

4323 FIRST AID KIT- 4323 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.

0498-4323: First Aid Kit (Eye Wash, Hand Sanitizer, BZK wipe, FABC, Neomycin, Sting rel- SF00004664)

Active ingredient

Sterile Water 99%

Purpose

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 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.

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

Inactive Ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Questions?

Call **1-800-430-5490**

Honeywell Safety Products USA, Inc. Smithfield, RI 02917

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

Other information

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

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

Questions

1-800-430-5490

BZK Antiseptic Wipe***Active ingredient***

Benzalkonium chloride 0.13%

BZK***Purpose***

First aid antiseptic

BZK***Uses***

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

BZK

Warnings

For external use only

BZK

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

Other information

- store at room temperature 15 ° to 30 ° C (59 ° - 86 °F)
- do not reuse towelette

BZK

Inactive ingredients

water

BZK

Questions

1-800-430-5490

Sting Relief

Active ingredient

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

Sting relief

Questions or Comments

1-800-430-5490

Neomycin Antibiotic Ointment

Active ingredient

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

Neomycin Antibiotic Ointment

Purpose

First aid antibiotic

Neomycin Antibiotic Ointment

Uses

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

Neomycin Antibiotic Ointment

Warnings

For external use only

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

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 reach of children

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

Neomycin Antibiotic Ointment

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

Neomycin Antibiotic Ointment

Other information

store at 15 ° to 25 °C (59 ° to 77 °F)

Neomycin Antibiotic Ointment

Inactive ingredient

petrolatum

Neomycin Antibiotic Ointment

Questions

1-800-430-5490

Hand Sanitizer

Active ingredient

Ethyl alcohol 62%

Hand Sanitizer

Purpose

Antiseptic handwash

Hand Sanitizer

Uses

- for hand washing to decrease bacteria on skin
- recommended for repeated use

Hand Sanitizer

Warnings

For external use only

Flammable, keep away from fire or flame

When using this product

- do not use in the eyes
- discontinue use if irritation and redness develops
- If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children

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

Hand Sanitizer

Directions

- wet hands thoroughly with product and allow to dry without wiping

Hand Sanitizer

Other information

- place a quarter size amount into one hand, spread over both hands to wrist and rub into skin until dry
- store at 15 ° to 25 ° C (59 ° to 77 ° F)

Hand Sanitizer

Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, dl-alpha tocopheryl acetate, fragrance, PEG-60 almond glycerides, propylene glycol, purified water .

Hand Sanitizer

Questions or Comments

1-800-275-3433 info@waterjel.com

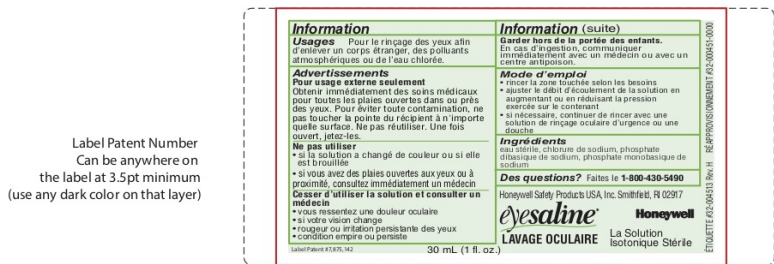
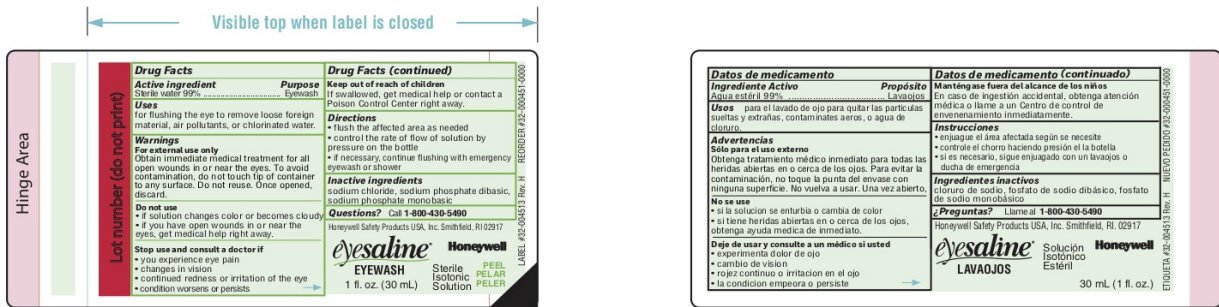
4323

SF00004664 Kit Contents

1 1X3 PLASTIC 100/BOX
1 EYE DRESS PKT W/4 ADH STRIPS
1 TWEEZER PLASTICS 4"
1 FIRST AID GUIDE ASHI
10 HAND SANITIZER 0.9G WJ BULK
3 GAUZE CLEAN-WRAP BDGE N/S 2"
2 GAUZE CLEAN-WRAP BDGE N/S 3"
2 ABD COMBINE PAD 5" X 9"
1 GZE PADS STERILE 3"X 3" 10'S
1 CO-FLEX BANDAGE 2"X 5YDS TAN
1 CPR FILTERSHIELD 77-100
1 1 OZ, BUFF EYEWASH
1 SCISSOR BDGE 4" RED PLS HDL
LBL STOCK 6-3/8"X4"
LBL STOCK 4"X2-7/8"
1 LBL STOCK 3"x1-7/8"
10 BZK ANTISEPTIC WIPE, BULK

4 PR LRG NITRILE GLVES ZIP BAG
10 FIRST AID BURN CREAM 1.0GR PKT EACH
2 TAPE ADHESIVE 1/2 X 2.5 125133
10 POUCH NEOMYCIN ANTIBIOTIC .9 G
1 WATER-JEL BURN DRESSING 4 X 4
2 ADH BNDG PLASTIC EX-LG 4"X 2"
1 KIT PP 24 UNIT FA
1 LBL CONTENTS ANSI 2015 CL A
1 BAG ZIPPER POLY 6 X 6 2 MIL
6 SAFETEC STING RELIEF WIPES BULK
1 TRI BNDG NON WOVEN 40"X40"X56"
1 COLD PACK UNIT 4"X6" BULK
4 WOVEN FINGERTIP BANDAGE 2"
3 WOVEN KNUCKLE BANDAGE

Eye Wash Package label



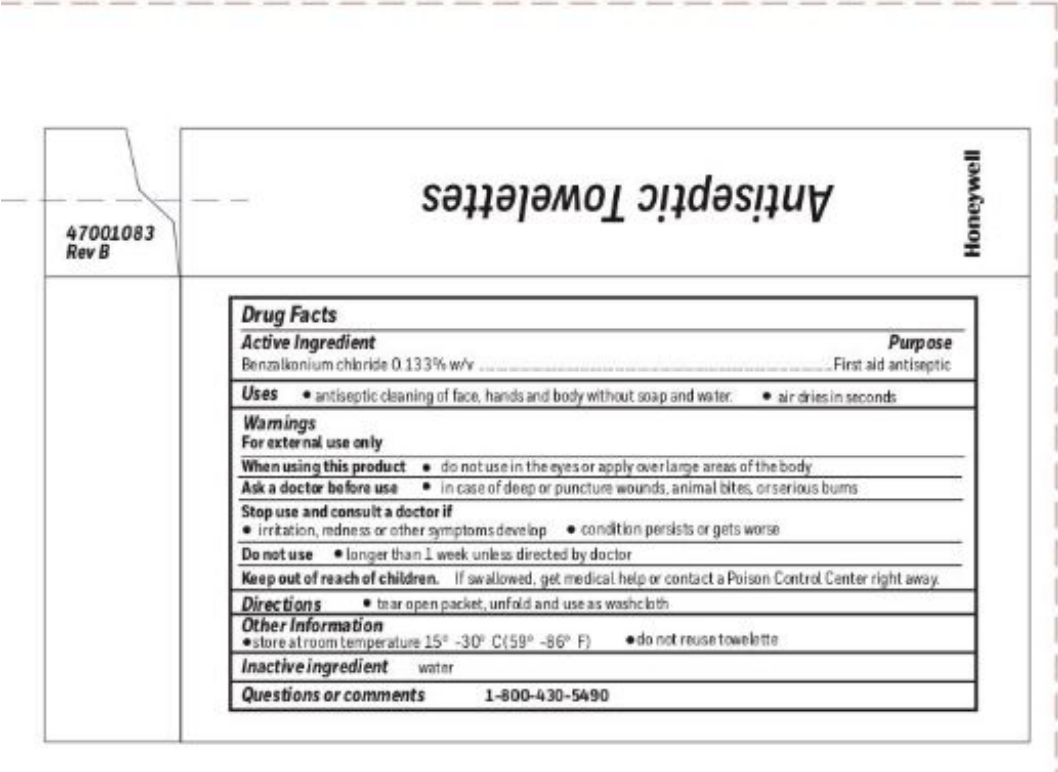
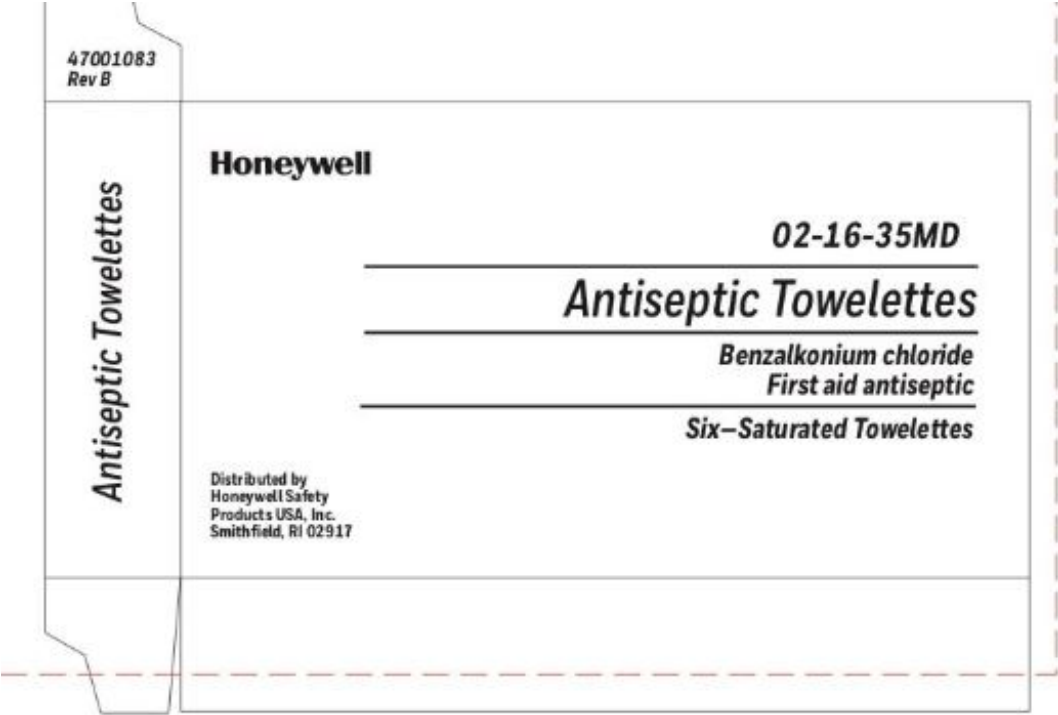
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First Aid Burn Cream Principal Display Panel

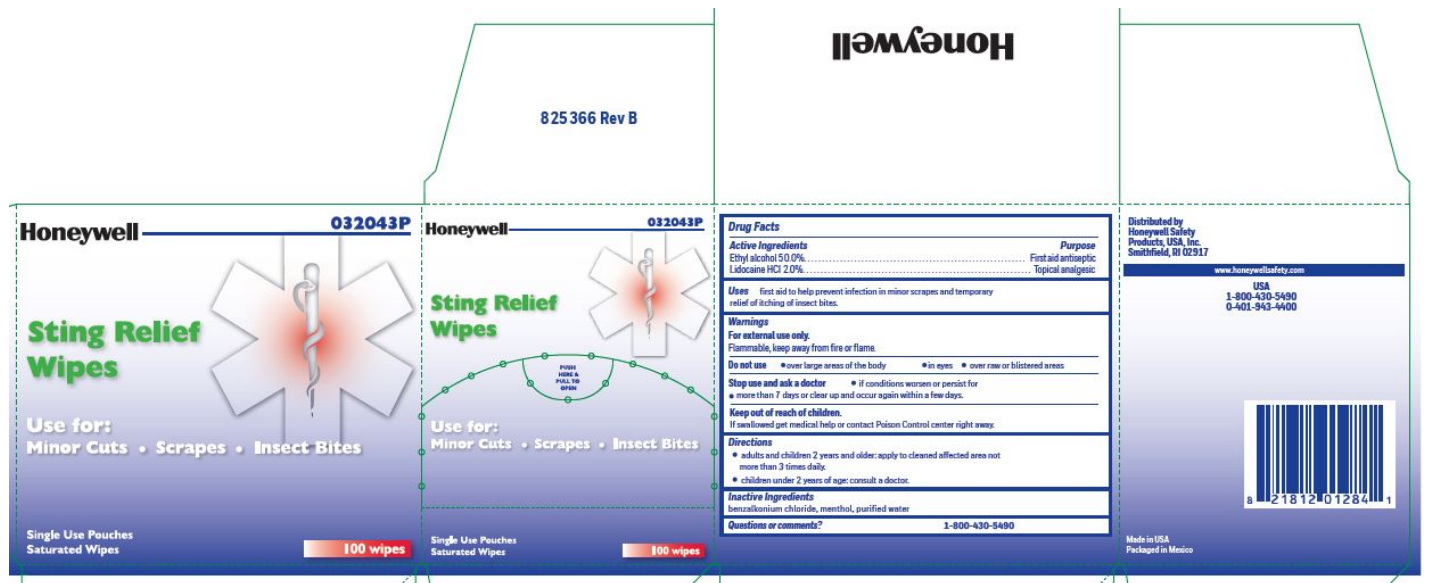
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Principal Display Panel



Sting Relief
Principal Display Panel



Neomycin Antibiotic Ointment Principal Display Panel

822568-25

**Neomycin First
aid antibiotic**

020126-25

Neomycin
First aid antibiotic
Neomycin sulfate
First aid antibiotic

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

25 Packets, Net Wt 1/32 oz (0.9 g) each

Neomycin First aid antibiotic

822568-25

Drug Facts

Active ingredient (in each gram) **Purpose**
Neomycin sulfate (equivalent to 3.5 mg neomycin) First aid antibiotic

Use 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 • over large areas of the body

Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns

Stop use and ask a doctor if • conditions persists or gets worse • rash or other allergic reaction develops
• you need to use longer than one week

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

Directions • clean the affected area

• apply a small amount of product (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 store at 15° to 25°C (59° to 77°F)

Inactive ingredient petrolatum

Questions or comments? 1-800-430-5490

Hand Sanitizer
Principal Display Panel



INSTANT

Hand Sanitizer

**Antiseptic Gel
With Vitamin E & Aloe**

Kills 99.9% of Germs

Without Water

240mL - (8 fl oz)

4323 Kit Label
SF00004664

 **First Aid** 

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

4323 FIRST AID KIT

4323 first aid kit kit

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4323
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4323-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	
Quantity of Parts				
Part #	Package Quantity		Total Product Quantity	
Part 1	1 BOTTLE		30 mL	
Part 2	6 POUCH		2.4 mL	
Part 3	10 PACKET		9 g	
Part 4	10 PACKET		9 g	
Part 5	10 PACKET		14 mL	
Part 6	1 PACKET		9 mL	
Part 1 of 6				
EYESALINE EMERGENCY EYEWASH				
purified water liquid				
Product Information				
Item Code (Source)		NDC:0498-0100		
Route of Administration		OPHTHALMIC		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)			WATER	98.6 mL in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)				
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:0498-0100-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
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Marketing Information

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

Part 2 of 6

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
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL 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		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 3 of 6

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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z 83H0X)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z 65AP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z 8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ 3MPW4)	

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 4 of 6

NEOMYCIN
antibiotic ointment

Product Information	
Item Code (Source)	NDC:0498-0730
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0730-01	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		03/31/2010	

Part 5 of 6

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		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
unapproved drug other		12/21/2017	

Part 6 of 6

INSTANT HAND SANITIZER

alcohol liquid

Product Information	
Item Code (Source)	NDC:59898-420
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-420-36	9 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
unapproved drug other		04/15/2011	

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 (118768815)

Revised: 1/2024

Honeywell Safety Products USA, INC