**

- in the eyes or other mucous membranes
- in cases of serious burns
- in case of deep orpuncture wounds
- for a prolonged period of time
- on large portion of the body

**Stop use and ask a doctor if**

- condition worsens or symptoms persist for more than 7 days
- condition clears up and occurs again within a few days
- redness, swelling, or irritation occurs

**Keep out of reach of children**

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

**Water Soluble 1st Aid Spray**
***Directions***

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only
- not intended for use on children

**Water Soluble 1st Aid pray**
***Other information***

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

**Water Soluble 1st Aid pray**
***Inactive ingredients***

dipropylene glycol, isobutane, N-butane, propane

**Burn Relief Water Soluble**
***Active ingredients***

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

**Burn Relief Water Soluble**
***Purpose***

Topical antiseptic

Topical anesthetic

Topical anesthetic

**Burn Relief Water Soluble**
***Uses***

for the temporary relief of pain and itching and helps protect against infection in:
- minor cuts and scrapes
- burns
- sunburn
- insect bites
- minor skin irritations

**Burn Relief Water Soluble**
***Warnings***

**For external use only**

**Flammable keep away from fire or flame**
- contents under pressure
- do not puncture or incinerate container
- do not expose to temperatures above 120 0 F

**Do not use**

- in or near the eyes or other mucous membranes
- in case of serious burns
- in case of deep or puncture wounds
- for prolonged period of time
- on large portion of the body

**Stop use and ask a doctor if**

- condition worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling, or irritation occurs

**Keep out of the reach of children**

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

**Burn Relief Water Soluble**
***Directions***

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage, if bandaged let dry first
- for adult institutional use only
- not intended for use on children

**Burn Relief Water Soluble**
***Other information***

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating or inhaling the contents may be harmful or fatal

**Burn Relief Water Soluble**
***Inactive ingredients***

dipropylene glycol, isobutane, n-butane, propane

**Triple**
***Active ingredients***

Bacitracin zinc 400 units

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Polymyxin B sulfate 5000 units

**Triple**

***Purpose***

First aid antibiotic

First aid antibiotic

First aid antibiotic

**Triple**
***Uses***

- first aid to help prevent infection in
- minor cuts
- scrapes
- burns

**Triple**
***Warnings***

**For external use only**

**Allergy alert** do not use if you are allergic to any of the ingredients

**Do not use**

- in the eyes
- over large areas of the body

**Ask a doctor before use if you have**

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

**Stop use and ask a doctor if**

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

**Keep out of the reach of children**

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

**Triple**
***Directions***

- clean the affected area
- apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

**Triple**

***Other information***

- store at 15 $^{0}$ to 25 $^{0}$ C (59 $^{0}$ to 77 $^{0}$ F)
- tamper evident sealed packets
- do not use if packet is torn or opened

**Triple**
***Inactive ingredient***

petrolatum

**Alcohol**
***Active ingredient***

Isopropyl alcohol 70%

**Alcohol**
***Purpose***

First aid antiseptic

**Alcohol**
***Uses***

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

**Alcohol**
***Warnings***

**For external use only**

**Flammable, keep away from fire or flame.**

**Do not use**

- in the eyes
- over large areas of the body

**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 consult a doctor**

- if condition persists or gets worse

**Keep out of reach of children**
- If swallowed, get medical help or contact a Poison Control Center right away.

**Alcohol**
***Directions***
- clean the affected area
- apply wipe to affected area 1 to 3 times daily
- may be covered with a sterile bandage
- discard wipe after single use

**Alcohol**
***Other information***

store at room temperature 15 $^{0}$ to 25 $^{0}$ C (59 $^{0}$ to 77 $^{0}$ F)

**Alcohol**
***Inactive ingredient***

water

**4221**
**340460F KIT CONTENTS**

1 3/4 X 3 PLAS 100/BOX

1 AMMONIA INHALANTS 10 PER

1 INSTANT COLD PACK 4" X 6"

1 ADHESIVE TAPE W/P 1/2"X 5 YD

1 FIRST AID GUIDE ASHI

1 GAUZE CLEAN-WRAP BDGE N/S 2"

1 BLOODSTOPPER

1 CO-FLEX BANDAGE 2"X 5YDS TAN

1 FIRST AID SPRAY AEROSOL 3 OZ

1 BURN SPRAY 3 OZ

1 4OZ BFS EYEWASH TRILINGUAL BOTTLE

1 SCISSOR BDGE 4" RED PLS HDL

1 KIT TWEEZER 3 1/2" SLANTED

LBL STOCK 6-3/8"X4"

LBL STOCK 4"X2-7/8"

1 LBL STOCK 3"x1-7/8"

1 PR LRG NITRILE GLVES ZIP BAG

2 BAYER 12 PACK PER ZIP BAG

2 TRIPLE BIOTIC FOIL PACK EACH

6 WIPE ALCOHOL PREP IPA 70% (DUKAL)

1 KIT STL 36 UN WHT 01 HOR SHELF

1 TRI BNDG NON WOVEN 40"X40"X56"

10 NON ADHERENT PAD 2" X 3"

1 WOVEN KNUCKLE 8'S

1 FINGERTIP "T" 8/BX

**4232**
**68180LBR Kit Contents**

1 3/4 X 3 PLAS 100/BOX

1 1X3 PLASTIC 100/BOX

1 FINGERTIP "T" WOVEN 40/BOX

1 1X3 WOVEN SING 50/BOX

1 SWIFT KNUCKLE 40/BX

1 AMMONIA INHALANTS 10 PER

1 INSTANT COLD PACK 4" X 6"

1 ELASTIC TAPE 1" X 5YD

1 FORCEPS POINTED METAL

1 O/H TAPE ADHESIVE TRI-CUT

1 FIRST AID GUIDE ASHI

4 GAUZE CLEAN-WRAP BDGE N/S 2"

2 BLOODSTOPPER

1 NON ADHERENT PADS 2"X3" 50'S

2 GZE PADS STERILE 2"X 2" 25'S

1 GZE PADS STERILE 4"X 4" 25'S

1 CO-FLEX BANDAGE 2"X 5YDS TAN

1 COTTON TIPS 100 PER VIAL

1 ANTISEPTIC WIPES BZK CHL 20'S

1 FIRST AID SPRAY AEROSOL 3 OZ

1 ALCOHOL WIPES 50'S

1 ASPIRIN IND PK 5 GR 2/ENV 250

1 BURN SPRAY 3 OZ

1 TRIPLE BIOTIC .5 GRAM PKT 20

4 1 OZ, BUFF EYEWASH

1 SCISSOR BDGE 4" RED PLS HDL

1 180 EMPTY BLANK NO LOGO

1 POCKET INSERT RED #180 KIT 4R

1 TONGUE BLADES SR WRAPPED 6'S

1 LBL STOCK 6-3/8"X4"

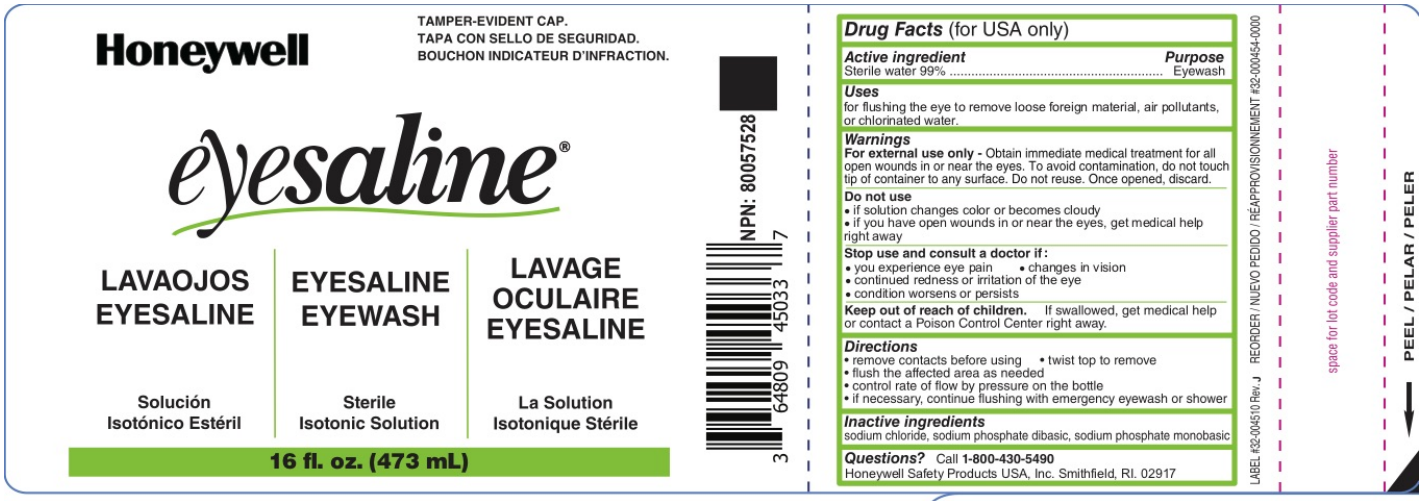
1 LBL STOCK 4"X2-7/8"

2 PR LRG NITRILE GLVES ZIP BAG

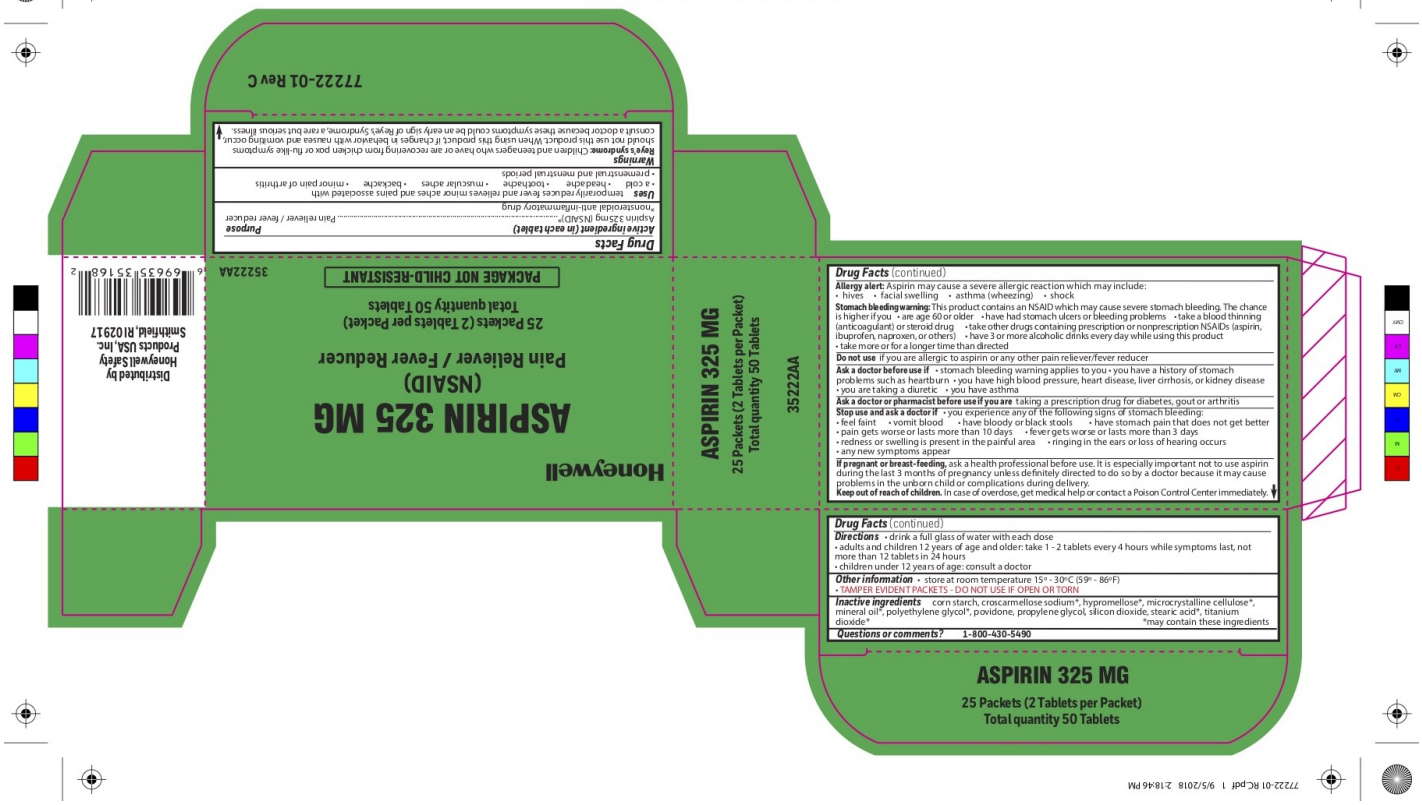
2 TRI BNDG NON WOVEN 40"X40"X56"

1 RED BIO BAGS 2/BX

**Eyewash**
***Principal Display Panel***



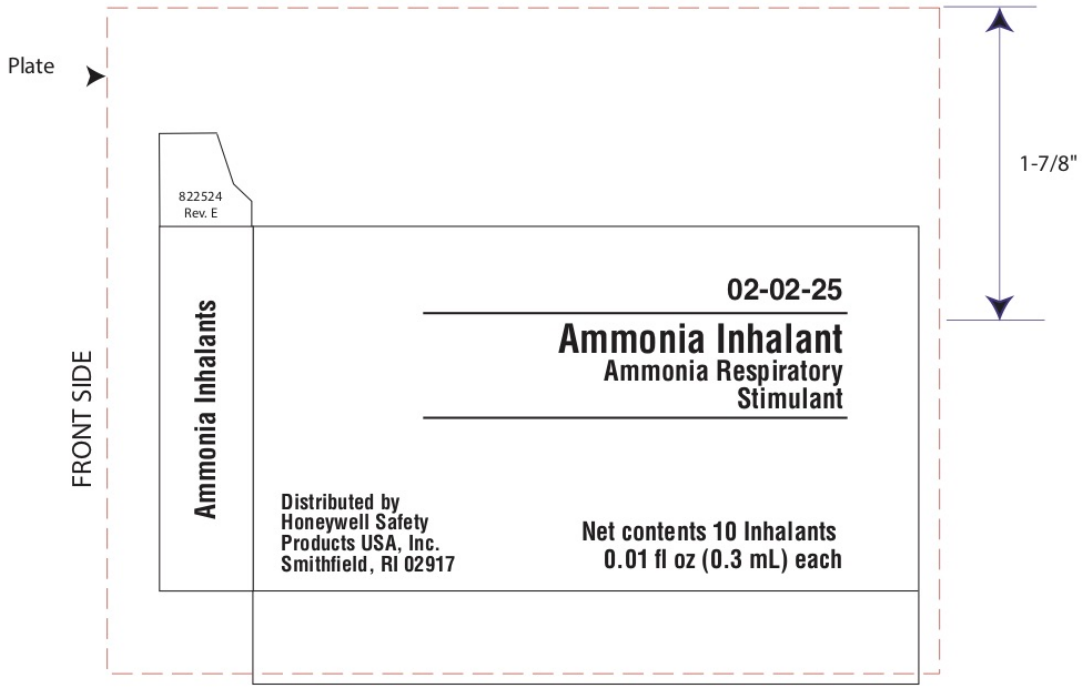

**Aspirin**
***Principal Display Panel***

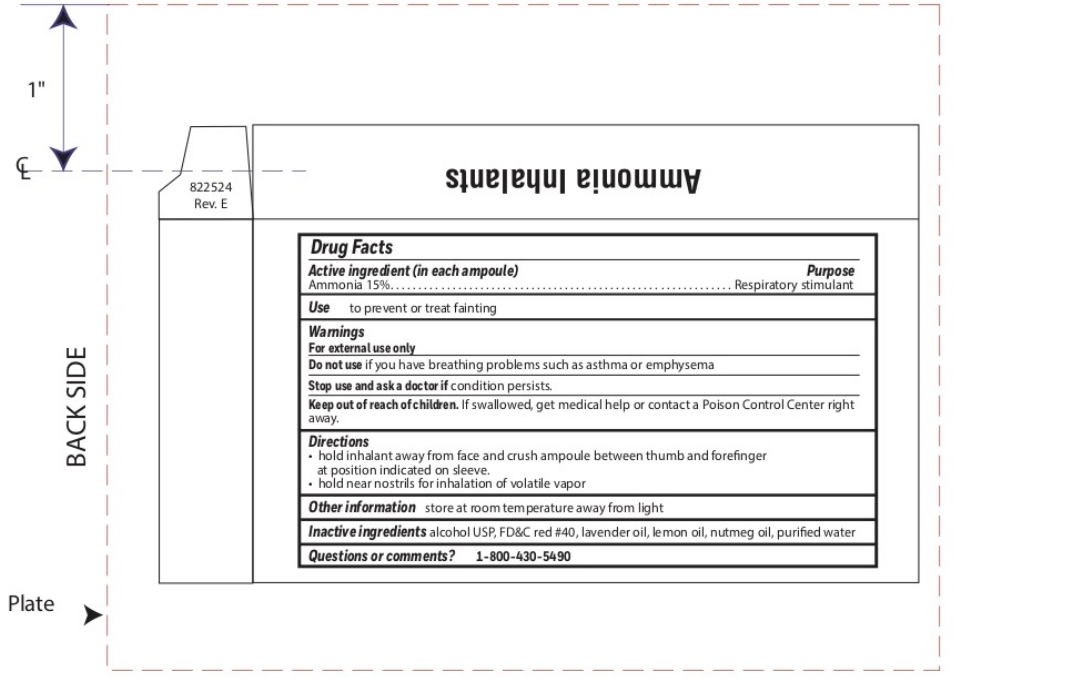



**Ammonia**

***Principal Display Panel***

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



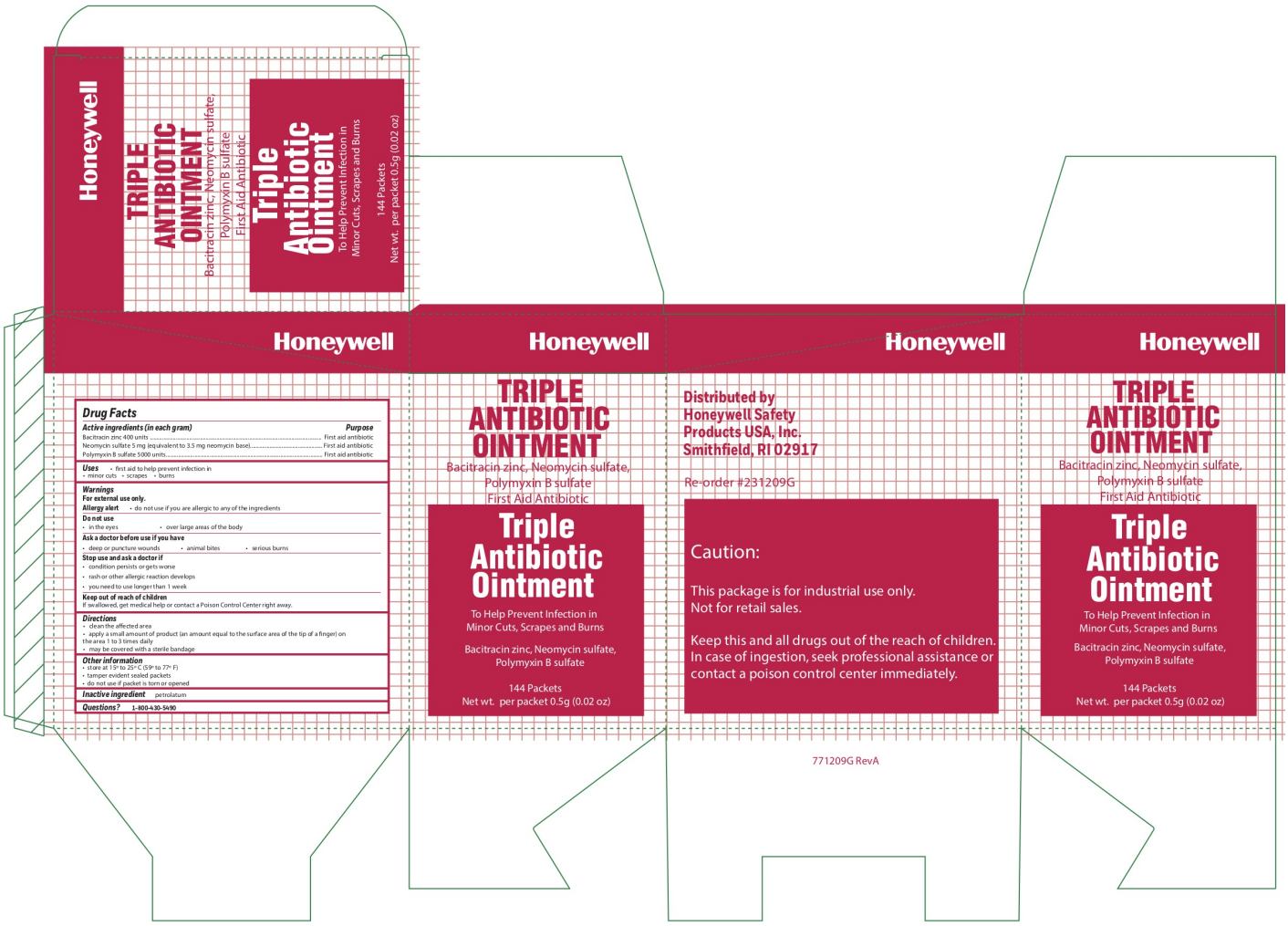

796006 Rev. E (page 3 of 3)

**Water Soluble 1st Aid Spray**

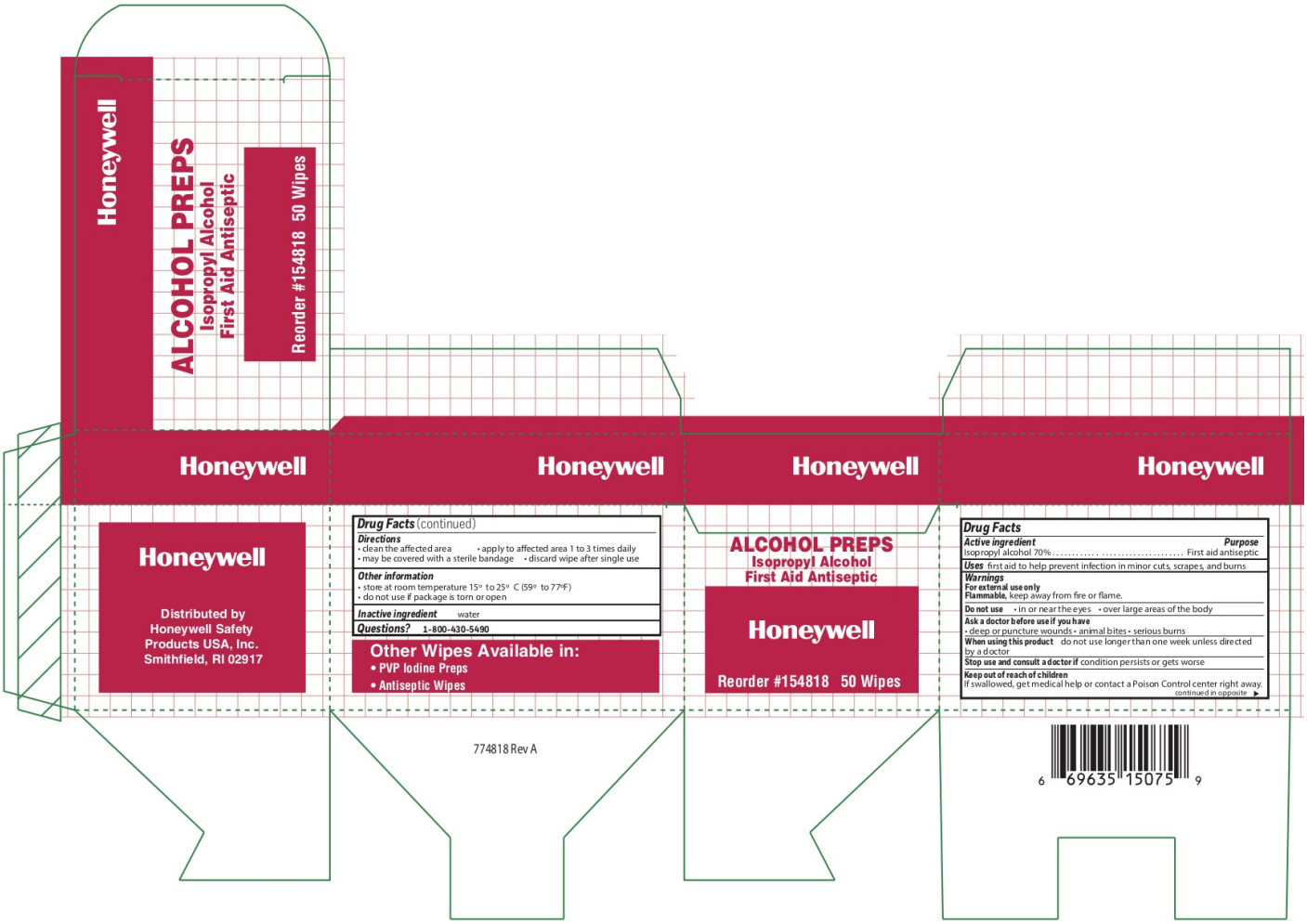
## Principal Display Panel


## Burn Relief Water Soluble
## *Principal Display Panel*

## Triple
## *Principal Display Panel*




## Alcohol
## *Principal Display Panel*




**4221 Kit Label**
**340460F**


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

**4232 Kit Label**
**68180LBR**

BRONER
GLOVE AND SAFETY
1-800-521-1318

First Aid

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

# 4232 FIRST AID KIT

4232 first aid kit kit

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:0498-4232 |

| Packaging | | | | |
|---|---|---|---|---|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0498-4232-01 | 1 in 1 KIT; Type 0: Not a Combination Product | 10/18/2018 | |

| Quantity of Parts | | |
|---|---|---|
| Part # | Package Quantity | Total Product Quantity |
| **Part 1** | 4 BOTTLE | 120 mL |
| **Part 2** | 125 PACKET | 250 |
| **Part 3** | 10 AMPULE | 3 mL |
| **Part 4** | 1 CAN | 85 g |
| **Part 5** | 1 CAN | 85 g |
| **Part 6** | 20 PACKET | 10 g |
| **Part 7** | 50 POUCH | 20 mL |

## Part 1 of 7

## EYESALINE EMERGENCY EYEWASH

purified water liquid

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0100 |
| **Route of Administration** | OPHTHALMIC |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| Ingredient Name | Basis of Strength | Strength |
| **WATER** (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) | WATER | 98.6 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| Ingredient Name | Strength |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **SODIUM PHOSPHATE, DIBASIC** (UNII: GR686LBA74) | |
| **SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE** (UNII: 593YOG76RN) | |

| Packaging | | | | |
|---|---|---|---|---|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0498-0100-01 | 30 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M018 | 12/18/2018 | |

## Part 2 of 7

# ASPIRIN

aspirin tablet

### Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:0498-0114 |
| **Route of Administration** | ORAL |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **ASPIRIN** (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E) | ASPIRIN | 325 mg |

### Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CELLULOSE, MICROCRYSTALLINE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **CROSCARMELLOSE SODIUM** (UNII: M28OL1HH48) | |
| **HYPROMELLOSE 2208 (100 MPA.S)** (UNII: B1QE5P712K) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |

### Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white | **Score** | 2 pieces |
| **Shape** | ROUND | **Size** | 10mm |
| **Flavor** | | **Imprint Code** | FR21 |
| **Contains** | | | |

### Packaging

| | | | Marketing Start | Marketing End |
|---|---|---|---|---|

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| 1 | NDC:0498-0114-01 | 2 in 1 PACKET; 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 7

## 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 4 of 7

# FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0031 |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZETHONIUM CHLORIDE** (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP) | BENZETHONIUM CHLORIDE | 0.2 g in 100 g |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 10 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **ISOBUTANE** (UNII: BXR49TP611) | |
| **BUTANE** (UNII: 6LV4FOR43R) | |
| **PROPANE** (UNII: T75W9911L6) | |
| **DIPROPYLENE GLYCOL** (UNII: E107L85C40) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:0498-0031-40 | 85 g in 1 CAN; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 09/19/2018 | |

## Part 5 of 7

# BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0021 |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZETHONIUM CHLORIDE** (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP) | BENZETHONIUM CHLORIDE | 0.2 g in 100 g |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 10 g in 100 g |
| **MENTHOL** (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) | MENTHOL | 0.33 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **ISOBUTANE** (UNII: BXR49TP611) | |
| **BUTANE** (UNII: 6LV4FOR43R) | |
| **PROPANE** (UNII: T75W9911L6) | |
| **DIPROPYLENE GLYCOL** (UNII: E107L85C40) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:0498-0021-40 | 85 g in 1 CAN; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 11/12/2018 | |

## Part 6 of 7

## TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0750 |
| **Route of Administration** | TOPICAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **POLYMYXIN B SULFATE** (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) | POLYMYXIN B | 5000 [iU] in 1 g |
| **BACITRACIN ZINC** (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I) | BACITRACIN | 400 [iU] in 1 g |
| **NEOMYCIN SULFATE** (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297) | NEOMYCIN | 3.5 mg in 1 g |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **PETROLATUM** (UNII: 4T6H12BN9U) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0498-0750-36 | 0.5 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 09/19/2018 | |

# Part 7 of 7

# ALCOHOL WIPE

isopropyl alcohol swab

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:0498-0143 |
| **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:0498-0143-04 | 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 | | 09/18/2018 | |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 10/18/2018 | |

## 4221 FIRST AID KIT

4221 first aid kit kit

### Product Information

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0498-4221 |
|---|---|---|---|

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0498-4221-01 | 1 in 1 KIT; Type 0: Not a Combination Product | 10/18/2018 | |

### Quantity of Parts

| Part # | Package Quantity | Total Product Quantity |
|---|---|---|
| **Part 1** | 1 BOTTLE | 118 mL |

| | | |
|---|---|---|
| **Part 2** | 12 PACKET | 24 |
| **Part 3** | 1 AMPULE | 0.3 mL |
| **Part 4** | 1 CAN | 85 g |
| **Part 5** | 1 CAN | 85 g |
| **Part 6** | 2 PACKET | 1.8 g |
| **Part 7** | 6 POUCH | 2.4 mL |

## Part 1 of 7

## EYESALINE EMERGENCY EYEWASH

purified water liquid

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0100 |
| **Route of Administration** | OPHTHALMIC |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) | WATER | 98.6 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE** (UNII: 593YOG76RN) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **SODIUM PHOSPHATE, DIBASIC** (UNII: GR686LBA74) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:0498-0100-02 | 118 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M018 | 12/18/2018 | |

## Part 2 of 7

## ASPIRIN

aspirin tablet

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:0498-0114 |
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **ASPIRIN** (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E) | ASPIRIN | 325 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CELLULOSE, MICROCRYSTALLINE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **STARCH, CORN** (UNII: O8232NY3SJ) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **CROSCARMELLOSE SODIUM** (UNII: M28OL1HH48) | |
| **HYPROMELLOSE 2208 (100 MPA.S)** (UNII: B1QE5P712K) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white | **Score** | 2 pieces |
| **Shape** | ROUND | **Size** | 10mm |
| **Flavor** | | **Imprint Code** | FR21 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0498-0114-01 | 2 in 1 PACKET; 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 7

# 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 4 of 7

# FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0031 |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZETHONIUM CHLORIDE** (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP) | BENZETHONIUM CHLORIDE | 0.2 g in 100 g |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 10 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **ISOBUTANE** (UNII: BXR49TP611) | |
| **BUTANE** (UNII: 6LV4FOR43R) | |
| **PROPANE** (UNII: T75W9911L6) | |
| **DIPROPYLENE GLYCOL** (UNII: E107L85C40) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:0498-0031-40 | 85 g in 1 CAN; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 09/19/2018 | |

## Part 5 of 7

## BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0021 |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZETHONIUM CHLORIDE** (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP) | BENZETHONIUM CHLORIDE | 0.2 g in 100 g |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 10 g in 100 g |

| | | |
|---|---|---|
| **MENTHOL** (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) | MENTHOL | 0.33 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **ISOBUTANE** (UNII: BXR49TP611) | |
| **BUTANE** (UNII: 6LV4FOR43R) | |
| **PROPANE** (UNII: T75W9911L6) | |
| **DIPROPYLENE GLYCOL** (UNII: E107L85C40) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:0498-0021-40 | 85 g in 1 CAN; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 11/12/2018 | |

## Part 6 of 7

## TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0750 |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **POLYMYXIN B SULFATE** (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) | POLYMYXIN B | 5000 [iU] in 1 g |
| **BACITRACIN ZINC** (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I) | BACITRACIN | 400 [iU] in 1 g |
| **NEOMYCIN SULFATE** (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297) | NEOMYCIN | 3.5 mg in 1 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |

| | |
|---|---|
| **PETROLATUM** (UNII: 4T6H12BN9U) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | white | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | | **Imprint Code** | |
| **Contains** | | | |

**Packaging**

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:0498-0750-35 | 0.9 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 09/19/2018 | |

## Part 7 of 7

## ALCOHOL WIPE

isopropyl alcohol swab

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:0498-0143 |
| **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:0498-0143-04 | 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 | | 09/18/2018 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 10/18/2018 | 04/22/2020 |

## Labeler - Honeywell Safety Products USA, INC (118768815)

Revised: 1/2024 Honeywell Safety Products USA, INC