

**4357 FIRST AID KIT- 4357 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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**4357 First Aid Kit (Burn Jel, alcohol wipe, Burn Sp WS, Triple, EW, HC cr, Foille- 6832649 )**

**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

**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

## **Hydrocortisone**

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

Hydrocortisone acetate (equivalent to Hydrocortisone 1%)

## **Hydrocortisone**

### ***Purpose***

Anti-itch cream

## **Hydrocortisone**

### ***Uses***

- for the temporary relief of itching associated with minor skin irritations and rashes

## **Hydrocortisone**

### ***Warnings***

#### **For external use only**

#### **Ask a doctor before use if**

- you are using any other hydrocortisone product

#### **When using the product**

- avoid contact with eyes
- do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash

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

- condition worsens
- condition persists for more than 7 days
- condition clears up and recurs within a few days

#### **Keep out of reach of children**

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

## **Hydrocortisone**

### ***Directions***

- adults and children 2 years and older:
- clean the affected area
- apply to the area not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor

## **Hydrocortisone**

### ***Other information***

- store at room temperature (do not freeze)

## **Hydrocortisone**

### ***Inactive ingredients***

cetyl alcohol, citric acid, diazolidinyl urea, edetate disodium, glycerin, glyceryl monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol, propylparaben, purified water, stearic acid, trolamine

## **Hydrocortisone**

### ***Questions or Comments?***

1-800-430-5490

### **Foille**

#### ***Active ingredient***

Benzocaine 5.0% (w/w)

Chloroxylenol 0.1% (w/w)

### **Foille**

#### ***Purpose***

External analgesic

Antiseptic

### **Foille**

#### ***Uses***

- For the temporary relief of pain associated with burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritations.
- First aid to help prevent infection in minor cuts, scrapes and burns.

### **Foille**

#### ***Warnings***

#### **For external use only**

#### **When using this product**

- avoid contact with the eyes

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

- condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days.
- Do not apply over large areas of the body. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.

#### **Keep out of reach of children.**

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

### **Foille**

#### ***Directions***

- clean the affected area.
- adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- children under 2 years of age: consult a physician.

### **Foille**

#### ***Other information***

Avoid contact with clothing  
Foilie may stain certain fabrics

## **Foilie**

### ***Inactive ingredients***

beeswax, benzyl alcohol, calcium disodium EDTA, calcium hydroxide, ceresin, eugenol, hydrogenated vegetable oil, maleic anhydride, mono- and di-glycerides, PEG-32, purified water, sodium borate, sodium lauryl sulfate, zea mays (corn) oil.

## **Burn Jel**

### ***Active ingredient***

Lidocaine HCl 2.0 %v

## **Burn Jel**

### ***Purpose***

External analgesic

## **Burn Jel**

### ***Uses***

- temporarily relieves pain due to minor burns

## **Burn Jel**

### ***Warnings***

#### **For external use only**

#### **Do not use**

- on large areas of the body, particularly over raw surfaces or blistered areas

#### **When using this product**

- avoid contact with eyes

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

- the condition gets worse
- symptoms persist for more than 7 days
- condition clears up and recurs within a few days

#### **Keep out of reach of children**

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

## **Burn Jel**

### ***Directions***

- adults and children 2 years of age and older; apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor
- you may report a serious reaction to this product to 800-430-5490

## **Burn Jel**

### ***Other information***

- store at room temperature - do not use if opened or torn

## **Burn Jel**

### ***Inactive ingredients***

carbopol 940, carbopol 1342, diazolidinyl urea, glycerin, melaleuca alternifolia (tea tree) leaf oil, methylparaben, octoxynol-9, propylene glycol, propylparaben, trolamine, water

## **Burn Jel**

### ***Questions***

1-800-430-5490

**4357**

### **6832649 KIT CONTENTS**

1 BLUE DETEC FNGERTP 8 WVN 25/B  
1 BLUE DETEC KNUCKLE WVN 40/BX  
1 BLUE DETEC 1X3 WVN 100/BX  
1 INSTANT COLD PACK 4" X 6"  
5 BURN JEL 1/8 OZ, 6 PER  
1 ADHESIVE TAPE W/P 1/2"X 5 YD  
1 FIRST AID GUIDE ASHI  
1 GAUZE CLEAN-WRAP BDGE N/S 2"  
1 GZE PADS STERILE 2"X 2" 25'S  
1 GZE PADS STERILE 3"X 3" 25'S  
1 CO-FLEX BANDAGE 2"X 5YDS TAN  
1 ALCOHOL WIPES 50'S  
1 TRIPLE BIOTIC .5 GRAM PKT 20  
1 HYDROCORTISONE 1% .9 GRM 20'S  
1 4OZ BFS EYEWASH TRILINGUAL BOTTLE  
1 SCISSOR BDGE 4" RED PLS HDL  
1 SPLINTER OUT 10 PIECES/PK  
1 KIT TWEEZER 3 1/2" SLANTED  
1 F A KIT EMPTY BLANK 140  
1 LBL STOCK 6-3/8"X4"  
1 LBL STOCK 4"X2-7/8"  
1 LBL STOCK 3"x1-7/8"  
2 PR LRG NITRILE GLVES ZIP BAG  
5 WATER-JEL BURN DRESSING 4 X 4

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

**Honeywell**

eyesaline®

**LAVAOJOS  
EYESALINE**

Solución  
Isotónico Estéril

**EYESALINE  
EYEWASH**

Sterile  
Isotonic Solution

**LAVAGE  
OCULAIRE  
EYESALINE**

La Solution  
Isotonique Stérile

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

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



NPN: 80057528  
64809 11 45033 117

**Drug Facts (for USA only)**

| Active ingredient | Purpose |
|-------------------|---------|
| 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

REORDER / NUEVO PEDIDO / REAPPROVISIONNEMENT #32-00045-0000

space for lot code and supplier part number

PEEL / PELAR / PELE

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

| Ingrediente Activo | Propósito |
|--------------------|-----------|
| Agua estéril 99%   | Lavaojos  |

**Usos**  
para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aéreos, 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 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**

- quítese 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

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

SHAKE WELL BEFORE USING

**NORTH**<sup>®</sup>

by Honeywell

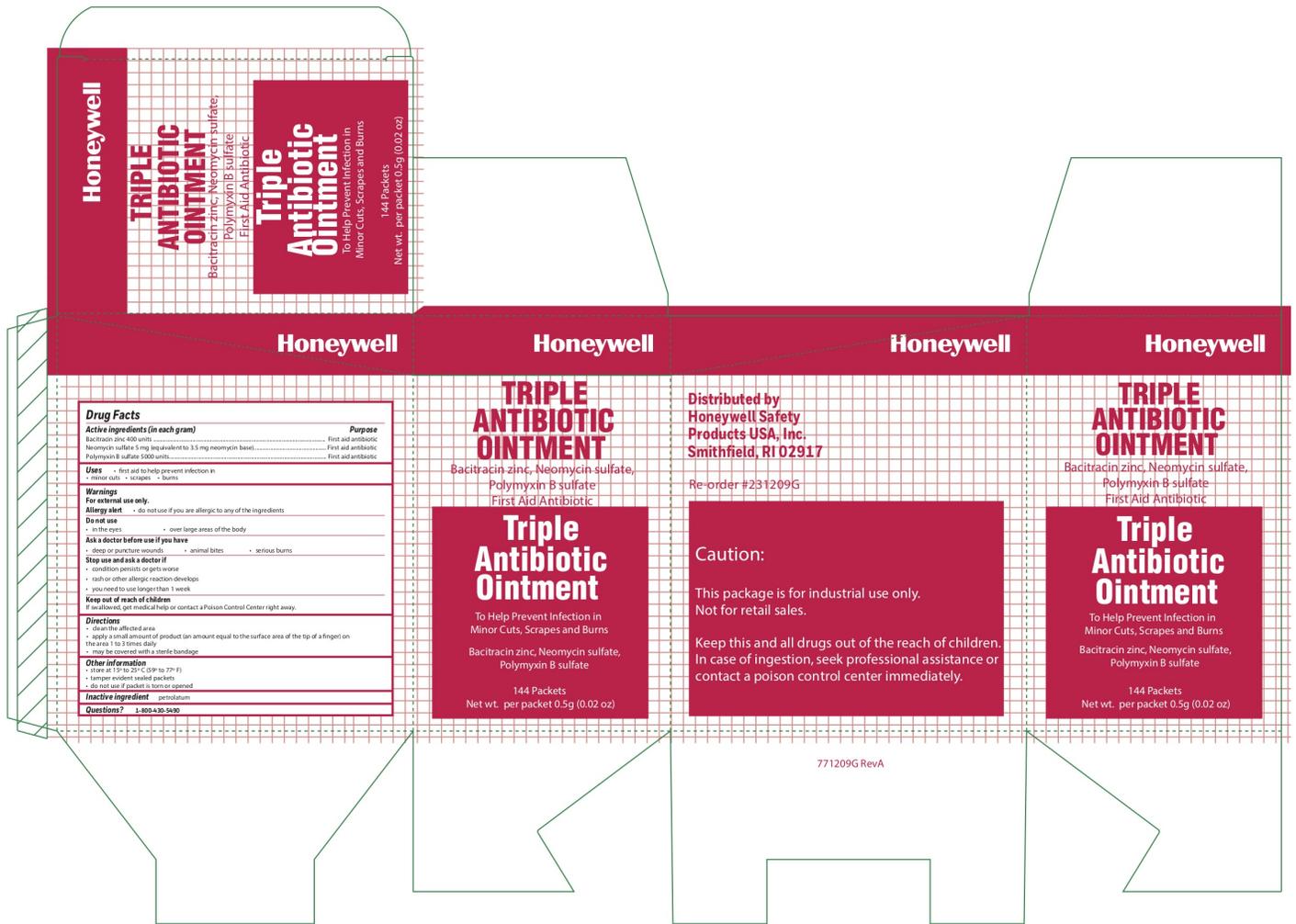
**BURN SPRAY**

*Water Soluble*

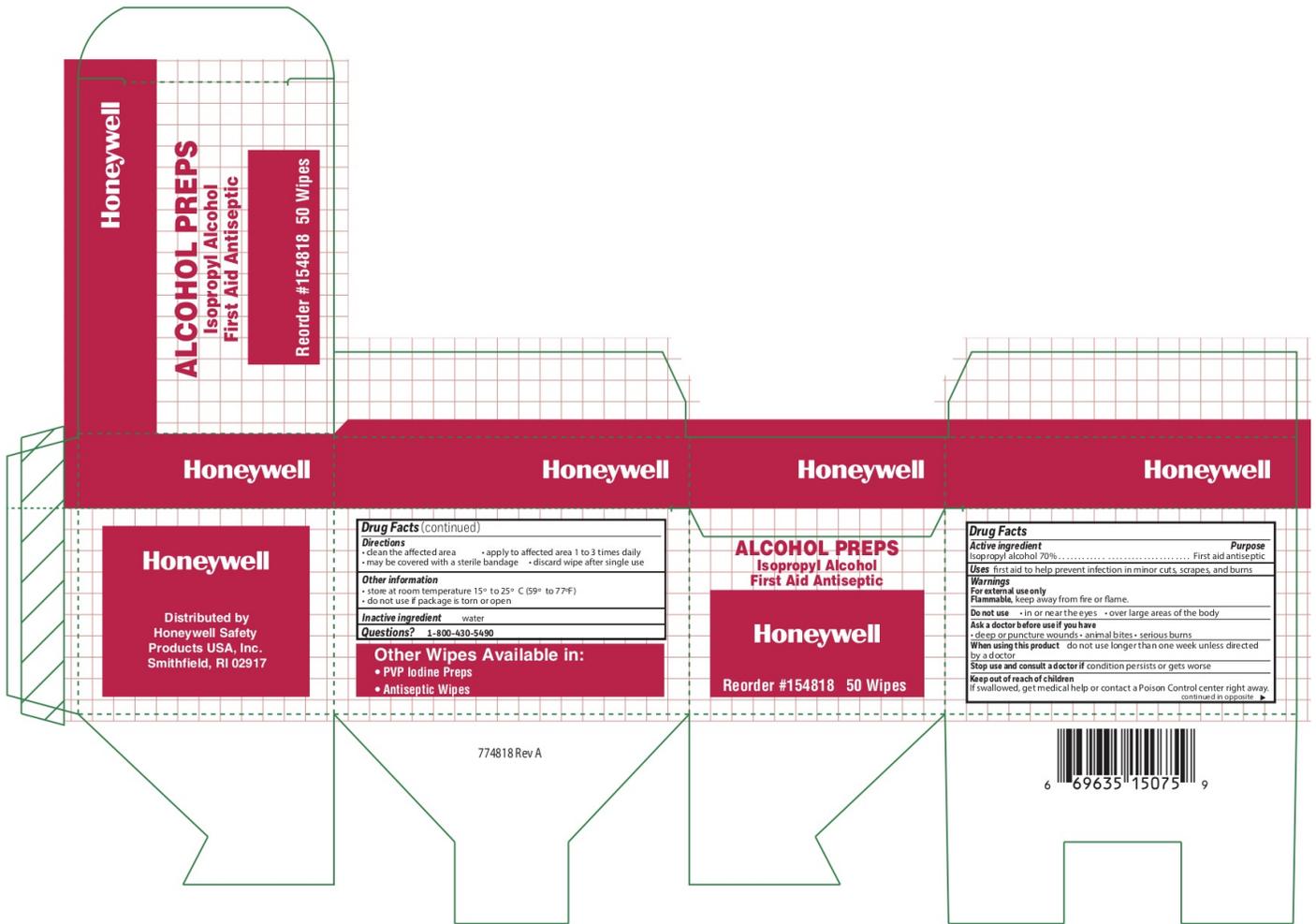
Provides antiseptic treatment  
and helps relieve the pain of minor burns  
and sunburn.

**CAUTION: FLAMMABLE**  
Contents under pressure  
Read warning on back panel.

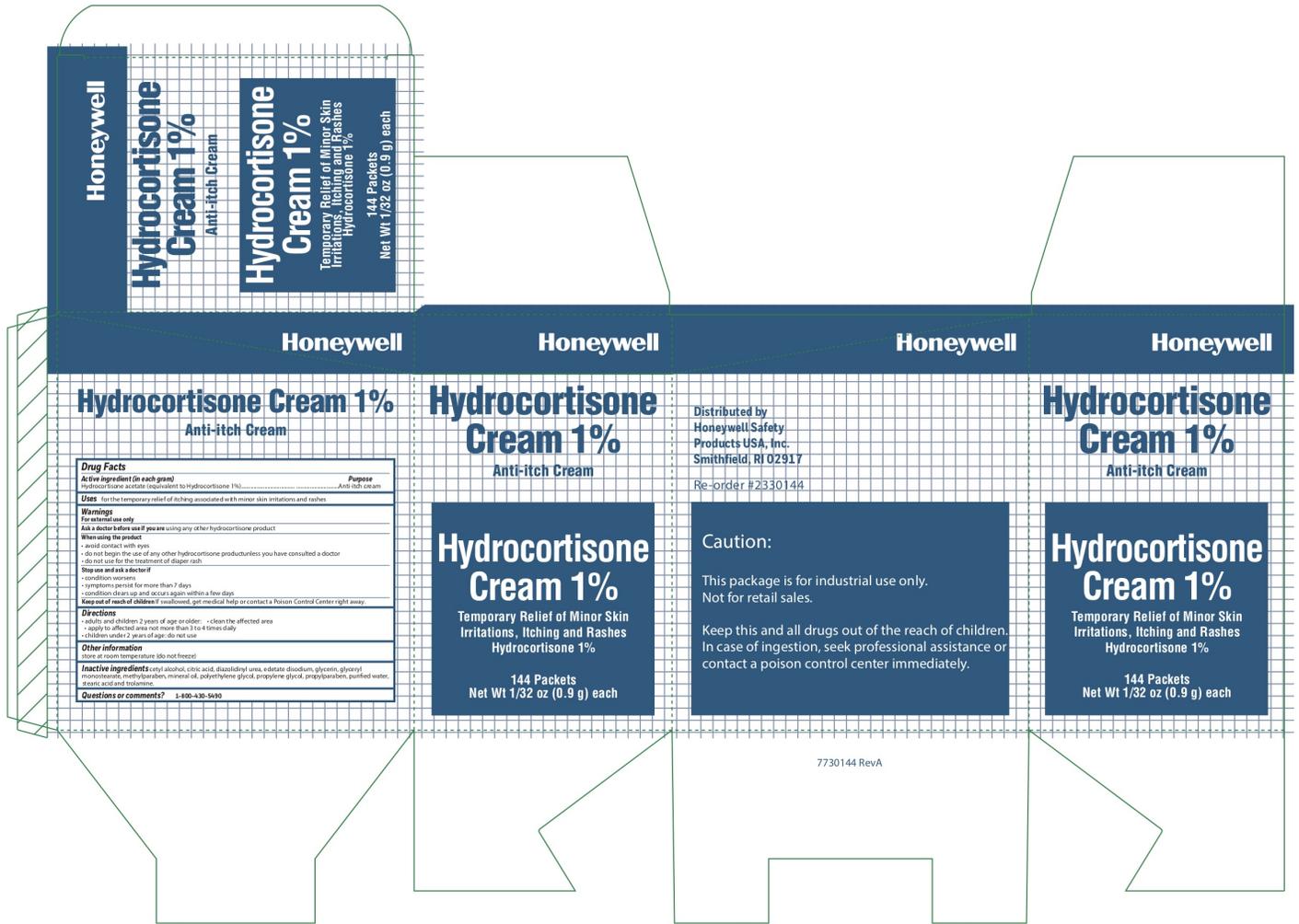
# Principal Display Panel



# Alcohol Principal Display Panel



**Hydrocortisone**  
*Principal Display Panel*



**Honeywell**  
**Hydrocortisone Cream 1%**  
 Anti-itch Cream

**Hydrocortisone Cream 1%**  
 Temporary Relief of Minor Skin Irritations, Itching and Rashes  
 Hydrocortisone 1%  
 144 Packets  
 Net Wt 1/32 oz (0.9 g) each

**Honeywell**

**Honeywell**

**Honeywell**

**Honeywell**

**Hydrocortisone Cream 1%**  
 Anti-itch Cream

| Drug Facts  | Purpose         |
|---|-----------------|
| <b>Active ingredient (in each gram)</b><br>hydrocortisone acetate (equivalent to hydrocortisone 1%)   | Anti-itch cream |
| <b>Uses:</b> For temporary relief of itching associated with minor skin irritations and rashes.   |                 |
| <b>Warnings:</b><br>For external use only.<br>Ask a doctor before use if you are using any other hydrocortisone product.  |                 |
| <b>When using the product:</b><br>- avoid contact with eyes<br>- do not begin the use of any other hydrocortisone product unless you have consulted a doctor<br>- do not use for the treatment of diaper rash   |                 |
| <b>Stop use and ask a doctor if:</b><br>- condition worsens<br>- symptoms persist for more than 7 days<br>- condition clears up and occurs again within a few days  |                 |
| <b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.   |                 |
| <b>Directions:</b><br>- adults and children 2 years of age or older: - clean the affected area<br>- apply to affected area not more than 3 to 4 times daily<br>- children under 2 years of age: do not use  |                 |
| <b>Other information:</b><br>Store at room temperature (do not freeze)  |                 |
| <b>Inactive ingredients:</b> cetyl alcohol, citric acid, disodium edetate, disodium glycinate, glycerin, glyceryl monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol, propylparaben, purified water, stearic acid and tolanolene |                 |
| <b>Questions or comments?</b> 1-800-430-5430  |                 |

**Hydrocortisone Cream 1%**  
 Anti-itch Cream

**Hydrocortisone Cream 1%**  
 Temporary Relief of Minor Skin Irritations, Itching and Rashes  
 Hydrocortisone 1%  
 144 Packets  
 Net Wt 1/32 oz (0.9 g) each

Distributed by  
**Honeywell Safety Products USA, Inc.**  
 Smithfield, RI 02917  
 Re-order #2330144

**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.

**Hydrocortisone Cream 1%**  
 Anti-itch Cream

**Hydrocortisone Cream 1%**  
 Temporary Relief of Minor Skin Irritations, Itching and Rashes  
 Hydrocortisone 1%  
 144 Packets  
 Net Wt 1/32 oz (0.9 g) each

7730144 RevA

**Foile**  
**Principal Display Panel**

**Drug Facts**

**Active ingredients Purpose**

Benzocaine 5.0% (w/w) External Analgesic  
Chloroxylenol 0.1% (w/w) Antiseptic

**Uses**

- For the temporary relief of pain associated with burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritations.
- First aid to help prevent infection in minor cuts, scrapes and burns.

**Warnings**

- For external use only.
- When using this product:
  - Avoid contact with the eyes.
- Stop use and ask a doctor if:
  - condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days.
  - Do not apply over large areas of the body. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.

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

**Directions**

- Clean the affected area.
- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 2 years of age: consult a physician.



**MEDICATED FIRST AID OINTMENT**

Fast, Soothing Relief Of Pain Due To:  
**Cuts & Scrapes • Minor Burns**  
Sunburn • Insect Bites

NDC 10157-9302-4



**MEDICATED FIRST AID OINTMENT**

Fast, Soothing Relief Of Pain Due To:  
**Cuts & Scrapes • Minor Burns**  
Sunburn • Insect Bites

NET WT.  
1 oz (28g)

**Drug Facts (continued)**

**Other information**

- Avoid contact with clothing. Foille may stain certain fabrics.

**Inactive ingredients**

beeswax, benzyl alcohol, calcium disodium EDTA, calcium hydroxide, cerasin, eugenol, hydrogenated vegetable oil, maleic anhydride mono- and di-glycerides, PEG-3-2, purified water, sodium borate, sodium lauryl sulfate, zea mays (corn) oil.



For the temporary relief of pain due to scrapes, cuts, minor burns, sunburn, non-poisonous insect bites, and minor skin irritation, Foille First Aid Ointment stops pain on contact and lets you resume normal activities right away. Foille has a medicated oil-based formula that helps heal and prevent infection.

SATISFACTION GUARANTEED  
**Blistex**

©2009 Blistex Inc.  
P.O. Box 5392  
Oak Brook, IL 60522-5392  
#39013

♻️ Carton is 100% Recyclable.

**Burn Jel**  
*Principal Display Panel*





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

## 4357 FIRST AID KIT

4357 first aid kit kit

### Product Information

|                     |                |                           |               |
|---------------------|----------------|---------------------------|---------------|
| <b>Product Type</b> | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:0498-4357 |
|---------------------|----------------|---------------------------|---------------|

### Packaging

| # | Item Code        | Package Description                           | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0498-4357-01 | 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> | 20 PACKET        | 18 g                   |
| <b>Part 2</b> | 1 BOTTLE         | 118 mL                 |
| <b>Part 3</b> | 1 CAN            | 85 g                   |
| <b>Part 4</b> | 20 PACKET        | 10 g                   |
| <b>Part 5</b> | 50 POUCH         | 20 mL                  |
| <b>Part 6</b> | 20 PACKET        | 18 g                   |
| <b>Part 7</b> | 6 TUBE           | 84 g                   |

**Part 1 of 8****HYDROCORTISONE**

anti-itch cream

**Product Information**

|                                |               |
|--------------------------------|---------------|
| <b>Item Code (Source)</b>      | NDC:0498-0801 |
| <b>Route of Administration</b> | TOPICAL       |

**Active Ingredient/Active Moiety**

| Ingredient Name  | Basis of Strength      | Strength        |
|--|------------------------|-----------------|
| HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ) | HYDROCORTISONE ACETATE | 1 g<br>in 100 g |

**Inactive Ingredients**

| Ingredient Name                                     | Strength |
|---|----------|
| STEARIC ACID (UNII: 4ELV7Z65AP)                     |          |
| TROLAMINE (UNII: 9O3K93S3TK)                        |          |
| METHYLPARABEN (UNII: A2I8C7HI9T)                    |          |
| WATER (UNII: 059QF0KO0R)                            |          |
| EDETATE DISODIUM (UNII: 7FLD91C86K)                 |          |
| LIGHT MINERAL OIL (UNII: N6K5787QVP)                |          |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)                 |          |
| DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)                |          |
| PROPYLPARABEN (UNII: Z8IX2SC1OH)                    |          |
| GLYCERIN (UNII: PDC6A3C0OX)                         |          |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)          |          |
| GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)            |          |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) |          |
| CETYL ALCOHOL (UNII: 936JST6JCN)                    |          |

**Packaging**

| # | Item Code        | Package Description                                  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0801-35 | 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 not final | part348                                  | 10/15/2019           |                    |

## Part 2 of 8

### 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 3 of 8

### 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 4 of 8****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>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:0498-0750-36 | 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 5 of 8

#### 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: 059QF0K00R) |          |

### 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           |                    |

## Part 6 of 8

### HYDROCORTISONE

anti-itch cream ointment

## Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:0498-0800 |
| Route of Administration | TOPICAL       |

## Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength      | Strength        |
|--|------------------------|-----------------|
| HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W14X0X7BPJ) | HYDROCORTISONE ACETATE | 1 g<br>in 100 g |

## Inactive Ingredients

| Ingredient Name                                     | Strength |
|---|----------|
| STEARIC ACID (UNII: 4ELV7Z65AP)                     |          |
| TROLAMINE (UNII: 9O3K93S3TK)                        |          |
| METHYL PARABEN (UNII: A2I8C7HI9T)                   |          |
| WATER (UNII: 059QF0KO0R)                            |          |
| EDETATE DISODIUM (UNII: 7FLD91C86K)                 |          |
| LIGHT MINERAL OIL (UNII: N6K5787QVP)                |          |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)                 |          |
| DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)                |          |
| PROPYL PARABEN (UNII: Z8IX2SC1OH)                   |          |
| GLYCERIN (UNII: PDC6A3C0OX)                         |          |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)          |          |
| GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)            |          |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) |          |
| CETYL ALCOHOL (UNII: 936JST6JCN)                    |          |

## Packaging

| # | Item Code        | Package Description                                  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0800-35 | 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 not final | part348                                  | 03/06/2013           | 10/15/2019         |

## Part 7 of 8

### BLISTEX FOILLE MEDICATED FIRST AID

benzocaine and chloroxylenol ointment

#### Product Information

**Item Code (Source)** NDC:10157-9302

**Route of Administration** TOPICAL

#### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength       |
|--|-------------------|----------------|
| BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)       | BENZOCAINE        | 5 g in 100 g   |
| CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q) | CHLOROXYLENOL     | 0.1 g in 100 g |

#### Inactive Ingredients

| Ingredient Name                                       | Strength |
|---|----------|
| SODIUM LAURYL SULFATE (UNII: 368GB5141J)              |          |
| BENZYL ALCOHOL (UNII: LKG8494WBH)                     |          |
| EDETATE CALCIUM DISODIUM ANHYDROUS (UNII: 8U5D034955) |          |
| CERESIN (UNII: Q1LS2UJO3A)                            |          |
| EUGENOL (UNII: 3T8H1794QW)                            |          |
| MALEIC ANHYDRIDE (UNII: V5877ZJZ25)                   |          |
| POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)           |          |
| CORN OIL (UNII: 8470G57WFM)                           |          |
| SODIUM BORATE (UNII: 91MBZ8H3QO)                      |          |
| YELLOW WAX (UNII: 2ZA36H0S2V)                         |          |
| CALCIUM HYDROXIDE (UNII: PF5DZW74VN)                  |          |
| WATER (UNII: 059QF0K00R)                              |          |

#### Packaging

| # | Item Code | Package Description                               | Marketing Start Date | Marketing End Date |
|---|-----------|---|----------------------|--------------------|
| 1 |           | 14 g in 1 TUBE; 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                                  | 03/05/2013           |                    |

## Part 8 of 8

### BURN JEL

gel for burns gel

#### Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:0498-0203 |
| Route of Administration | TOPICAL       |

#### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength                 | Strength        |
|---|-----------------------------------|-----------------|
| LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98P200987) | LIDOCAINE HYDROCHLORIDE ANHYDROUS | 2 g<br>in 100 g |

#### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| TROLAMINE (UNII: 9O3K93S3TK)   |          |
| CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E) |          |
| CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31) |          |
| PROPYLPARABEN (UNII: Z8IX2SC1OH)   |          |
| OCTOXYNOL-9 (UNII: 7JPC6Y25QS)   |          |
| TEA TREE OIL (UNII: VIF565UC2G)  |          |
| DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)   |          |
| EDETATE DISODIUM (UNII: 7FLD91C86K)  |          |
| DIPROPYLENE GLYCOL (UNII: E107L85C40)  |          |
| GLYCERIN (UNII: PDC6A3C0OX)  |          |
| METHYLPARABEN (UNII: A2I8C7HI9T)   |          |

#### Packaging

| # | Item Code        | Package Description                                  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0498-0203-00 | 3.5 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 not final | part348                                  | 09/19/2018           |                    |

#### 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      |
|--------------|---------|-----------|--------------------------|
| Blistex Inc. |         | 005126354 | manufacture(10 157-9302) |

### Establishment

| Name                               | Address | ID/FEI    | Business Operations |
|------------------------------------|---------|-----------|---------------------|
| Honeywell Safety Products USA, INC |         | 079287321 | pack(0498-4357)     |

### Establishment

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

### Establishment

| Name                   | Address | ID/FEI    | Business Operations                                     |
|------------------------|---------|-----------|---|
| Water-Jel Technologies |         | 155522589 | manufacture(0498-0750, 0498-0800, 0498-0203, 0498-0801) |

### 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) |

Revised: 10/2019

Honeywell Safety Products USA, INC