4109 FIRST AID KIT- 4109 first aid kit
4111 FIRST AID KIT- 4111 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.

0498-4109 & 4111: First Aid Kit (Bagged Components- 019701-0001L, 019743-0030L, 019759-0035L, 019760-0036L, SF00001231, Z019842, Z019760-0036L)

First Aid Burn Cream
Active ingredient

Benzalkonium chloride o.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 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 º to 30 º C (59 º - 86 ºF)
- do not reuse towelette

BZK
Inactive ingredients

water

BZK
Questions

1-800-430-5490

Aypanal
Active igredient

Acetaminophen 325 mg

Aypanal
Purpose

Pain reliever/fever reducer

Aypanal
Uses

- temporarily relieves minor aches and pains due to the common cold and headache
- temporarily reduces fever

Ask a doctor before use if you have

liver disease

Aypanal
Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: more than 4,000 mg in 24 hours, which is the maximum daily amount - child takes

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

skin reddening
blisters
rash

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if

you are taking the blood thinning drug warfarin

if preganat or breast feeding

ask a health professional before use

Keep out of rech of children

Keep out of reach of children

Overdose Warning

Overdose warning: In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for

children even if you do not notice any signs or symptoms.

Aypanal
Directions

- do not take more than directed (see overdose warning) adults and children 12 years of age or older
- take two tablets every 4-6 hours while symptoms last
- do not take more than directed (see overdose warning)

adults and children 12 years of age or older

- take two tablets every 4-6 hours while symptoms last
- do not take more than 12 tablets in 24 hours
- children 6 to under 12 years of age
- take 1 tablet every 4-6 hours while symptoms last
- do not take more than 5 tablets in 24 hours
- children under 6 years
- consult a doctor

Aypanal
Other information

store at room temperature 15 º to 30 º C (59 º - 86 ºF) TAMPER EVIDENT PACKETS- DO NOT USE IF OPEN OR TORN

Aypanal
Inactive ingredients

corn starch, microcrystalline cellulose, povidone, sodium starch glycolate, stearic acid.

Aypanal
Questions or Comments?

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 º to 25 ºC (59 º to 77 ºF)

Neomycin Antibiotic Ointment
Inactive ingredient

petrolatum

Neomycin Antibiotic Ointment
Questions

1-800-430-5490

4107
019701-0001L Kit Contents

1 TWEEZER PLASTICS 4"
1 FIRST AID GUIDE ASHI
1 GAUZE CLEAN-WRAP BDGE N/S 2"
1 GAUZE CLEAN-WRAP BDGE N/S 3"
1 ABD COMBINE PAD 5" X 9"
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
1 1" X 3" PLASTIC BANDS 16/BAG
2 TAPE ADHESIVE 1/2 X 2.5 125133
1 ADH BNDG PLASTIC EX-LG 4"X 2"
1 KIT STL 10 UN WHITE 01
1 TRI BNDG NON WOVEN 40"X40"X56"
1 COLD PACK UNIT 4"X6" BULK
4 GAUZE PADS 2"X2" 12PLY
4 GAUZE PADS 3"X3" 12PLY
1 WOVEN FINGERTIP BANDAGE 2"

4108
019743-0030L Kit Contents

1 TWEEZER PLASTICS 4"
2 GAUZE CLEAN-WRAP BDGE N/S 3"
1 ABD COMBINE PAD 5" X 9"
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
2 1" X 3" PLASTIC BANDS 16/BAG
1 TAPE ADHESIVE 1/2 X 2.5 125133
1 KIT, PP 16 UNIT FA
1 TRI BNDG NON WOVEN 40"X40"X56"
1 COLD PACK UNIT 4"X6" BULK
3 EYE PADS STD OVAL STERILE
6 GAUZE PADS 3"X3" 12PLY

4109

019759-0035L Kit Contents

1 GAUZE PADS, 3" X 3", 4 PER
1 FLEXICON 2"X 4.1 YD 12/BAG
1 FIRST AID GUIDE ASHI
1 ABD COMBINE PAD 5" X 9"
2 BUGX TOWELETTE, BULK, 300/CS
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
1 1" X 3" PLASTIC BANDS 16/BAG
1 TAPE ADHESIVE 1/2 X 2.5 125133
1 KIT STL 10 UN WHITE 01

4110
019760-0036L Kit Contents

1 FNGRTIP-5 PER, KNCKL BDG-4 PER
1 GAUZE PADS, 3" X 3", 4 PER
1 FLEXICON 2"X 4.1 YD 12/BAG
1 FIRST AID GUIDE ASHI
1 EMERGENCY SURVIVAL BLANKET
1 ABD COMBINE PAD 5" X 9"
2 BUGX TOWELETTE, BULK, 300/CS
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
1 1" X 3" PLASTIC BANDS 16/BAG
1 TAPE ADHESIVE 1/2 X 2.5 125133
1 KIT STL 16 UN (HORIZONTAL)
1 TRI BNDG NON WOVEN 40"X40"X56"

4111
SF00001231 Kit Contents

1 TWEEZER PLASTICS 4"
1 FIRST AID GUIDE ASHI
1 GAUZE CLEAN-WRAP BDGE N/S 2"
1 GAUZE CLEAN-WRAP BDGE N/S 3"
1 ABD COMBINE PAD 5" X 9"
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
1 1" X 3" PLASTIC BANDS 16/BAG
2 TAPE ADHESIVE 1/2 X 2.5 125133
1 ADH BNDG PLASTIC EX-LG 4"X 2"
1 KIT, PP 10 UNIT FA
1 BAG ZIPPER POLY 6 X 6 2 MIL
1 TRI BNDG NON WOVEN 40"X40"X56"

1 COLD PACK UNIT 4"X6" BULK
4 GAUZE PADS 2"X2" 12PLY
1 GAUZE PADS 3"X3" 12PLY
1 WOVEN FINGERTIP BANDAGE 2"

4119
Z019842 Kit Contents

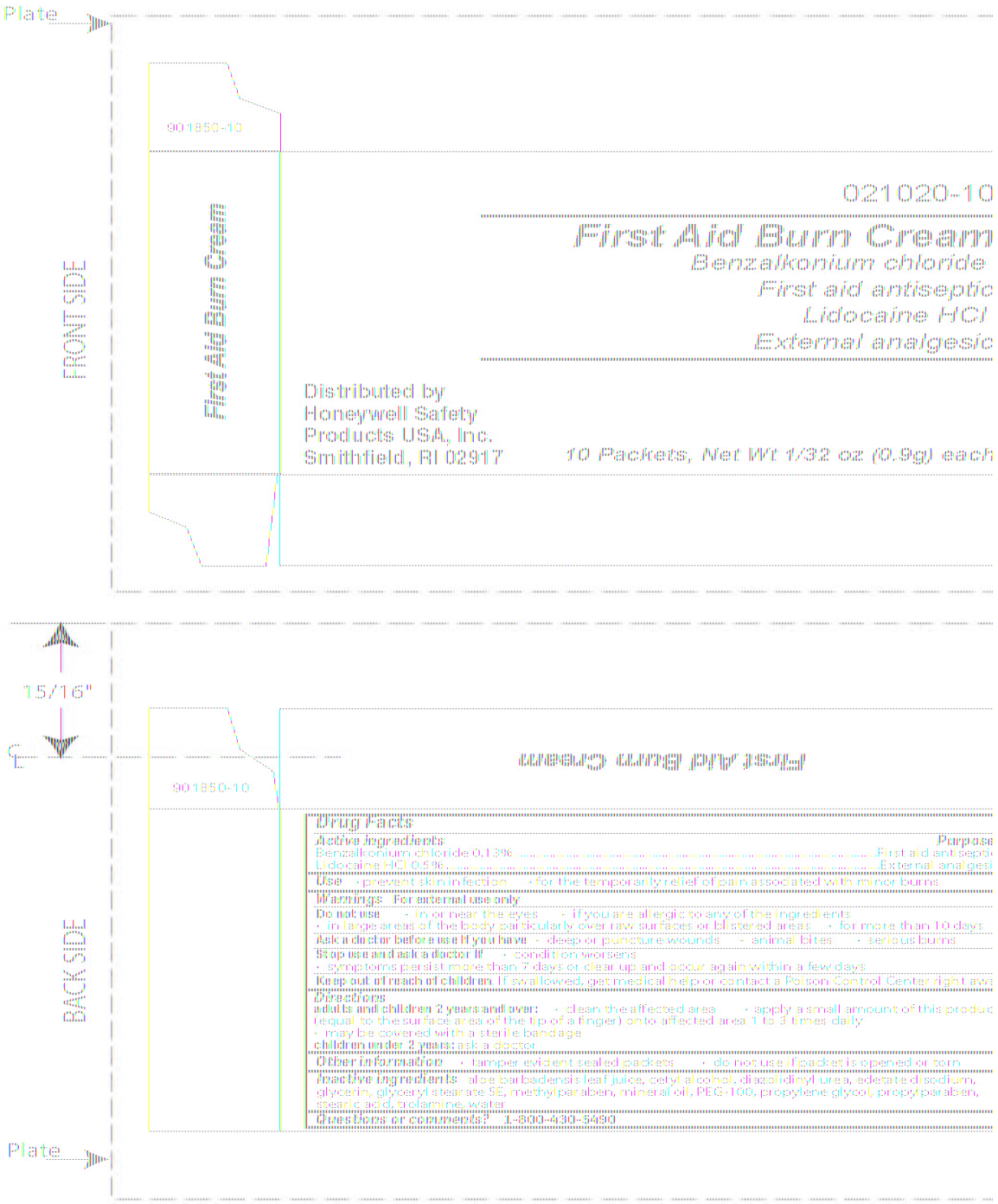
1 GAUZE PADS, 3" X 3", 4 PER
1 ADHESIVE BDG,PLSTIC,1"X3"16PER
1 FLEXICON 2"X 4.1 YD 12/BAG
1 FIRST AID GUIDE ASHI
1 ABD COMBINE PAD 5" X 9"
2 BUGX TOWELETTE, BULK, 300/CS
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 LABEL COVER, GRAINGER Z019842
1 TAPE ADHESIVE 1/2 X 2.5 125133
1 KIT STL 10 UN WHITE 01

4120
Z019760-0036L Kit Contents

1 FNGRTIP-5 PER, KNCKL BDG-4 PER
1 GAUZE PADS, 3" X 3", 4 PER
1 ADHESIVE BDG,PLSTIC,1"X3"16PER
1 FLEXICON 2"X 4.1 YD 12/BAG
1 FIRST AID GUIDE ASHI
1 EMERGENCY SURVIVAL BLANKET
1 ABD COMBINE PAD 5" X 9"
2 BUGX TOWELETTE, BULK, 300/CS
1 BAGGED COMP MISC
1 SCISSOR BDGE 4" RED PLS HDL
1 2 PR LRG NITRILE GLVES ZIP BAG
1 TAPE ADHESIVE 1/2 X 2.5 125133
1 KIT STL 16 UN (HORIZONTAL)
1 LABEL, VEHICLE KIT (6 Person)

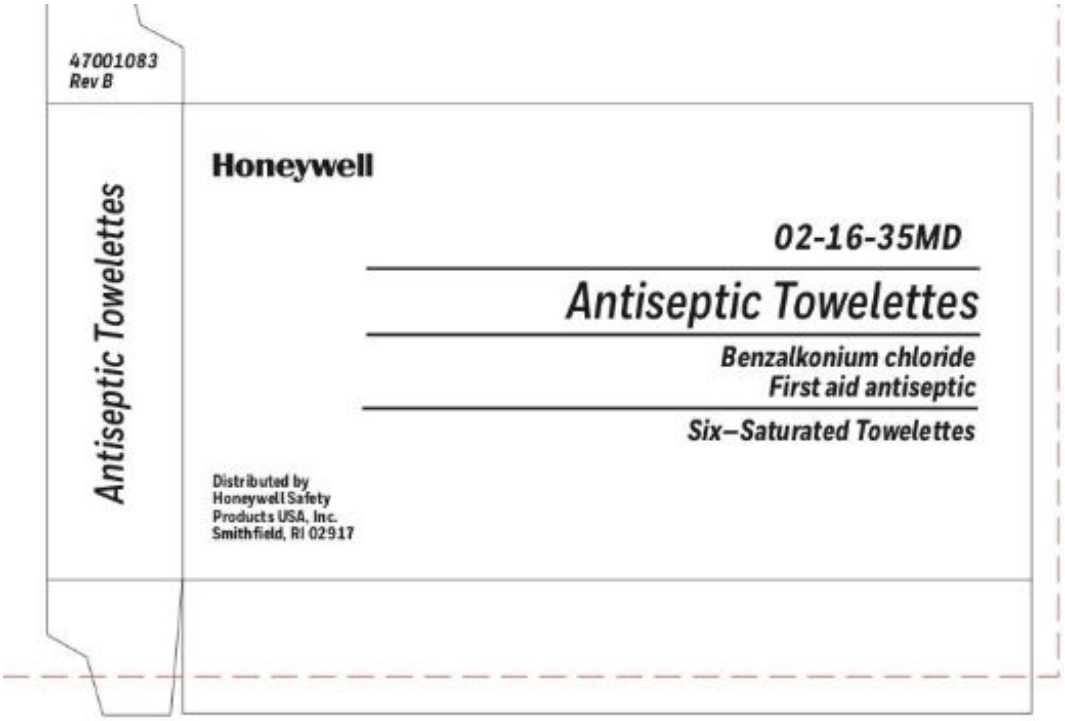
First Aid Burn Cream
Principal Display Panel

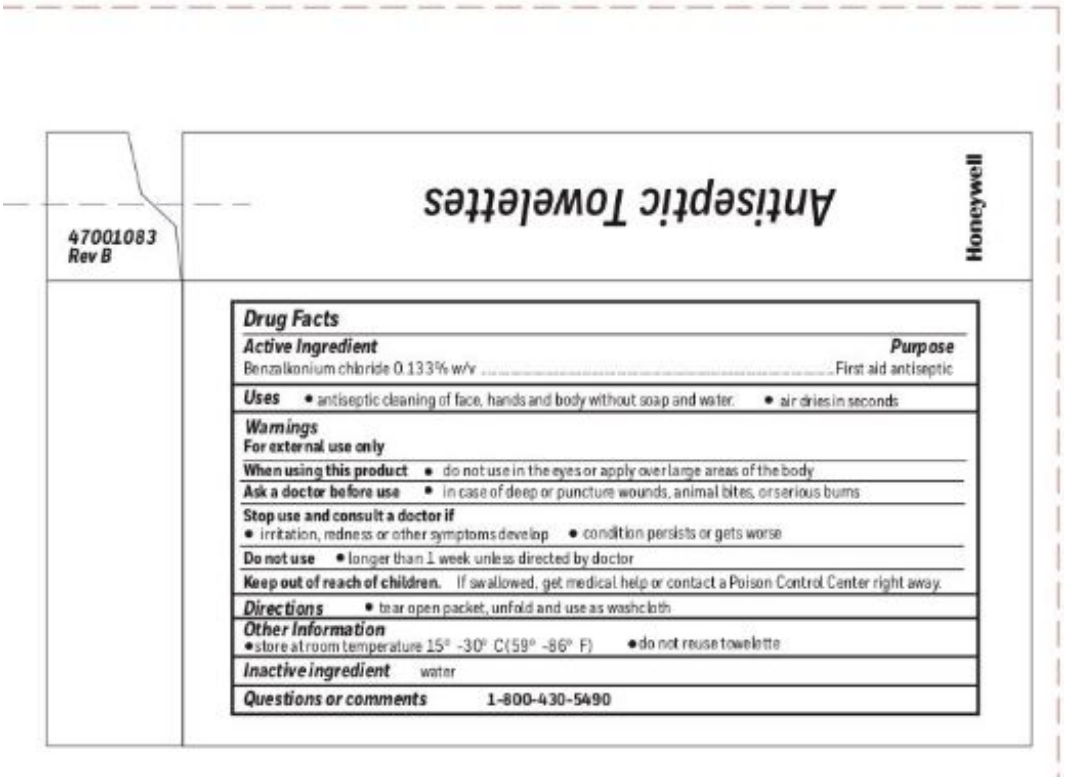
796353-10 Rev. A Unit Carton Printing Plate for "B" size



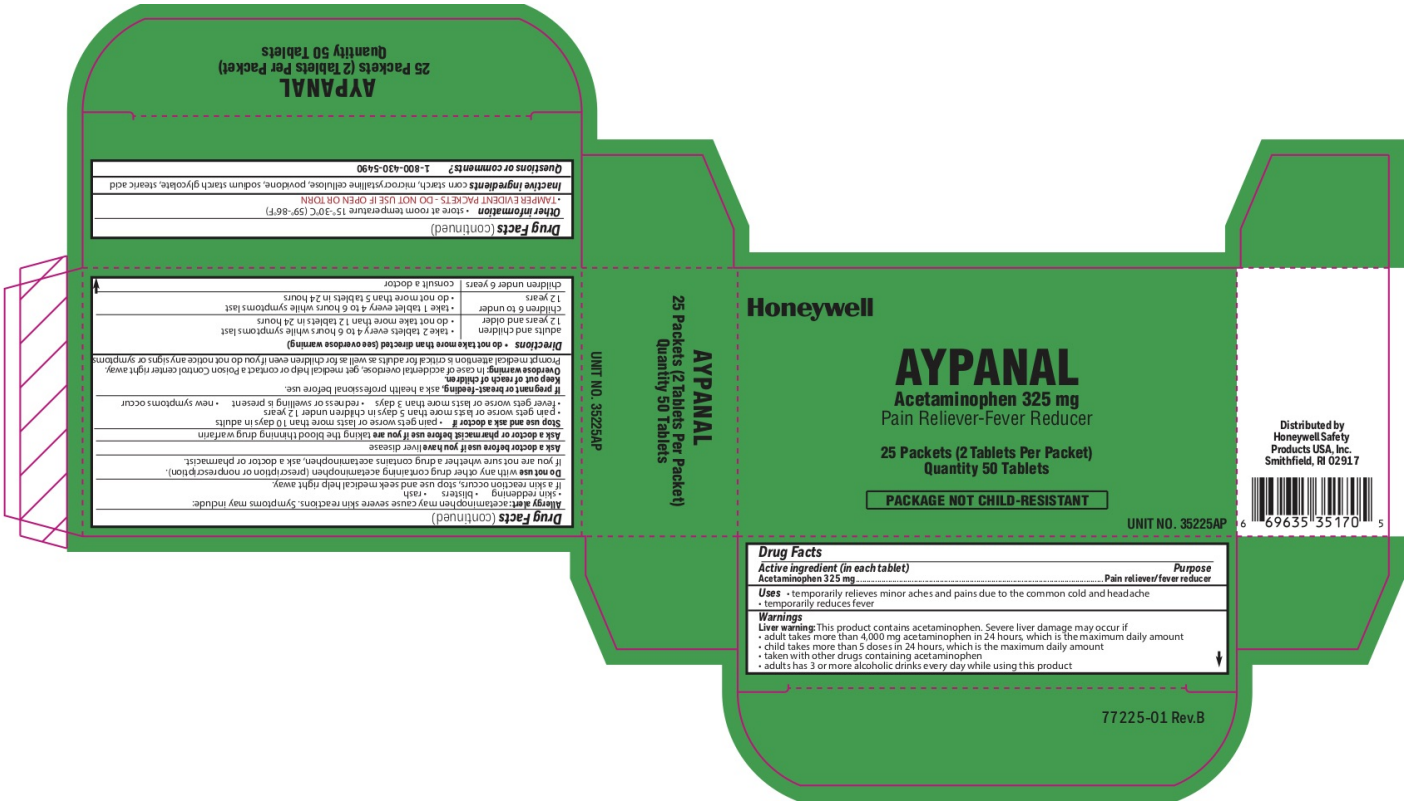
BZK

Principal Display Panel





Aypanal
Principal Display Panel



Sting Relief
Principal Display Panel

Neomycin Antibiotic Ointment
Principal Display Panel

822568-25

020126-25

Neomycin
First aid antibiotic
Neomycin sulfate
First aid antibiotic

Neomycin 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

4107 Kit Label
019701-0029L

777030 Rev C
white printed 5 colors
4 color process and pms 072 blue

4108 Kit Label
019743-0030L

Standard Kit
Printed 4 color process with PMS 072 blue

4109 Kit Label
019759-0035L

777046
4 color process, pms 072 blue, and pms 186 red

4110 Kit Lael
019760-0036L

printed 5 color process and PMS blue 072

4111 Kit Label
SF00001231

Rapid Label PICS 10u/10p
white printed 4 color process and red (pms 185c)

4119 Kit Label
Z019842

4120 Kit Label
Z019760-0036L

printed 5 color process and PMS blue 072

4109 FIRST AID KIT

4109 first aid kit kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4109

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-4109-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	3 PACKET	6
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 1 of 5

AYPANAL NON-ASPIRIN

acetaminophen tablet

Product Information	
Item Code (Source)	NDC:0498-2001
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics			
Color	white	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	circle;U
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-2001-01	2 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		04/10/2012	

Part 2 of 5

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information	
Item Code (Source)	NDC:0498-0733
Route of Administration	TOPICAL

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

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1		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 5

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

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

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
unapproved drug other		12/20/2017	

Part 4 of 5

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 5

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	

Marketing Information

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

4111 FIRST AID KIT

4111 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4111

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4111-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	3 PACKET	6
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 1 of 5

AYPANAL NON-ASPIRIN

acetaminophen tablet

Product Information	
Item Code (Source)	NDC:0498-2001
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics			
Color	white	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	circle;U
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-2001-01	2 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		04/10/2012	

other		04/10/2012	

Part 2 of 5

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information	
Item Code (Source)	NDC:0498-0733
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
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 5

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

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

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
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)	

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 5

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 5

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	

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)

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