

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

**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 breast-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 EVIDENT 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 or puncture 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<sup>0</sup> to 25<sup>0</sup> C (59<sup>0</sup> to 77<sup>0</sup> 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<sup>0</sup> to 25<sup>0</sup> C (59<sup>0</sup> to 77<sup>0</sup> 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"

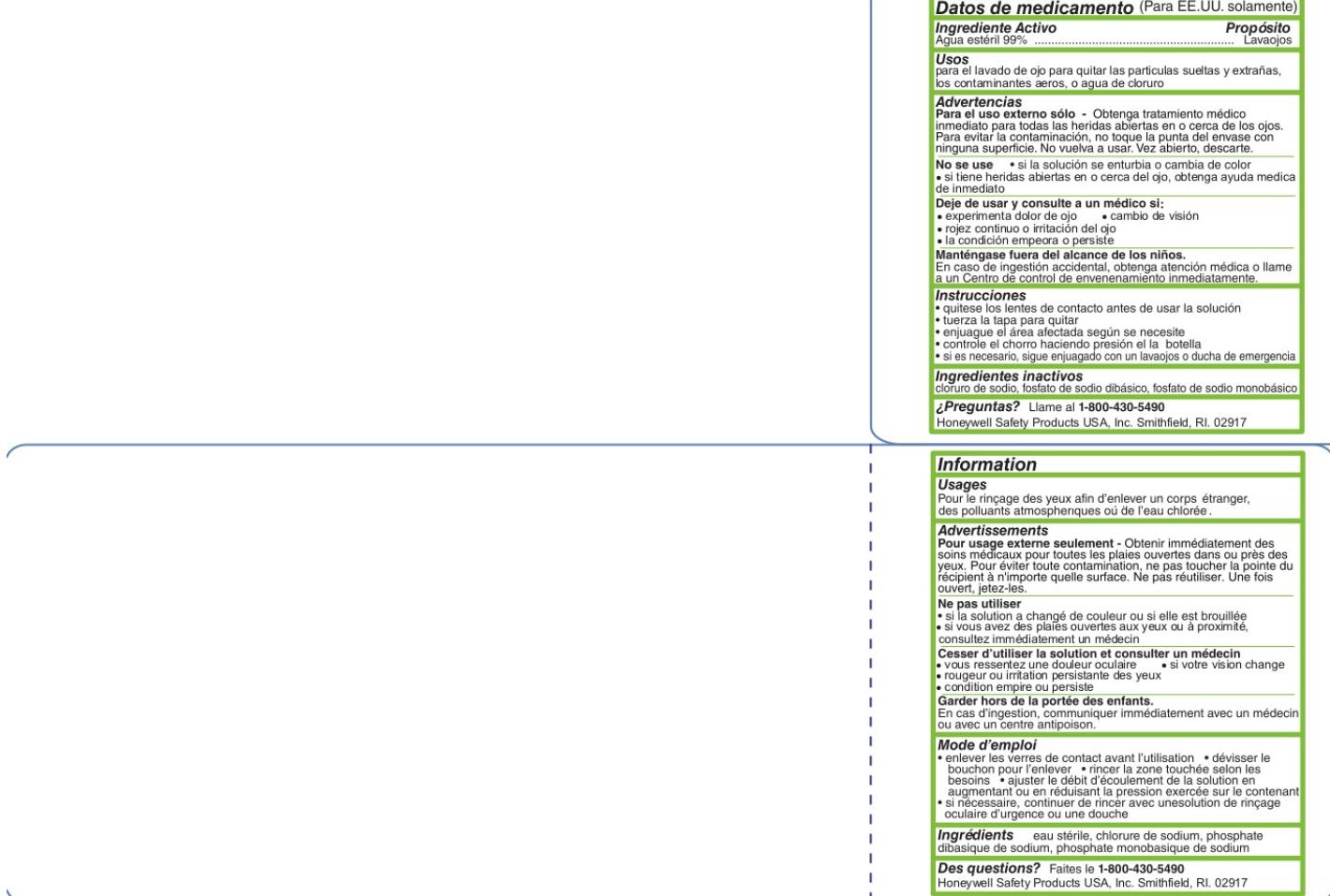
1 RED BIO BAGS 2/BX

## **Eyewash**

### *Principal Display Panel*

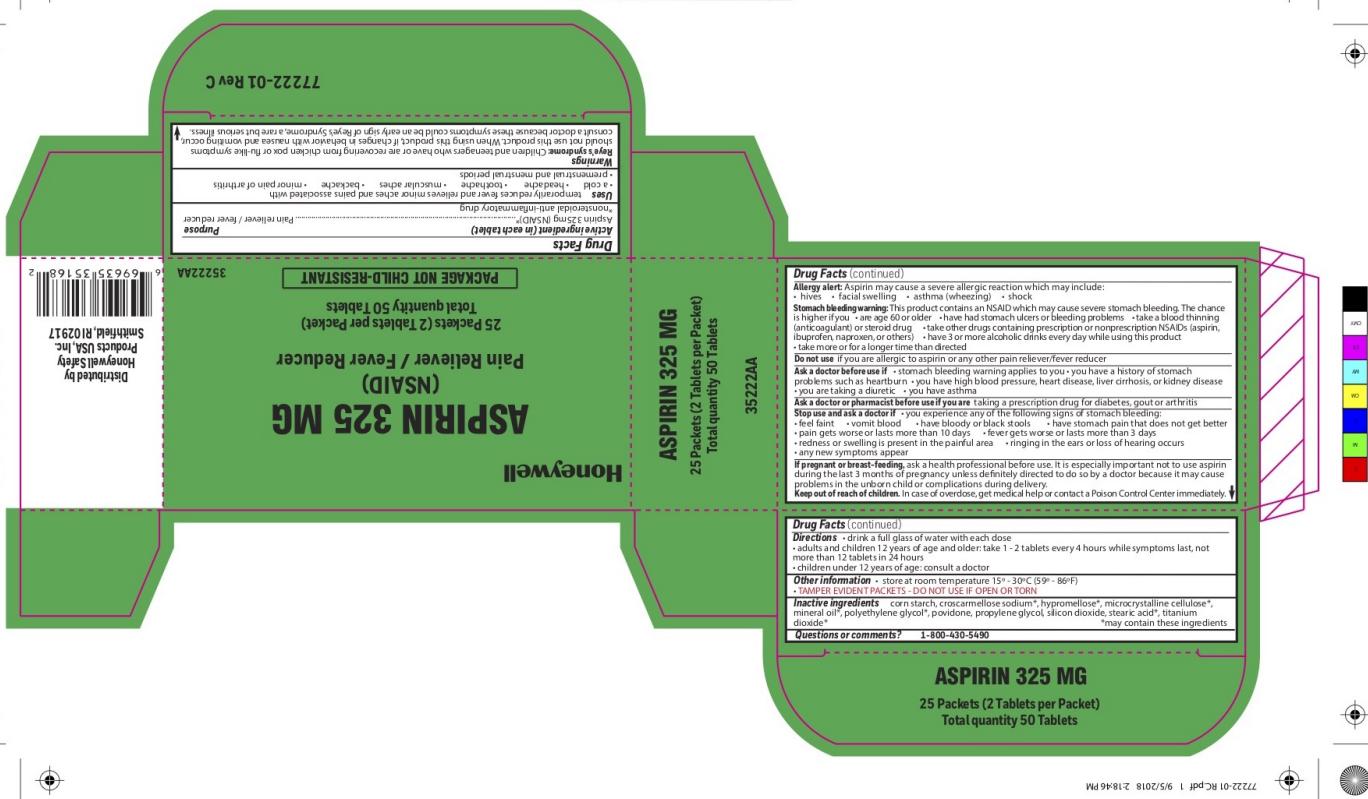


ANSWER



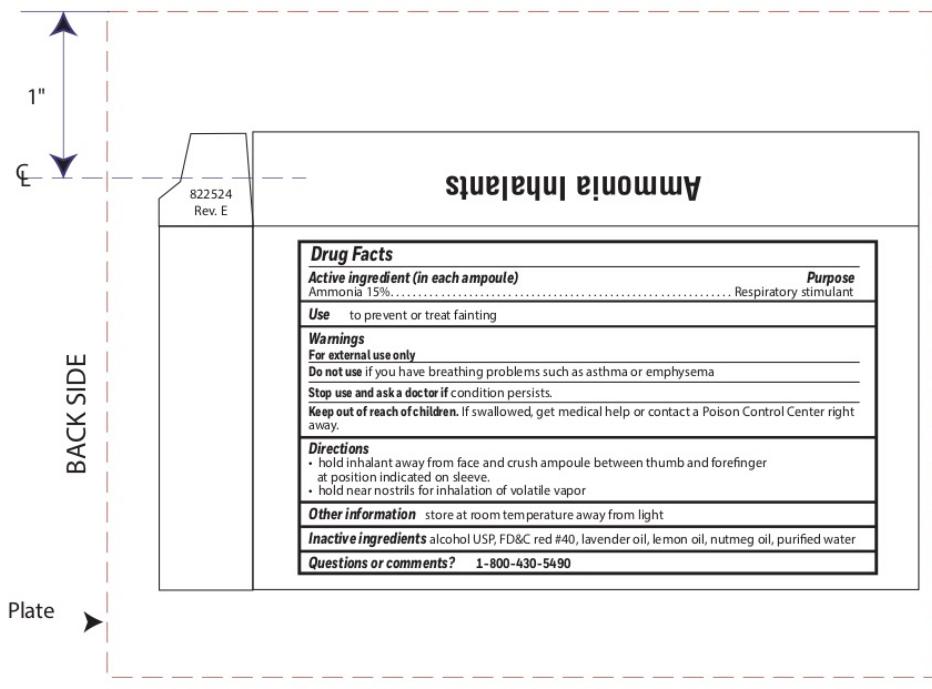
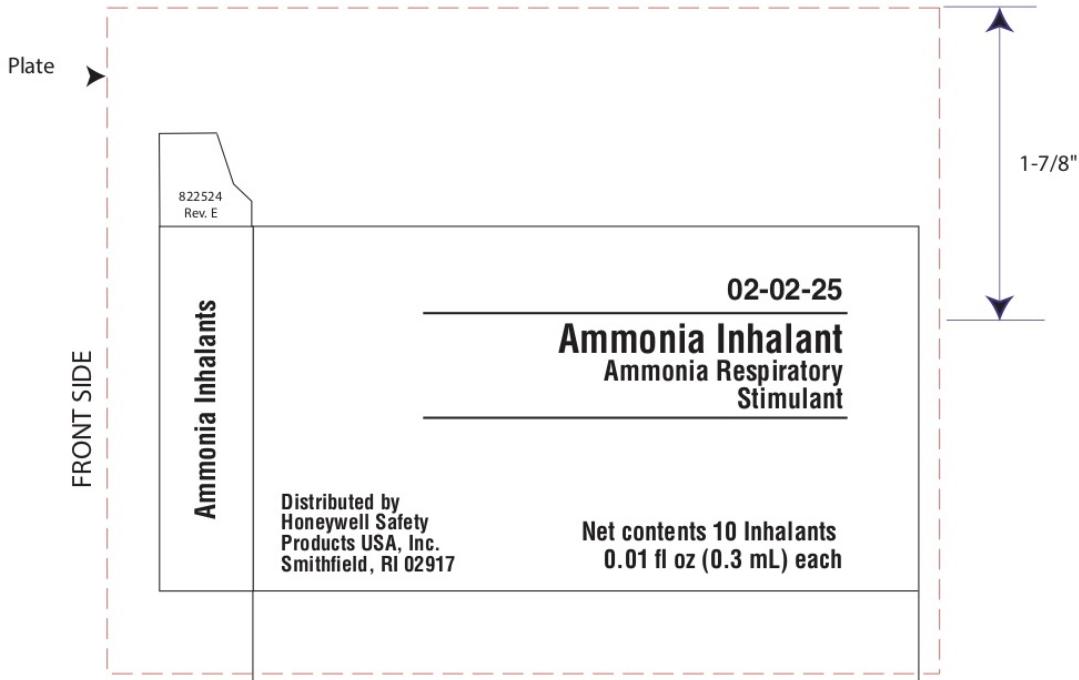
# **Aspirin**

## *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)

## Principal Display Panel



47-0303

Cat. No. 151019

### DRUG FACTS

#### Active ingredients

	Purpose
Benzethonium chloride 0.2% w/w.....	Topical antiseptic
Benzocaine 10% w/w.....	Topical anesthetic

#### Uses

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

#### 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°F

Do not use	• in or near eyes or other mucus membranes	• in case of serious burns
	• in case of deep or puncture wounds	• for a prolonged period of time
	• on large portion of the body	

#### Stop use and ask a doctor if:

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

#### 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

#### Other information

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

#### Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

#### Questions or comments?

1-800-430-5490

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Products USA, Inc.  
Smithfield, RI 02917

**Honeywell**

## Burn Relief Water Soluble Principal Display Panel



47-0306

Cat. No. 201005

### DRUG FACTS

#### Active ingredients

	Purpose
Benzethonium chloride 0.2% w/w.....	Topical antiseptic
Benzocaine 10% w/w.....	Topical anesthetic
Menthol .33%.....	Topical anesthetic

#### Uses

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

#### 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°F

Do not use	• in or near eyes or other mucus membranes	• in case of serious burns
	• in case of deep or puncture wounds	• for a prolonged period of time
	• on large portion of the body	

#### Stop use and ask a doctor if:

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

#### 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

#### Other information

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

#### Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

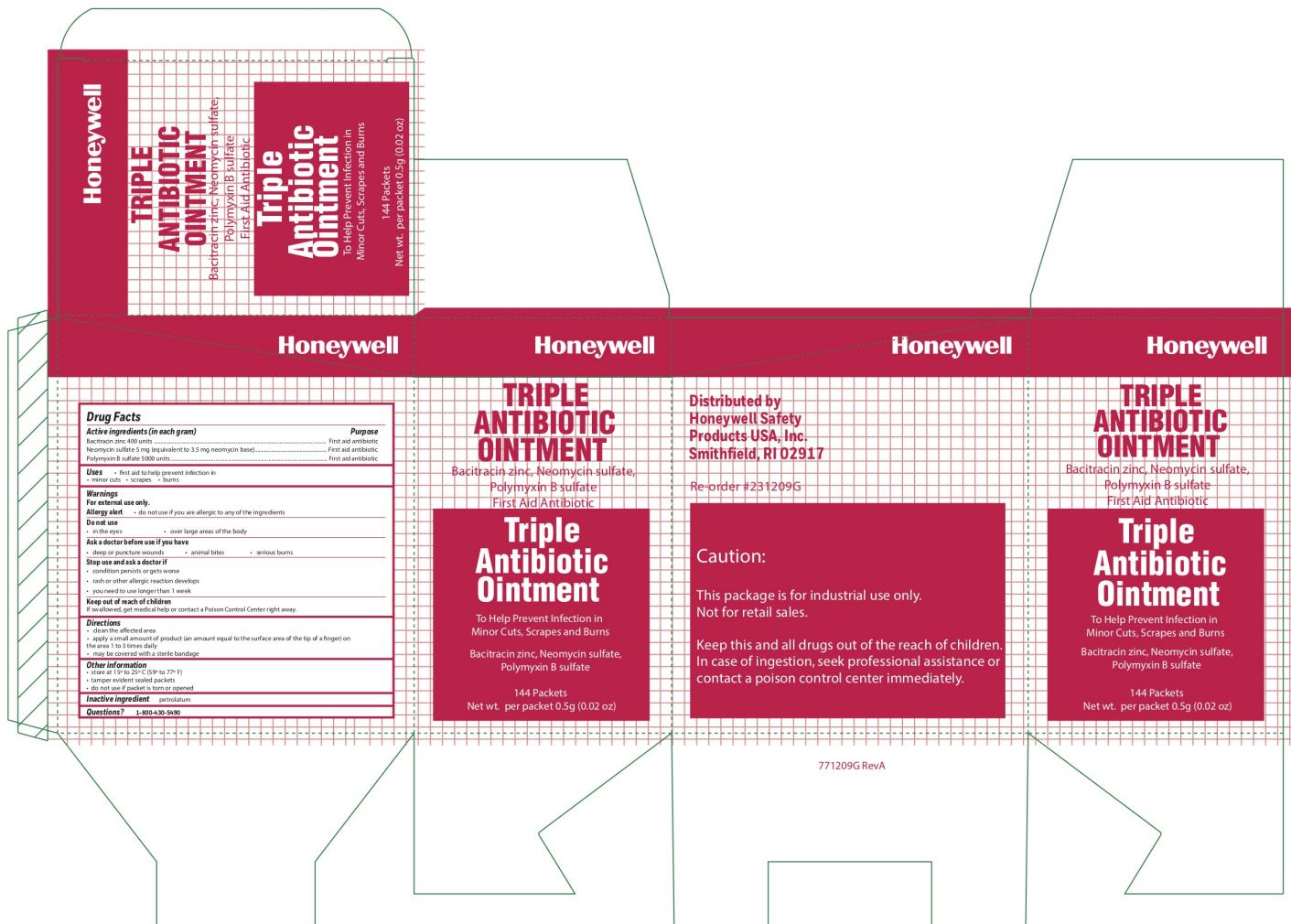
#### Questions or comments?

1-800-430-5490

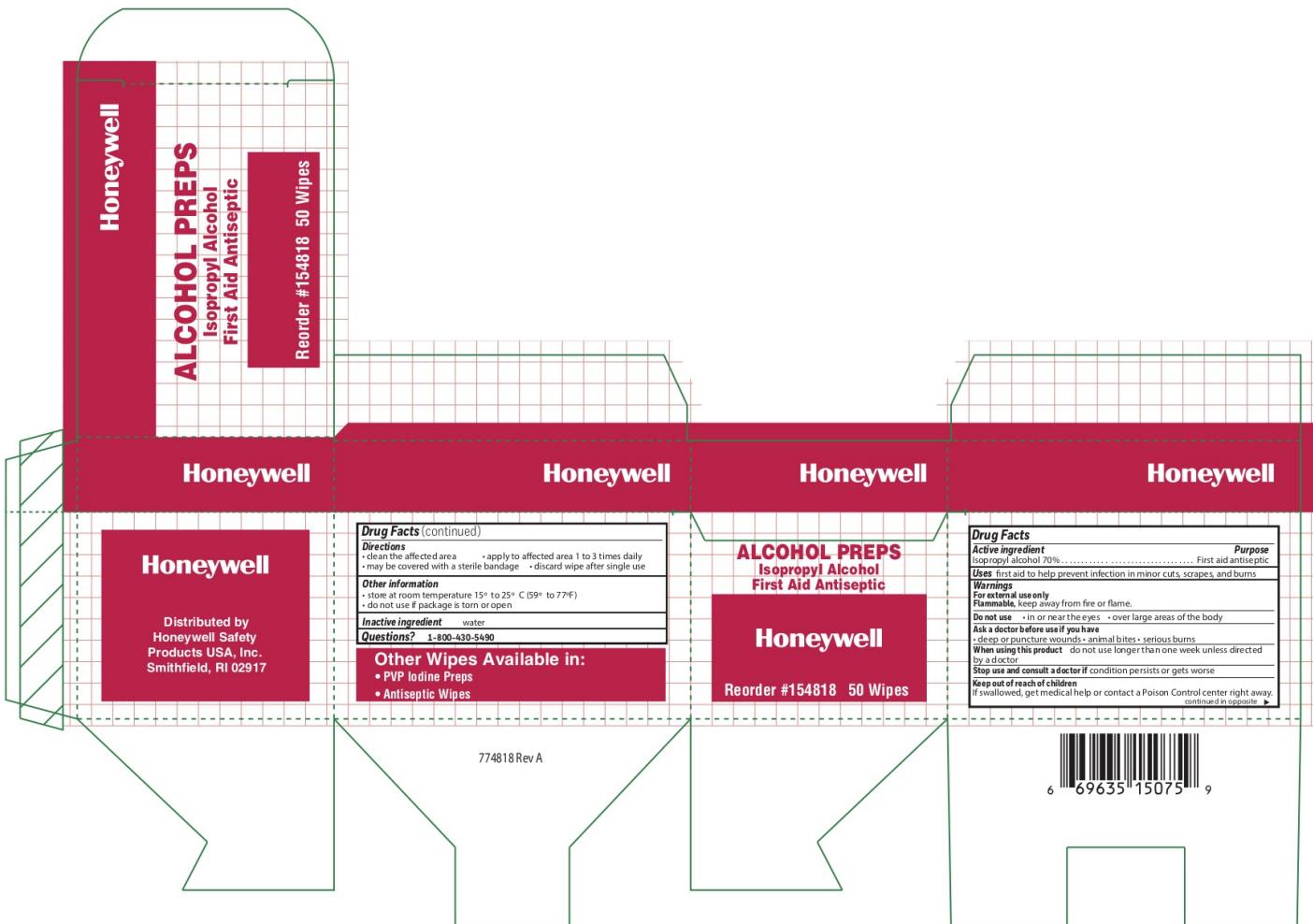
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Products USA, Inc.  
Smithfield, RI 02917

**Honeywell**

## Triple Principal Display Panel



## Alcohol Principal Display Panel



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



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**4232 Kit Label  
68180LBR**



**BRONER**  
**GLOVE AND SAFETY**  
**1-800-521-1318**



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 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:0498-0100-01	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
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**Part 2 of 7****ASPIRIN**

aspirin tablet

**Product Information**

<b>Item Code (Source)</b>	NDC:0498-0114
<b>Route of Administration</b>	ORAL

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE 2208 (100 MPAS) (UNII: B1QE5P712K)	
MINERAL OIL (UNII: T5L8T28FGP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	10 mm
<b>Flavor</b>		<b>Imprint Code</b>	FR21
<b>Contains</b>			

**Packaging**

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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
OTC monograph not final	part343	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
<b>BENZETHONIUM CHLORIDE</b> (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>ISOBUTANE</b> (UNII: BXR49TP611)	
<b>BUTANE</b> (UNII: 6LV4FOR43R)	
<b>PROPANE</b> (UNII: T75W9911L6)	
<b>DIPROPYLENE GLYCOL</b> (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
OTC monograph not final	part333E	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
<b>BENZETHONIUM CHLORIDE</b> (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g
<b>MENTHOL</b> (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
OTC monograph not final	part348	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
OTC monograph final	part333B	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
OTC monograph not final	part333A	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
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### 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 final	part349	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: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE 2208 (100 MPAS) (UNII: B1QE5P712K)	
MINERAL OIL (UNII: T5L8T28FGP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

## Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	10 mm

<b>Flavor</b>		<b>Imprint Code</b>	FR21
<b>Contains</b>			

## 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
OTC monograph not final	part343	09/18/2018	

## Part 3 of 7

### AMMONIA INHALENT

ammonia inhalent inhalant

## Product Information

<b>Item Code (Source)</b>	NDC:0498-3334		
<b>Route of Administration</b>	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
OTC monograph not final	part333E	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
<b>BENZETHONIUM CHLORIDE</b> (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
<b>BENZOCAINE</b> (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.33 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>ISOBUTANE</b> (UNII: BXR49TP611)	
<b>BUTANE</b> (UNII: 6LV4FOR43R)	
<b>PROPANE</b> (UNII: T75W9911L6)	
<b>DIPROPYLENE GLYCOL</b> (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
OTC monograph not final	part348	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
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
<b>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>NEOMYCIN SULFATE</b> (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
OTC monograph final	part333B	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
OTC monograph not final	part333A	09/18/2018	

## Marketing Information

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

**Labeler** - Honeywell Safety Products USA, INC (079287321)

## Establishment

Name	Address	ID/FEI	Business Operations
James Alexander		040756421	manufacture(0498-3334)

## Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4221, 0498-4232)

## Establishment

Name	Address	ID/FEI	Business Operations
Ultra Seal Corporation		085752004	manufacture(0498-0114)

## Establishment

Name	Address	ID/FEI	Business Operations
Dixon Investments		115315822	manufacture(0498-0031, 0498-0021)

## Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0750)

## Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc.		167518617	manufacture(0498-0100)

## Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0143)