

**4223 FIRST AID KIT - 4223 first aid kit**

**4275 FIRST AID KIT - 4275 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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**4223, 4275 First Aid Kit (FABC, HC cr, BZK wipes - 34575H, SF00001582)**

### **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

### **BZK Wipes**

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

Benzalkonium chloride 0.13% w/v

### **BZK Wipes**

#### ***Purpose***

First aid antiseptic

### **BZK**

#### ***Uses***

Antiseptic cleansing of face, hands, and body without soap and water

### **BZK**

#### ***Warnings***

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

#### **Do not use**

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

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

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

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

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

### **BZK**

#### ***Directions***

- tear open packet and use as a washcloth

### **BzK**

#### ***Inactive ingredient***

water

### **BZK**

#### ***Questions***

1-800-430-5490

### **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**

- 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

## **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

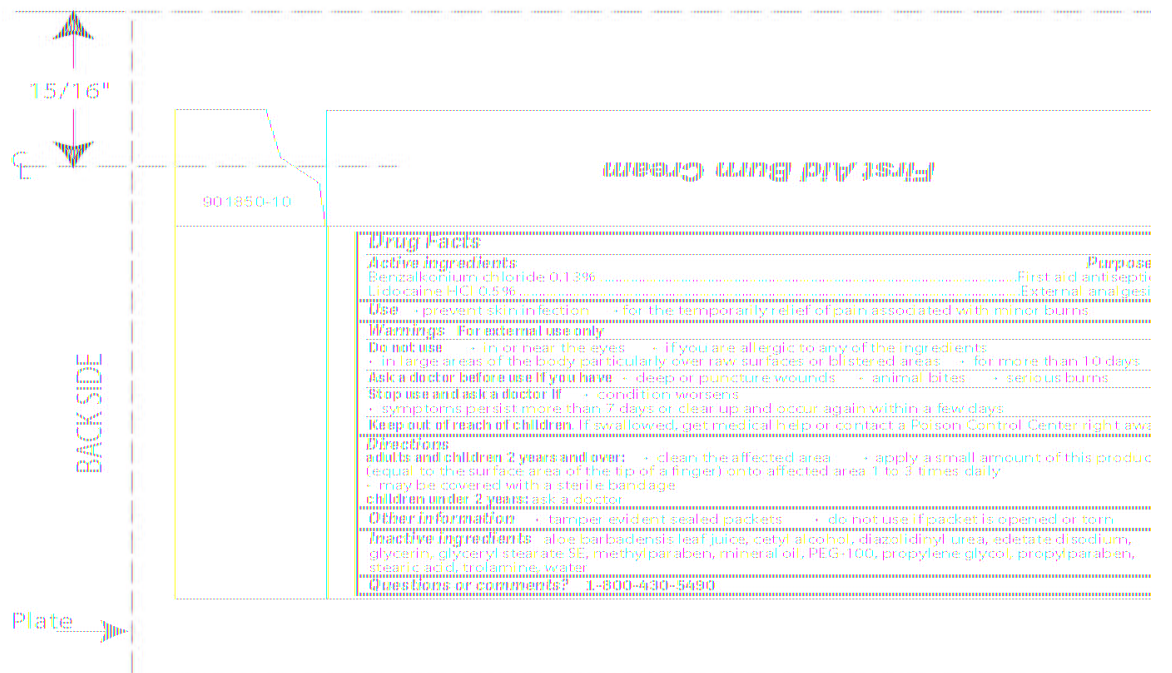
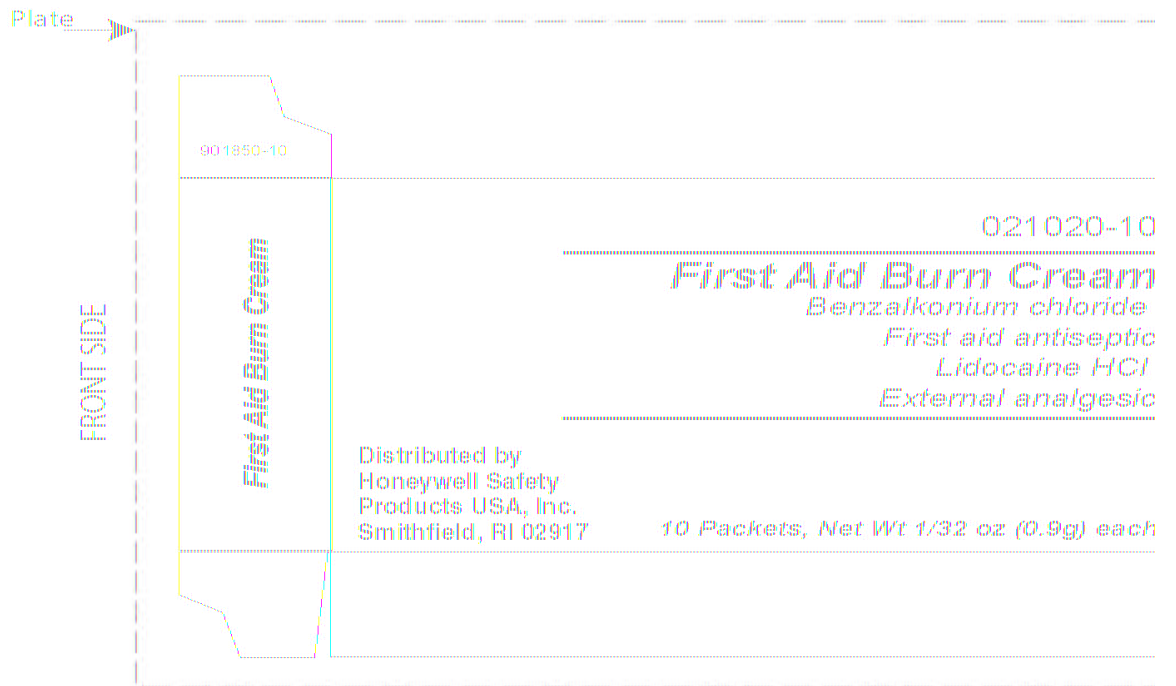
**4275**

### **SF00001582 kit contents**

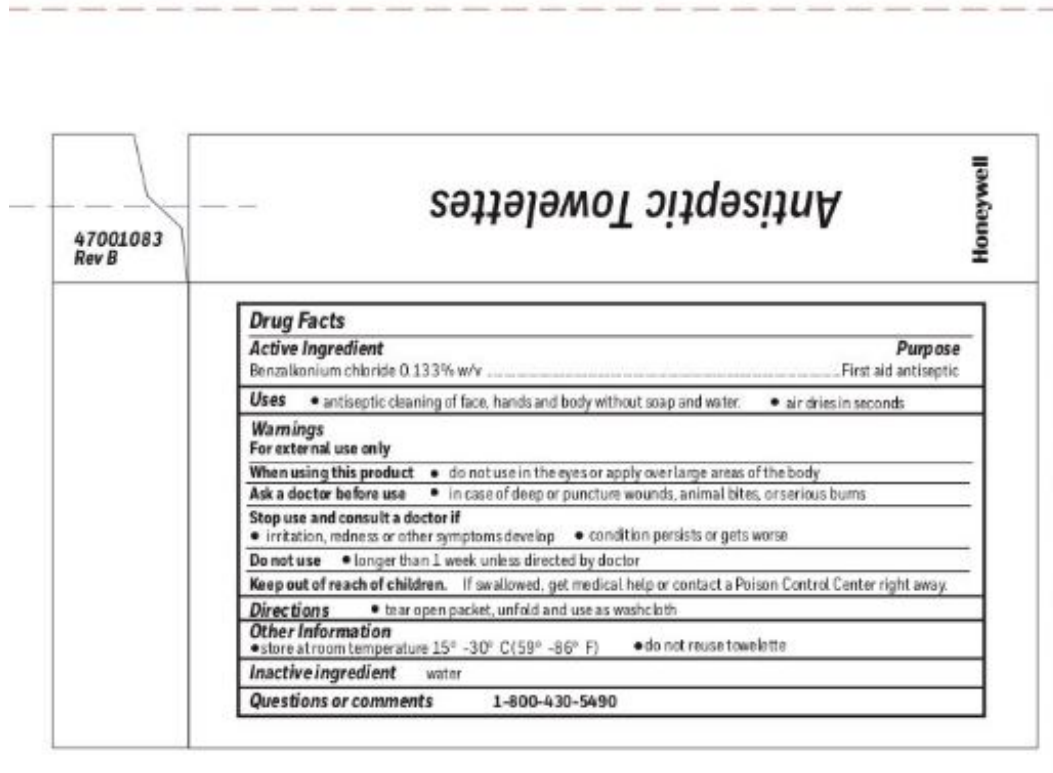
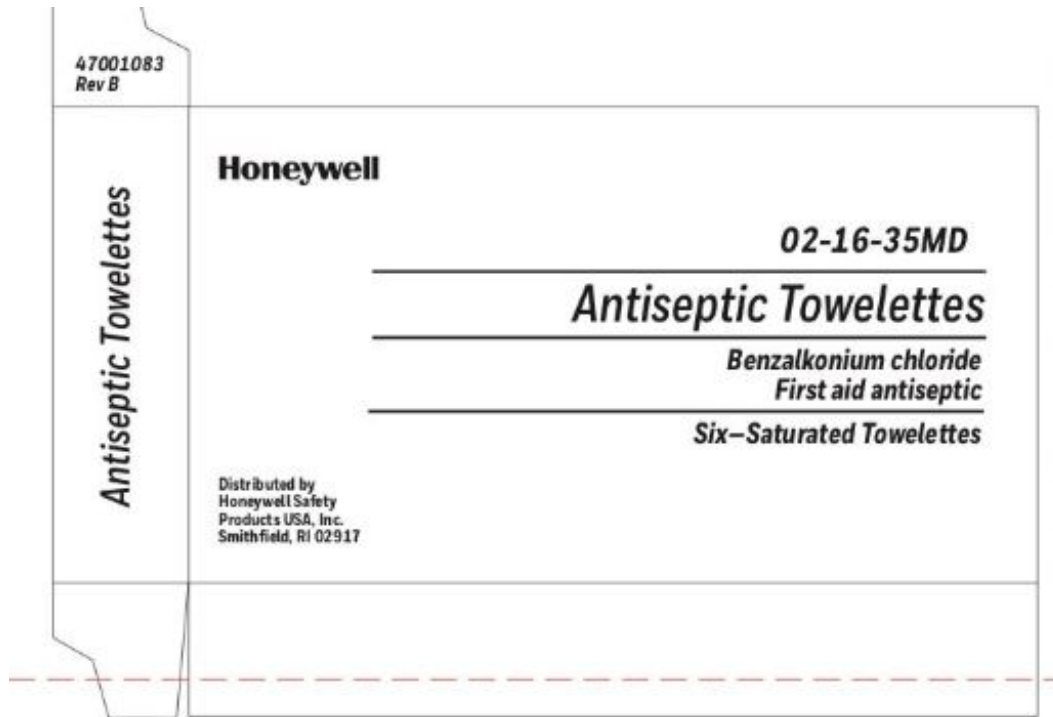
1 ADH TAPE W/P 1/2" X 2 1/2 YD  
1 GAUZE CLEAN-WRAP BDGE N/S 2"  
1 SCISSOR BDGE 4" RED PLS HDL  
1 KIT TWEEZER 3 1/2" SLANTED  
1 600H EMPTY BOX BLANK  
1 F. A. INST CHART SM (INDIVIDUAL LBL)  
LBL STOCK 6-3/8" X 4"  
1 LBL STOCK 3" x 1-7/8"  
4 BZK ANTISEPTIC WIPE, BULK  
4 FIRST AID CREAM 1.0GR PKT EACH  
4 HYDROCORTISONE 1% 0.9GM  
4 GAUZE PADS 2" X 2" 12PLY  
2 EYE PADS STD OVAL STERILE  
4 NON ADHERENT PAD 2" X 3"

## **First Aid Burn Cream**

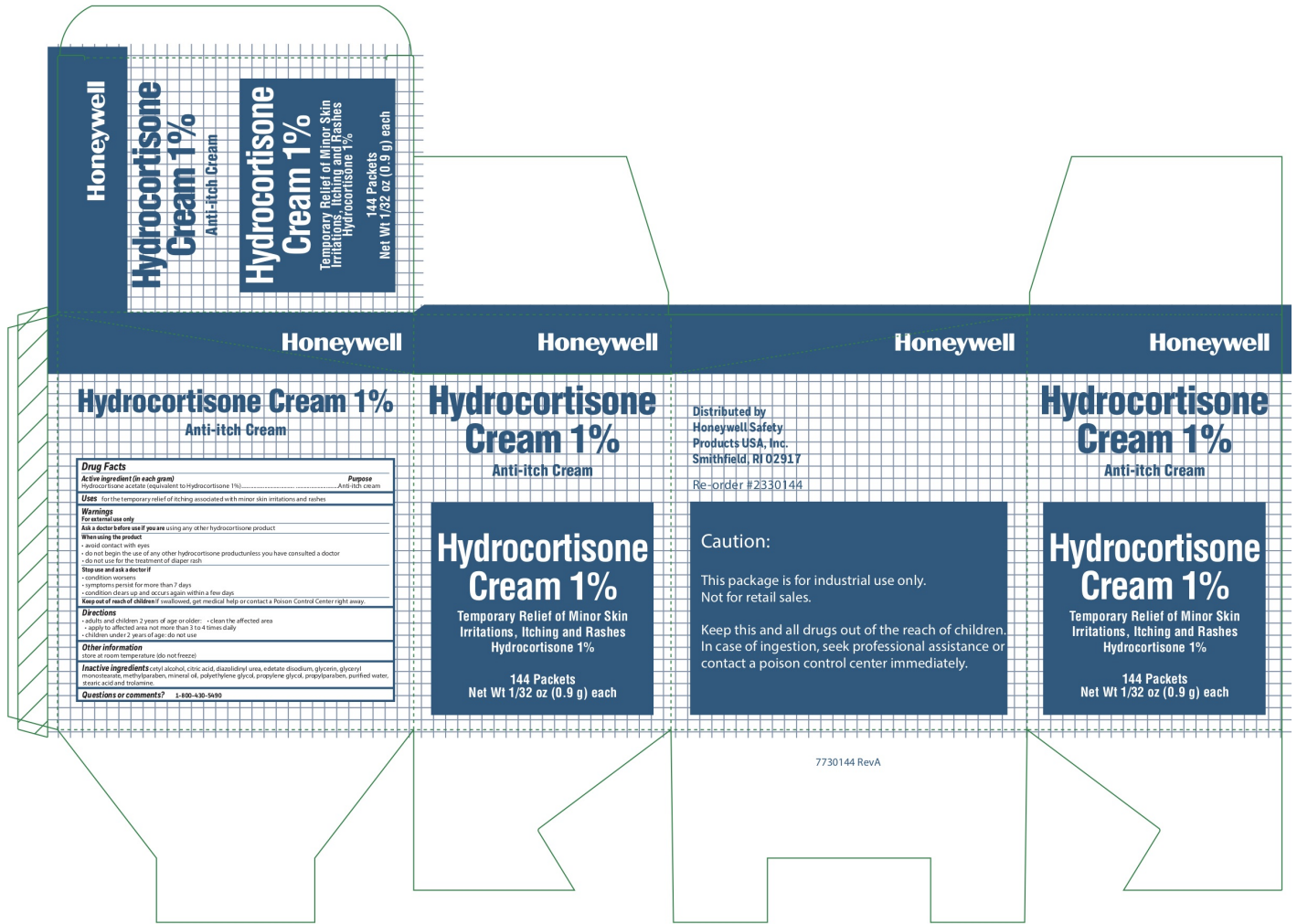
### **Principal Display Panel**



**Principal Display Panel**



**Hydrocortisone  
Principal Display Panel**



4223 Kit Label  
 34575H

# ***First Aid*** ***Handy Kit Jr.***

## *Contents List*

<b>10</b>	<b>1" x 3" Woven Bandages</b>	<b>1</b>	<b>2" x 5 Yd Gauze</b>
<b>2</b>	<b>2" x 3" Woven Bandages</b>	<b>4</b>	<b>2" x 3" Non-Stick Pads</b>
<b>2</b>	<b>Knuckle Bandages</b>	<b>4</b>	<b>Hydrocortisone Foil Packs</b>
<b>4</b>	<b>2" x 2" Gauze Pads</b>	<b>1</b>	<b>1/2" x 2 1/2 Yd Tape</b>
<b>4</b>	<b>First Aid Cream Foil Packs</b>	<b>1</b>	<b>Scissors &amp; Tweezers</b>
<b>4</b>	<b>Antiseptic Wipes</b>		

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

**4275 Kit Label**  
**SF00001582**





# FIRST AID KIT

*Red Rooster*® – Professional Tools  
Toll Free (866) 725-3627  
[www.redroosterag.com](http://www.redroosterag.com)

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

## 4223 FIRST AID KIT

4223 first aid kit kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-4223
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4223-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 PACKET	3.6 g
Part 2	4 PACKET	3.6 g
Part 3	4 PACKET	5.6 mL

**Part 1 of 4****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:W14X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

**Inactive Ingredients**

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

**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 4

### 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: 059QF0K00R)	
<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	NDC:0498-0903-34	10 in 1 BOTTLE, UNIT-DOSE		
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
OTC monograph not final	part333A	12/20/2017	

## Part 3 of 4

### ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

#### Product Information

Item Code (Source) NDC:0498-0501

Route of Administration TOPICAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL 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	part333E	12/21/2017	

## Part 4 of 4

### 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:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

### Inactive Ingredients

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

### 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

### Marketing Information

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

## 4275 FIRST AID KIT

4275 first aid kit kit

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4275

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4275-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 PACKET	3.6 g
Part 2	4 PACKET	3.6 g
Part 3	4 PACKET	5.6 mL
Part 4	4 PACKET	3.6 g

### Part 1 of 4

## HYDROCORTISONE

anti-itch cream

### Product Information

Item Code (Source)	NDC:0498-0801
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 in 100 g

### Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
TROLAMINE (UNII: 9O3K93S3TK)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	

### 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 4

### 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: 059QF0K00R)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>METHYL PARABEN</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>PROPYL PARABEN</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
OTC monograph not final	part333A	12/20/2017	

### Part 3 of 4

#### ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

### Product Information

Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL 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	part333E	12/21/2017	

### Part 4 of 4

#### 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 in 100 g

**Inactive Ingredients**

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

**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

**Marketing Information**

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

## Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4223, 0498-4275)

## Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0903, 0498-0800, 0498-0801)

## Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0501)

Revised: 10/2019

Honeywell Safety Products USA, INC