

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.

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

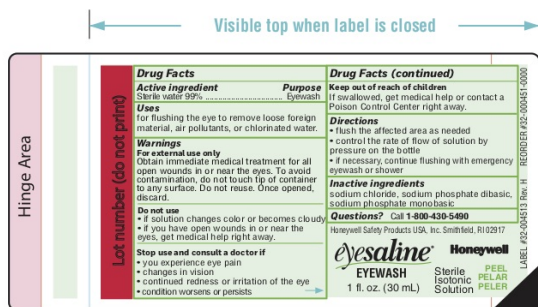
SF00004664 Kit Contents

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

#32-004513 Rev. H

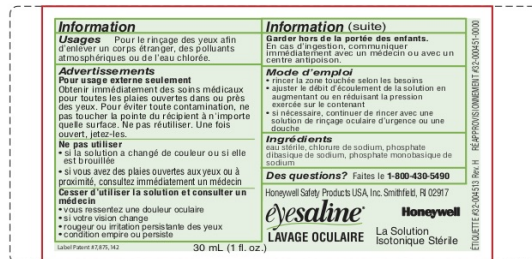


Top Panel
3/32" from die edges



Back
1/8" from die edges

Label Patent Number
Can be anywhere on
the label at 3.5pt minimum
(use any dark color on that layer)



Base
3/32" from die edges

Printable Text Area

**First Aid Burn Cream
Principal Display Panel**

796353-10 Rev. A Unit Carton Printing Plate for "B" size

Plate →

FRONT SIDE

901850-10

021020-10

First Aid Burn Cream
Benzalkonium chloride
First aid antiseptic
Lidocaine HCl
External analgesic

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

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

15/16"

f

BACK SIDE

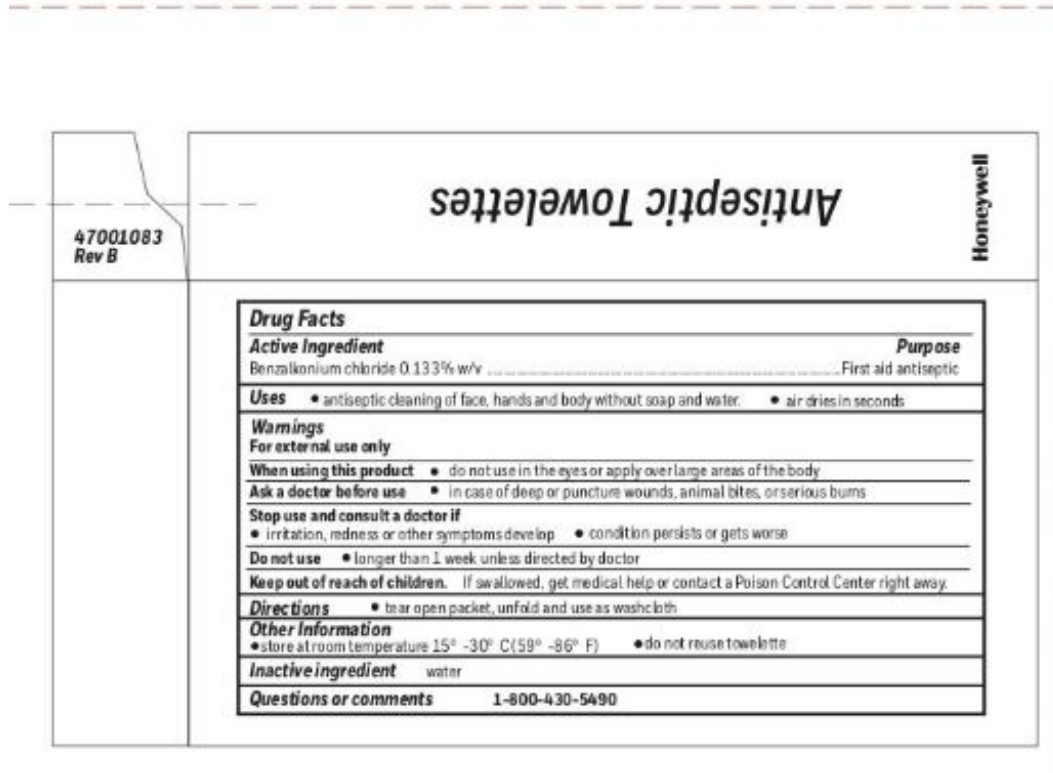
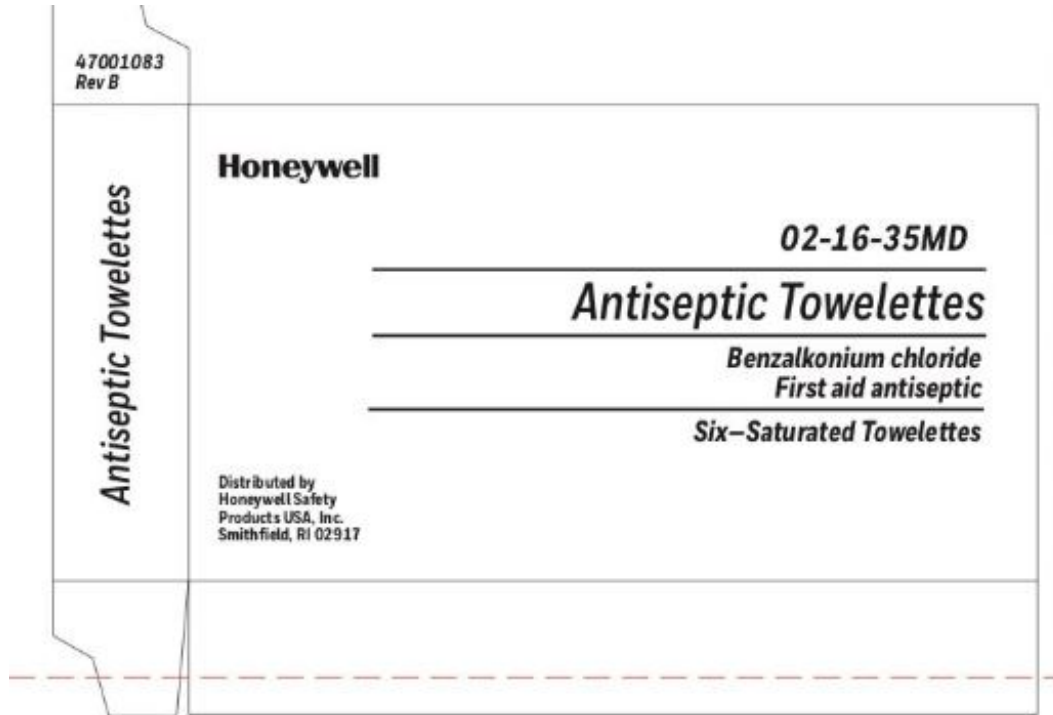
901850-10

First Aid Burn Cream

Drug Facts	
Active ingredients	Purpose
Benzalkonium chloride 0.13%	First aid antiseptic
Lidocaine HCl 0.5%	External analgesic
Use - prevent skin infection - for the temporary relief of pain associated with minor burns	
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 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	
Directions	
adults and children 2 years and over: - 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: ask a doctor	
Other information - tamper evident sealed packets - do not use if packet is opened or torn	
Inactive ingredients aloe barbadensis leaf juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, triethylamine, water	
Questions or comments? 1-800-430-5490	

Plate →

**BZK
Principal Display Panel**



**Sting Relief
Principal Display Panel**

Honeywell

825366 Rev B

Honeywell

032043P

Honeywell

032043P

Sting Relief Wipes



Use for:
Minor Cuts • Scrapes • Insect Bites

Single Use Pouches
Saturated Wipes

100 wipes

Sting Relief Wipes



Use for:
Minor Cuts • Scrapes • Insect Bites

Single Use Pouches
Saturated Wipes

100 wipes

POUR
HERE &
PULL TO
OPEN

Drug Facts

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

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

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

www.honeywellsafety.com

USA
1-800-430-5490
0-401-343-4400



Made in USA
Packaged in Mexico

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**



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
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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	10 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: 059QF0K00R) (WATER - UNII:059QF0K00R)	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		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	12/18/2018	

Part 2 of 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
OTC monograph not final	part348	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: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
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: 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
OTC monograph not final	part333A	12/20/2017	

Part 4 of 6

NEOMYCIN

antibiotic ointment

Product Information

Item Code (Source)	NDC:0498-0730
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Route of Administration	TOPICAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	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
OTC monograph final	part333B	03/31/2010	

Part 5 of 6**ANTISEPTIC TOWELETTE**

benzalkonium chloride liquid

Product Information

Item Code (Source)	NDC:0498-0501
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Route of Administration	TOPICAL
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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
OTC monograph not final	part333E	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: 059QF0K00R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-420-36	0.9 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
OTC monograph not final	part333E	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 (079287321)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4323)

Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0903, 0498-0730, 59898-420)

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(0498-0501)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America Inc		874965262	manufacture(0498-0733)