

**4253 FIRST AID KIT- 4253 first aid kit**  
**4197 FIRST AID KIT- 4197 first aid kit**  
**4284 FIRST AID KIT- 4284 first aid kit**  
**4198 FIRST AID KIT- 4198 first aid kit**  
**4199 FIRST AID KIT- 4199 first aid kit**  
**4254 FIRST AID KIT- 4254 first aid kit**  
**4336 FIRST AID KIT- 4336 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.*

-----

**0498-4197, 4198, 4199, 4253, 4254, 4284, 4336: First Aid Kit (FABC, EW, triple, PVP wipes, Sting relief- 019710-0006L,019711-0006L, 019712-0007L, Z019710-0006L, Z019712-0007L, SF00002448, Z019821)**

**First Aid Burn Cream**  
***Active ingredient***

Benzalkonium chloride 0.13%

Lidocaine HCl 0.5%

**First Aid Burn Cream**  
***Purpose***

First aid antiseptic

External analgesic

**First Aid Burn Cream**  
***Uses***

- prevent skin infection
- for temporary relief of pain associated with minor burns

**First Aid Burn Cream**  
***Warnings***

**For external use only**

**Do not use**

- in or near the eyes
- if you are allergic to any of the ingredients
- in large areas of the body, particularly over raw surfaces or blistered areas
- for more than 10 days

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

- deep or puncture wounds
- animal bites
- serious burns

**Stop use and ask a doctor if**

- condition worsens
- symptoms persist for more than 7 days or clear up and occurs again within a few days

**First Aid Burn Cream*****Directions***

- **adults and children 2 years of age and older:**
- clean the affected area
- apply a small amount of this product (equal to the surface area of the tip of a finger) onto affected area 1 to 3 times daily
- may be covered with a sterile bandage
- children under 2 years of age: consult a doctor

**First Aid Burn Cream*****Other information***

- tamper evident sealed packets
- do not use if packet is opened or torn

**First Aid Burn Cream*****Inactive ingredients***

aloe barbadensis juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, trolamine, water

**First Aid Burn Cream*****Questions***

1-800-430-5490

**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

## **Triple**

### **Active ingredient (each gram contains)**

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

## **Triple**

### ***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.

**Sting 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

**PVP**

***Active ingredient***

Povidone-iodine 10% (equivalent to 1% titratable iodine)

**PVP**

***Purpose***

First aid antiseptic

**PVP**

***Uses***

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

**PVP**

***Warnings***

**For external use only.**

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

- deep or puncture wounds
- animal bites
- serious burns

**Stop use and ask a doctor if**

- condition worsens or persists for more than 72 hours
- 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***

- clean the affected area
- apply 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first

**PVP*****Other information***

- do not use on individuals who are allergic or sensitive to iodine
- store at controlled temperature 59-86°F (15-30°C)
- do not use if pouch is open or torn

**PVP*****Inactive ingredients***

nonoxynol 9, water

**PVP*****Questions***

1-800-430-5490

**4197****019710-0006L Kit Contents**

- 1 KNUCKLE BAND 8 PER
- 1 TRIPLE ANTIBIOTIC 10 PER
- 1 FIRST AID BURN CREAM 6 PER
- 1 TRIANGULAR BDG, NON-STERILE
- 1 GAUZE PADS, 3" X 3", 4 PER
- 1 ADH TAPE, .5" X 2.5 YD, 2 PER
- 1 FORCEPS & SCISSORS, 1 EA
- 1 GAUZE COMP, 1 SQ YARD, 1 PER

1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 BANDAGE COMP, 4" OFFSET, 1 PER  
2 ADHESIVE BDG, PLSTIC, 1"X3"16PER  
1 PVP IODINE WIPES 10 PER  
1 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI  
LBL STOCK 6-3/8"X4"  
LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 LBL NORTH CONTS 6.75X3.5 ID B  
1 KIT, PP 16 UNIT FA  
1 LBL CONTENTS ANSI Z308.1-2009 REV B  
1 LBL 16U CVR NORTH ID C  
1 STING Relief WIPES 10

## **4198**

### **019711-0006L Kit Contents**

1 KNUCKLE BAND 8 PER  
1 TRIPLE ANTIBIOTIC 10 PER  
1 FIRST AID BURN CREAM 6 PER  
1 TRIANGULAR BDG, NON-STERILE  
1 GAUZE PADS, 3" X 3", 4 PER  
1 ADH TAPE, .5" X 2.5 YD, 2 PER  
1 FORCEPS & SCISSORS, 1 EA  
1 GAUZE COMP, 1 SQ YARD, 1 PER  
1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 BANDAGE COMP, 4" OFFSET, 1 PER  
1 ADHESIVE BDG, PLSTIC, 1"X3"16PER  
1 PVP IODINE WIPES 10 PER  
1 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI



LBL STOCK 6-3/8"X4"  
LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 LBL NORTH CONTS 6.75X3.5 ID B  
1 KIT STL 16 UN (HORIZONTAL)  
1 LBL CONTENTS ANSI Z308.1-2009 REV B  
1 LBL 16U CVR NORTH ID C  
1 STING Relief WIPES 10

**4199**  
**019712-0007L Kit Contents**

1 KNUCKLE BAND 8 PER  
1 TRIPLE ANTIBIOTIC 10 PER  
2 FIRST AID BURN CREAM 6 PER  
1 TRIANGULAR BDG, NON-STERILE  
1 GAUZE PADS, 3" X 3", 4 PER  
1 ADH TAPE, .5" X 2.5 YD, 2 PER  
1 FORCEPS & SCISSORS, 1 EA  
1 GAUZE BANDAGE, 2" X 6 YD, 2 PER  
1 GAUZE COMP, 1 SQ YARD, 1 PER  
1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 BANDAGE COMP, 4" OFFSET, 1 PER  
3 ADHESIVE BDG, PLSTIC, 1"X3"16PER  
1 FINGERTIP BANDAGE, 10 PER  
2 PVP IODINE WIPES 10 PER  
2 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI  
LBL STOCK 6-3/8"X4"  
LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 LABEL NORTH CONTENTS 8X8 ID B  
1 KIT PP 24 UNIT FA

1 LBL CONTENTS ANSI Z308.1-2009 REV B

1 BBL 24U CVR NORTH ID C

1 STING Relief WIPES 10

**4253**

**Z019710-0006L**

1 KNUCKLE BAND 8 PER

1 TRIPLE ANTIBIOTIC 10 PER

1 FIRST AID BURN CREAM 6 PER

1 TRIANGULAR BDG, NON-STERILE

1 GAUZE PADS, 3" X 3", 4 PER

1 ADH TAPE, .5" X 2.5 YD, 2 PER

1 FORCEPS & SCISSORS, 1 EA

1 GAUZE COMP, 1 SQ YARD, 1 PER

1 INSTANT COLD PACK 4" X 6"

1 BUFFERED EYE WASH 1 OZ BTL

1 BANDAGE COMP, 4" OFFSET, 1 PER

2 ADHESIVE BDG, PLSTIC, 1" X 3" 16 PER

1 PVP IODINE WIPES 10 PER

1 STING RELIEF WIPES 10 PER BOX

1 NITRILE GLOVES 2PR BBP

1 FIRST AID GUIDE ASHI

LBL STOCK 6-3/8" X 4"

1 LBL STOCK 6-3/8" X 4"

LBL STOCK 4" X 2-7/8"

1 LBL STOCK 3" X 1-7/8"

1 KIT, PP 16 UNIT FA

1 LBL 16U CVR NORTH ID C

**4254**

**Z019712-0007L**

1 KNUCKLE BAND 8 PER

1 TRIPLE ANTIBIOTIC 10 PER

2 FIRST AID BURN CREAM 6 PER

1 TRIANGULAR BDG, NON-STERILE  
1 GAUZE PADS, 3" X 3", 4 PER  
1 ADH TAPE, .5" X 2.5 YD, 2 PER  
1 FORCEPS & SCISSORS, 1 EA  
1 GAUZE BANDAGE, 2" X 6 YD, 2 PER  
1 GAUZE COMP, 1 SQ YARD, 1 PER  
1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 BANDAGE COMP, 4" OFFSET, 1 PER  
3 ADHESIVE BDG, PLSTIC, 1" X 3" 16 PER  
1 FINGERTIP BANDAGE, 10 PER  
2 PVP IODINE WIPES 10 PER  
1 STING RELIEF WIPES 10 PER BOX  
2 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI  
LBL STOCK 6-3/8" X 4"  
1 LBL STOCK 6-3/8" X 4"  
LBL STOCK 4" X 2-7/8"  
1 LBL STOCK 3" X 1-7/8"  
1 KIT PP 24 UNIT FA  
1 LBL CONTENTS ANSI Z308.1-2009 REV B  
1 BBL 24U CVR NORTH ID C

#### **4284**

#### **SF00002448 kit contents**

1 KNUCKLE BAND 8 PER  
1 TRIPLE ANTIBIOTIC 10 PER  
1 FIRST AID BURN CREAM 6 PER  
1 TRIANGULAR BDG, NON-STERILE  
1 GAUZE PADS, 3" X 3", 4 PER  
1 ADH TAPE, .5" X 2.5 YD, 2 PER  
1 FORCEPS & SCISSORS, 1 EA  
1 GAUZE COMP, 1 SQ YARD, 1 PER

1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 ADHESIVE BDG,PLSTIC,1"X3"16PER  
1 PVP IODINE WIPES 10 PER  
1 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI  
LBL STOCK 6-3/8"X4"  
LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
1 KIT STL 16 UN (HORIZONTAL)  
1 STING Relief WIPES 10

### **4336**

#### **Z019821 kit contents**

1 KNUCKLE BAND 8 PER  
1 TRIPLE ANTIBIOTIC 10 PER  
2 FIRST AID BURN CREAM 6 PER  
1 TRIANGULAR BDG, NON-STERILE  
1 GAUZE PADS, 3" X 3", 4 PER  
1 ADH TAPE, .5" X 2.5 YD, 2 PER  
1 FORCEPS & SCISSORS, 1 EA  
1 GAUZE BANDAGE, 2" X 6 YD,2 PER  
1 GAUZE COMP, 1 SQ YARD, 1 PER  
1 INSTANT COLD PACK 4" X 6"  
1 BUFFERED EYE WASH 1 OZ BTL  
1 BANDAGE COMP, 4" OFFSET, 1 PER  
3 ADHESIVE BDG,PLSTIC,1"X3"16PER  
1 FINGERTIP BANDAGE, 10 PER  
2 PVP IODINE WIPES 10 PER  
1 STING RELIEF WIPES 10 PER BOX  
2 NITRILE GLOVES 2PR BBP  
1 FIRST AID GUIDE ASHI  
LBL STOCK 6-3/8"X4"

LBL STOCK 4"x2-7/8"

1 LBL STOCK 3"x1-7/8"

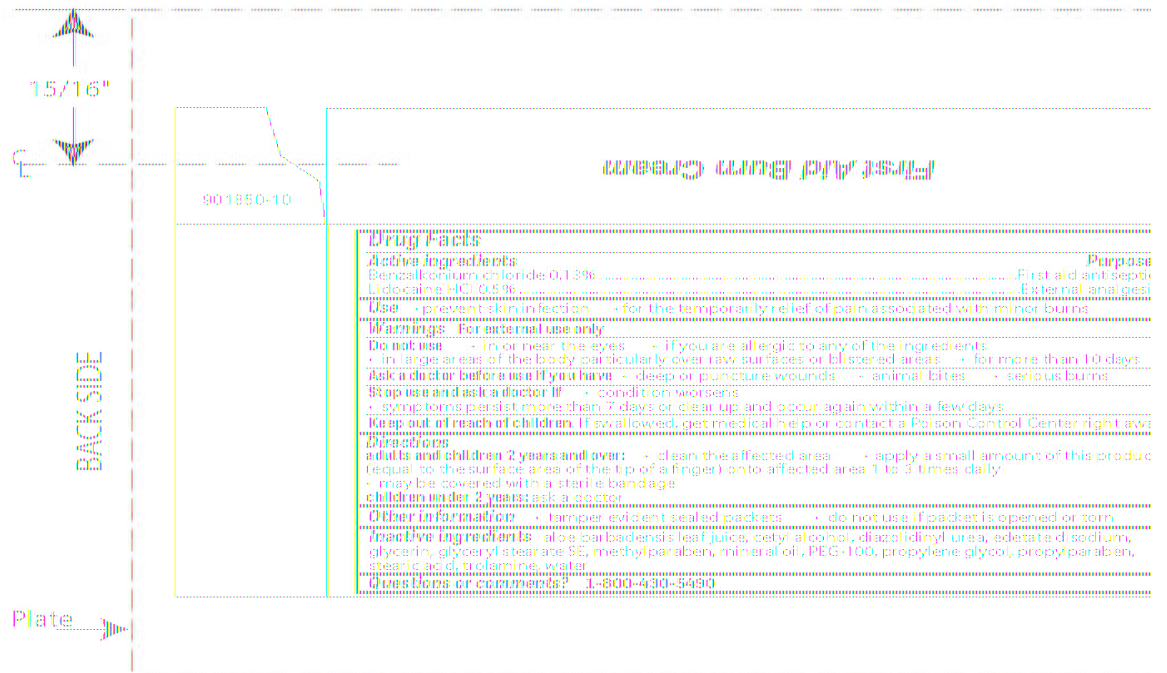
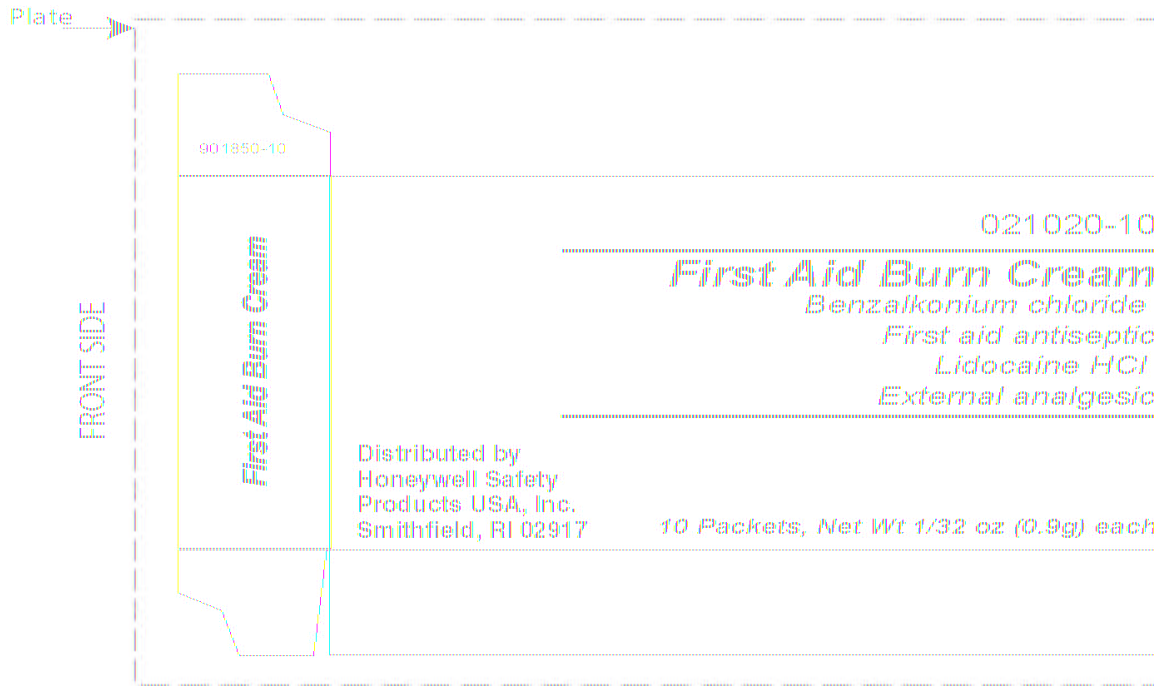
1 LBL CONTS 8"x8",CUSTOM ID B

1 LABEL COVER, GRAINGER Z019821


1 KIT PP 24 UNIT FA

1 LBL CONTENTS ANSI Z308.1-2009 REV B

**First Aid Burn Cream  
Principal Display Panel**



# Principal Display Panel



# eyesaline®

TAMPER-EVIDENT CAP.  
TAPA CON SELLO DE SEGURIDAD.  
BOUCHON INDICATEUR D'INFRACTION.

**LAVAOJOS  
EYESALINE**

Solución  
Isotónico Estéril

**EYESALINE  
EYEWASH**


Sterile  
Isotonic Solution

**LAVAGE  
OCULAIRE  
EYESALINE**

La Solution  
Isotonique Stérile

**16 fl. oz. (473 mL)**

NPN: 80057528



3 64809 45033 7

**Drug Facts (for USA only)**

<b>Active ingredient</b>	<b>Purpose</b>
Sterile water 99% .....	Eyewash

**Uses**  
for flushing the eye to remove loose foreign material, air pollutants, or chlorinated water.

**Warnings**  
**For external use only** - Obtain immediate medical treatment for all open wounds in or near the 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 consult 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.

**Directions**

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

**Inactive ingredients**  
sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

**Questions?** Call 1-800-430-5490  
Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

LABEL #E2-004510 Rev. J

RECORDER / NUEVO PEDIDO / RÉAPPROVISIONNEMENT #E2-00454-0000

space for lot code and supplier part number

PEEL / PELAR / PELEER

**Datos de medicamento (Para EE.UU. solamente)**

<b>Ingrediente Activo</b>	<b>Propósito</b>
Agua estéril 99% .....	Lavaojos

**Usos**  
para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aéros, o agua de cloruro

**Advertencias**  
**Para el uso externo sólo** - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con ninguna superficie. No vuelva a usar. Vez abierto, descarte.

**No se use**

- si la solución se enturbia o cambia de color
- si tiene heridas abiertas en o cerca del ojo, obtenga ayuda médica de inmediato

**Deje de usar y consulte a un médico si:**

- experimenta dolor de ojo
- cambio de visión
- rojez continuo o irritación del ojo
- la condición empeora o persiste

**Manténgase fuera del alcance de los niños.**  
En caso de ingestión accidental, obtenga atención médica o llame a un Centro de control de envenenamiento inmediatamente.

**Instrucciones**

- quítase los lentes de contacto antes de usar la solución
- tuerza la tapa para quitar
- enjuague el área afectada según se necesite
- controle el chorro haciendo presión en la botella
- si es necesario, sigue enjuagado con un lavaojos o ducha de emergencia

**Ingredientes inactivos**  
cloruro de sodio, fosfato de sodio dibásico, fosfato de sodio monobásico

**¿Preguntas?** Llame al 1-800-430-5490  
Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

**Information**

**Usages**  
Pour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmosphériques ou de l'eau chlorée.

**Advertissements**  
**Pour usage externe seulement** - Obtenir immédiatement des soins médicaux pour toutes les plaies ouvertes dans ou près des yeux. Pour éviter toute contamination, ne pas toucher la pointe du récipient à n'importe quelle surface. Ne pas réutiliser. Une fois ouvert, jetez-les.

**Ne pas utiliser**

- si la solution a changé de couleur ou si elle est brouillée
- si vous avez des plaies ouvertes aux yeux ou à proximité, consultez immédiatement un médecin

**Cesser d'utiliser la solution et consulter un médecin**

- vous ressentez une douleur oculaire
- si votre vision change
- rougeur ou irritation persistante des yeux
- condition empire ou persiste

**Garder hors de la portée des enfants.**  
En cas d'ingestion, communiquer immédiatement avec un médecin ou avec un centre antipoison.

**Mode d'emploi**

- enlever les verres de contact avant l'utilisation
- dévisser le bouchon pour l'enlever
- rincer la zone touchée selon les besoins
- ajuster le débit d'écoulement de la solution en augmentant ou en réduisant la pression exercée sur le contenant
- si nécessaire, continuer de rincer avec une solution de rinçage oculaire d'urgence ou une douche

**Ingédients** eau stérile, chlorure de sodium, phosphate dibasique de sodium, phosphate monobasique de sodium

**Des questions?** Faites le 1-800-430-5490  
Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

# Triple Principal Display Panel

**Honeywell**

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

**Honeywell**

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

**Honeywell**

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

**Honeywell**

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

**Drug Facts**

Active Ingredients (in each gram)	Purpose
Bacitracin zinc 400 units	First aid antibiotic
Neomycin sulfate 5 mg (equivalent to 3.5 mg neomycin base)	First aid antibiotic
Polymyxin B sulfate 5000 units	First aid antibiotic

**Uses** • first aid to help prevent infection in:  
• minor cuts • scrapes • burns

**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

**Ask a doctor before use if you have**  
• deep or puncture wounds • animal bites • serious burns  
• conditions such as psoriasis  
• rash or other allergic reaction develops  
• you need to use longer than 1 week

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

**Directions**  
• clean affected area  
• apply a small amount of product (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.

**Other information**  
• keep out of reach of children (39 to 79°F)  
• tamper evident sealed packets  
• do not use if packet is broken or opened

**Inactive ingredient** petrolatum

**Questions?** 1-800-430-5490

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

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

Re-order #231209G

**Caution:**  
  
This package is for industrial use only.  
Not for retail sales.

Keep this and all drugs out of the reach of children.  
In case of ingestion, seek professional assistance or  
contact a poison control center immediately.

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

**TRIPLE  
ANTIBIOTIC  
OINTMENT**  
Bacitracin zinc, Neomycin sulfate,  
Polymyxin B sulfate  
First Aid Antibiotic

**Triple  
Antibiotic  
Ointment**  
To Help Prevent Infection in  
Minor Cuts, Scrapes and Burns

144 Packets  
Net wt. per packet 0.5g (0.02 oz)

771209G Rev A

## Sting Relief Principal Display Panel

**Honeywell**

**032043P**

**Sting Relief  
Wipes**

**Use for:  
Minor Cuts • Scrapes • Insect Bites**

Single Use Pouches  
Saturated Wipes

100 Wipes

**Honeywell**

**032043P**

**Sting Relief  
Wipes**

**Use for:  
Minor Cuts • Scrapes • Insect Bites**

Single Use Pouches  
Saturated Wipes

100 Wipes

**Honeywell**

**825366 Rev B**

**Drug Facts**

Active Ingredients	Purpose
Ethyl alcohol 50.0%	First aid antiseptic
Lidocaine HCl 2.0%	Topical analgesic

**Uses** • first aid to help prevent infection in minor scrapes and temporary relief of itching of insect bites.

**Warnings**  
**For external use only.**  
Flammable, keep away from 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 Poison Control center right away.

**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.

**Inactive Ingredients**  
benzalkonium chloride, menthol, purified water

**Questions or comments?** 1-800-430-5490

**Honeywell**

**032043P**

**Sting Relief  
Wipes**

**Use for:  
Minor Cuts • Scrapes • Insect Bites**

Single Use Pouches  
Saturated Wipes

100 Wipes

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

USA  
1-800-430-5490  
0-401-943-4400

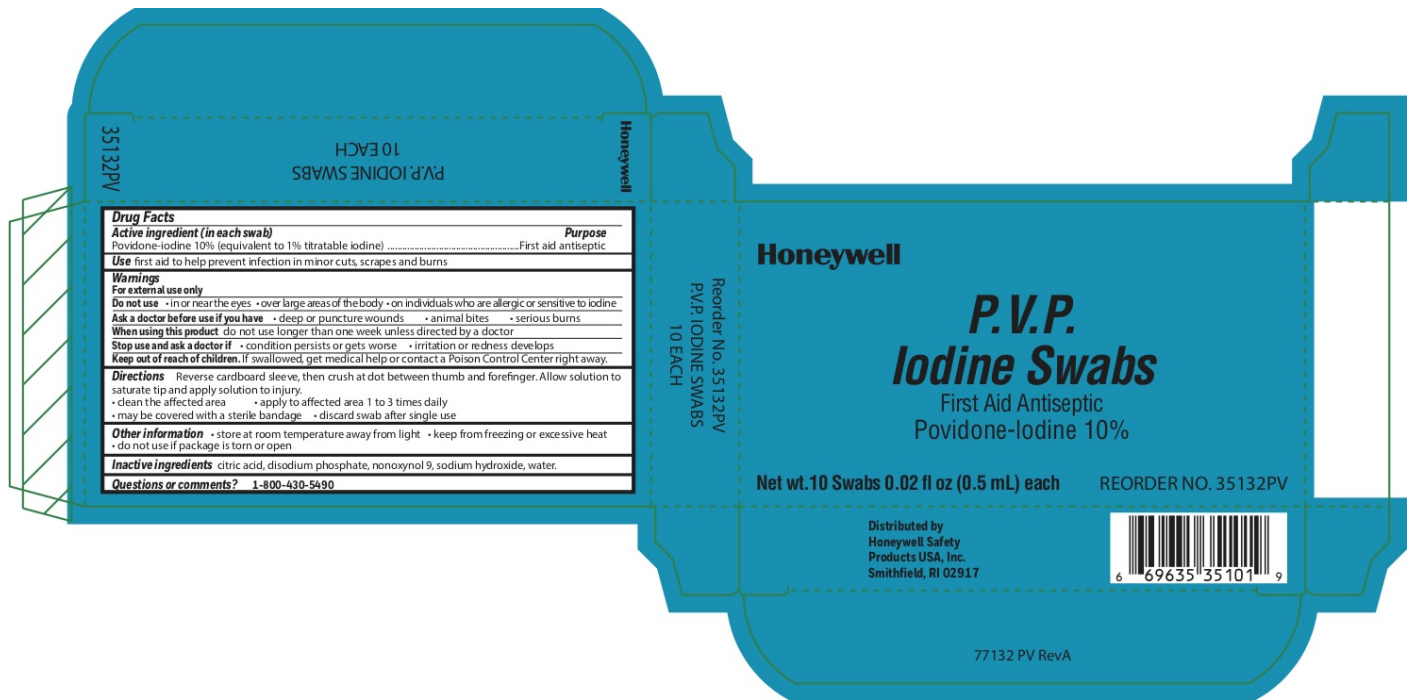
www.honeywellsafety.com

8 21812 01264 1

Made in USA  
Packaged in Mexico

## PVP Principal Display Panel





**4197 Kit Label**  
**019710-0006L**

**Honeywell**

**First Aid Kit**

**Unitized**

COMPLIANT

Latex Free

**For Up To 16 People**

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

771025 Rev. D

**4198 KIT LABEL**  
**019711-0006L**

# Honeywell

## First Aid Kit

### Unitized



For Up To 16 People



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

**4199 Kit Label**  
**019712-0007L**

# Honeywell

# First Aid Kit

## Unitized



For Up To 16 People



Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02887

777035 Dsw.D

4253 Kit Label  
Z019710-0006L

777025 Rev. D  
4 color process, pms 072 blue, and pms 186 red

# Honeywell

## First Aid Kit

### Unitized

ANSI Z308.1-2009  
COMPLIANT

Latex Free

For Up To 16 People



777025 Rev. D

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

**4254 kit Label**  
**Z019712-0007L**

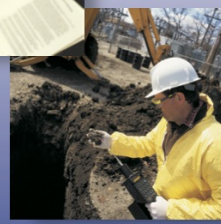
777026 Rev. D  
4 color process, pms 072 blue, and pms 186 red

# Honeywell



# First Aid Kit

## Unitized



For Up To 25 People

777026 Rev. D

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

**4284 Kit Label**  
**Sf00002448**

# **FIRST AID**

---



**(800) 442-8839**

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

**4336 Kit Label  
Z019821**

# FIRST AID

GENERAL PURPOSE, UNITIZED  
25 PERSON



**GRAINGER**<sup>®</sup>  
||| FOR THE ONES WHO GET IT DONE

GRAINGER.COM<sup>®</sup>

47001726A

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

## 4253 FIRST AID KIT

4253 first aid kit kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4253
---------------------	----------------	---------------------------	---------------

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4253-	1 in 1 KIT; Type 0: Not a Combination	10/19/2019	

**Quantity of Parts**

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	5.4 g
Part 2	1 BOTTLE	30 mL
Part 3	10 PACKET	9 g
Part 4	10 POUCH	4 mL
Part 5	10 POUCH	3 mL

**Part 1 of 5**

**FIRST AID BURN**  
 benzalkonium chloride, lidocaine hydrochloride cream

**Product Information**

<b>Item Code (Source)</b>	NDC:0498-0903
<b>Route of Administration</b>	TOPICAL

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	



**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

**Part 2 of 5****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
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (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	Application Number or Monograph	Marketing Start	Marketing End
-----------	---------------------------------	-----------------	---------------

Category	Citation	Date	Date
OTC Monograph Drug	M018	12/18/2018	

## Part 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

#### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

#### 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 4 of 5

### 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
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

#### Inactive Ingredients

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

#### 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		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (UNII: 059QF0K00R)	

**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	

**Marketing Information**

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

**4197 FIRST AID KIT**

4197 first aid kit kit

**Product Information****Product Type** HUMAN OTC DRUG**Item Code (Source)**

NDC:0498-4197

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4197-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	5.4 g
Part 2	1 BOTTLE	30 mL
Part 3	10 PACKET	9 g
Part 4	10 POUCH	4 mL
Part 5	10 POUCH	3 mL

### Part 1 of 5

#### FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

### Product Information

Item Code (Source)	NDC:0498-0903
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	

**DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)**

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

### Part 2 of 5

### 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
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (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
OTC Monograph Drug	M018	12/18/2018	

## Part 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (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	Application Number or Monograph	Marketing Start	Marketing End
-----------	---------------------------------	-----------------	---------------

Category	Citation	Date	Date
unapproved drug other		09/19/2018	

## Part 4 of 5

### 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
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>WATER</b> (UNII: 059QF0KO0R)	

#### 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		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (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	

### Marketing Information

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

## 4284 FIRST AID KIT

4284 first aid kit kit

### Product Information

**Product Type** HUMAN OTC DRUG

**Item Code (Source)**

NDC:0498-4284

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4284-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	5.4 g
Part 2	1 BOTTLE	30 mL
Part 3	10 PACKET	9 g
Part 4	10 POUCH	4 mL
Part 5	10 POUCH	3 mL

## Part 1 of 5

### FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

## Product Information

Item Code (Source)	NDC:0498-0903
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	

<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

<b>Part 2 of 5</b>
<b>EYESALINE EMERGENCY EYEWASH</b>
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
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)	

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

01	Product		
----	---------	--	--

## Marketing Information

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

## Part 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (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 4 of 5

### 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
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

## Inactive Ingredients

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

## 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		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (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	

#### Marketing Information

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

## 4198 FIRST AID KIT

4198 first aid kit kit

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4198
---------------------	----------------	---------------------------	---------------

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4198-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	5.4 g
Part 2	1 BOTTLE	30 mL
Part 3	10 PACKET	9 g
Part 4	10 POUCH	4 mL
Part 5	10 POUCH	3 mL

### Part 1 of 5

#### FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0903
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	

<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

## Part 2 of 5

### EYESALINE EMERGENCY EYEWASH

purified water liquid

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0100
<b>Route of Administration</b>	OPHTHALMIC

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
---	-----------	---------------------	----------------------	--------------------



#	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 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (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-	0.9 g in 1 PACKET; Type 0: Not a Combination		

35	Product		
----	---------	--	--

### Marketing Information

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

### Part 4 of 5

### 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
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

### Inactive Ingredients

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

### 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		12/23/2017	

## Part 5 of 5

### 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: 059QF0K00R)	

#### 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	

#### Marketing Information

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

## 4199 FIRST AID KIT

4199 first aid kit kit

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4199
---------------------	----------------	---------------------------	---------------

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4199-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
<b>Part 1</b>	12 PACKET	10.8 g
<b>Part 2</b>	1 BOTTLE	30 mL
<b>Part 3</b>	10 PACKET	9 g
<b>Part 4</b>	10 POUCH	4 mL
<b>Part 5</b>	20 POUCH	6 mL

## Part 1 of 5

### FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

## Product Information

<b>Item Code (Source)</b>	NDC:0498-0903
<b>Route of Administration</b>	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	

<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

## Part 2 of 5

### EYESALINE EMERGENCY EYEWASH

purified water liquid

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0100
<b>Route of Administration</b>	OPHTHALMIC

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (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
OTC Monograph Drug	M018	12/18/2018	

## Part 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

## Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

## Packaging

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

#	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 4 of 5

### 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
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

## Inactive Ingredients

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

## 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	Application Number or Monograph	Marketing Start	Marketing End
-----------	---------------------------------	-----------------	---------------

Category	Citation	Date	Date
unapproved drug other		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (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	

#### Marketing Information

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



# 4254 FIRST AID KIT

4254 first aid kit kit

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4254
---------------------	----------------	---------------------------	---------------

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4254-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	12 PACKET	10.8 g
Part 2	1 BOTTLE	30 mL
Part 3	10 PACKET	9 g
Part 4	10 POUCH	4 mL
Part 5	20 POUCH	6 mL

## Part 1 of 5

### FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

## Product Information

<b>Item Code (Source)</b>	NDC:0498-0903
---------------------------	---------------

<b>Route of Administration</b>	TOPICAL
--------------------------------	---------

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	

<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

## Part 2 of 5

### EYESALINE EMERGENCY EYEWASH

purified water liquid

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0100
<b>Route of Administration</b>	OPHTHALMIC

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (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
OTC Monograph Drug	M018	12/18/2018	

## Part 3 of 5

### 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>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ 07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (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 4 of 5

### 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
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## 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		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (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	

## Marketing Information

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

other

10/10/2010

**4336 FIRST AID KIT**

4336 first aid kit kit

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4336
---------------------	----------------	---------------------------	---------------

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4336-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

**Quantity of Parts**

Part #	Package Quantity	Total Product Quantity
<b>Part 1</b>	12 PACKET	10.8 g
<b>Part 2</b>	1 BOTTLE	30 mL
<b>Part 3</b>	10 PACKET	9 g
<b>Part 4</b>	10 POUCH	4 mL
<b>Part 5</b>	20 POUCH	6 mL

**Part 1 of 5****FIRST AID BURN**

benzalkonium chloride, lidocaine hydrochloride cream

**Product Information**

<b>Item Code (Source)</b>	NDC:0498-0903
---------------------------	---------------

<b>Route of Administration</b>	TOPICAL
--------------------------------	---------

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII: 98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
-----------------	----------

<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		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		12/20/2017	

### Part 2 of 5

### EYESALINE EMERGENCY EYEWASH

purified water liquid

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0100
<b>Route of Administration</b>	OPHTHALMIC

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE</b> (UNII: 593YOG76RN)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (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
OTC Monograph Drug	M018	12/18/2018	

### Part 3 of 5

### TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0750
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [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
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	



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

### 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 4 of 5

### STING RELIEF PAD

ethyl alcohol, lidocaine swab

### Product Information

<b>Item Code (Source)</b>	NDC:0498-0733
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0733-	0.4 mL in 1 POUCH; Type 0: Not a Combination		

00	Product		
----	---------	--	--

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/23/2017	

## Part 5 of 5

### 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
<b>POVIDONE-IODINE</b> (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>NONOXYNOL-9</b> (UNII: 48Q180SH9T)	
<b>WATER</b> (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	

## Marketing Information

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

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

Revised: 1/2024

Honeywell Safety Products USA, INC