4278 FIRST AID KIT- 4278 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.

0498-4278: First Aid Kit (Eye Wash, BCM SF00002177)

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 ey
- 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 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

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 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

4277
SF00002177 Kit Contents

1 GAUZE PADS, 3" X 3", 4 PER

1 TWEEZER PLASTICS 4"

1 FIRST AID GUIDE ASHI

2 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 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"

1 LBL NORTH CONTS 6.75X3.5 ID B

2 PR LRG NITRILE GLVES ZIP BAG

2 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 16 UNIT FA

1 LBL CONTENTS ANSI Z308.1-2009 REV B

1 TRI BNDG NON WOVEN 40"X40"X56"

1 COLD PACK UNIT 4"X6" BULK

4 GAUZE PADS 2"X2" 12PLY

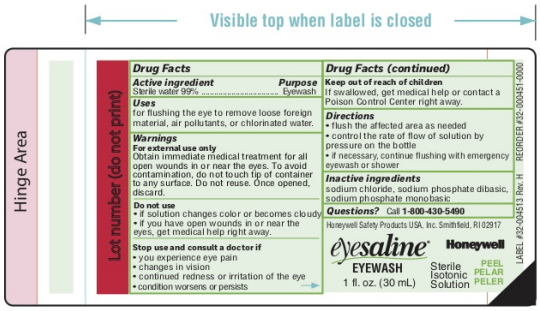
1 EYE PADS STD OVAL STERILE

3 WOVEN FINGERTIP BANDAGE 2"

2 WOVEN KNUCKLE BANDAGE

Eye Wash Package label

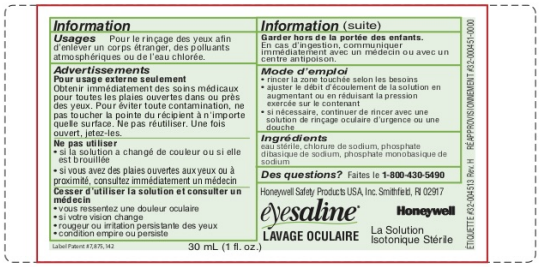
#32-004513 Rev. H



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)

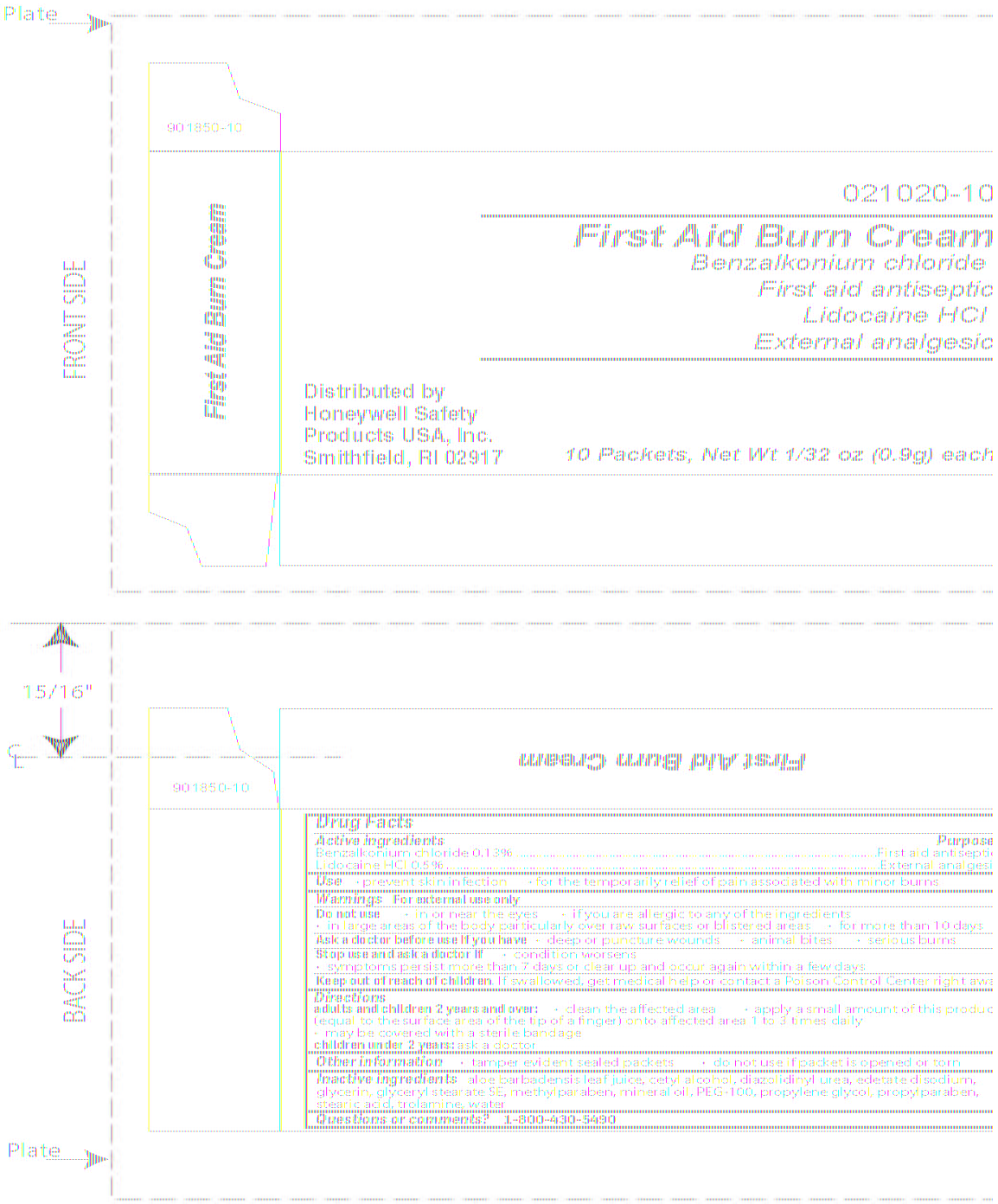


Base
3/32″ from die edges

Printable Text Area

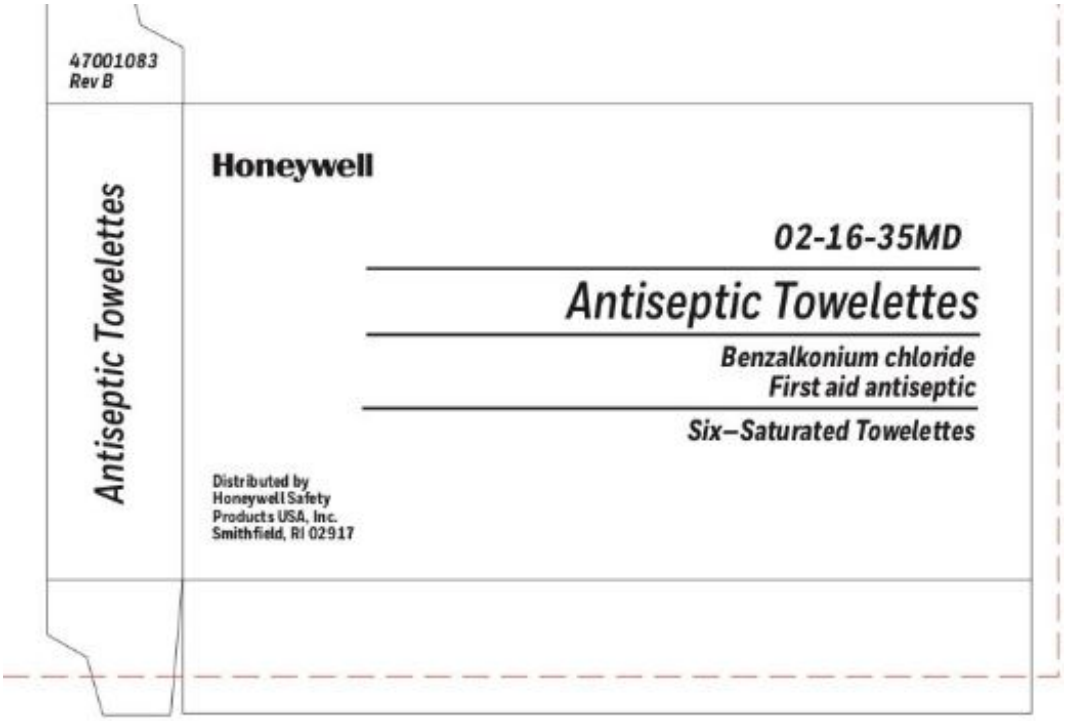
First Aid Burn Cream
Principal Display Panel

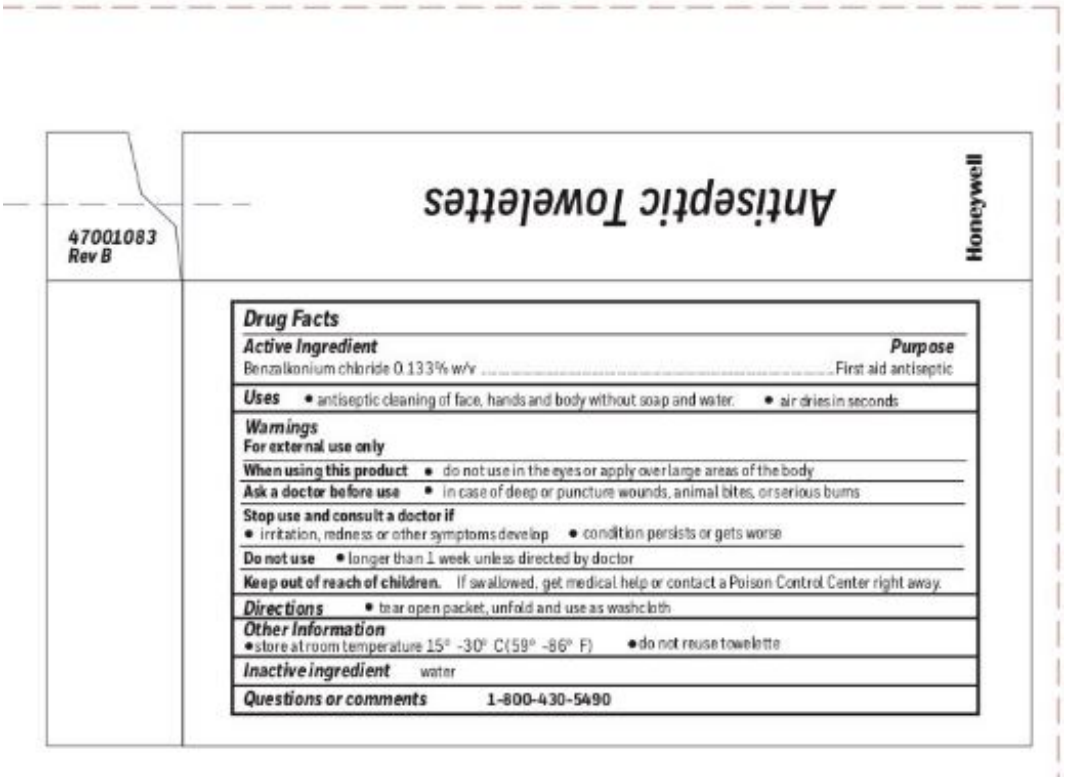
796353-10 Rev. A Unit Carton Printing Plate for "B" size



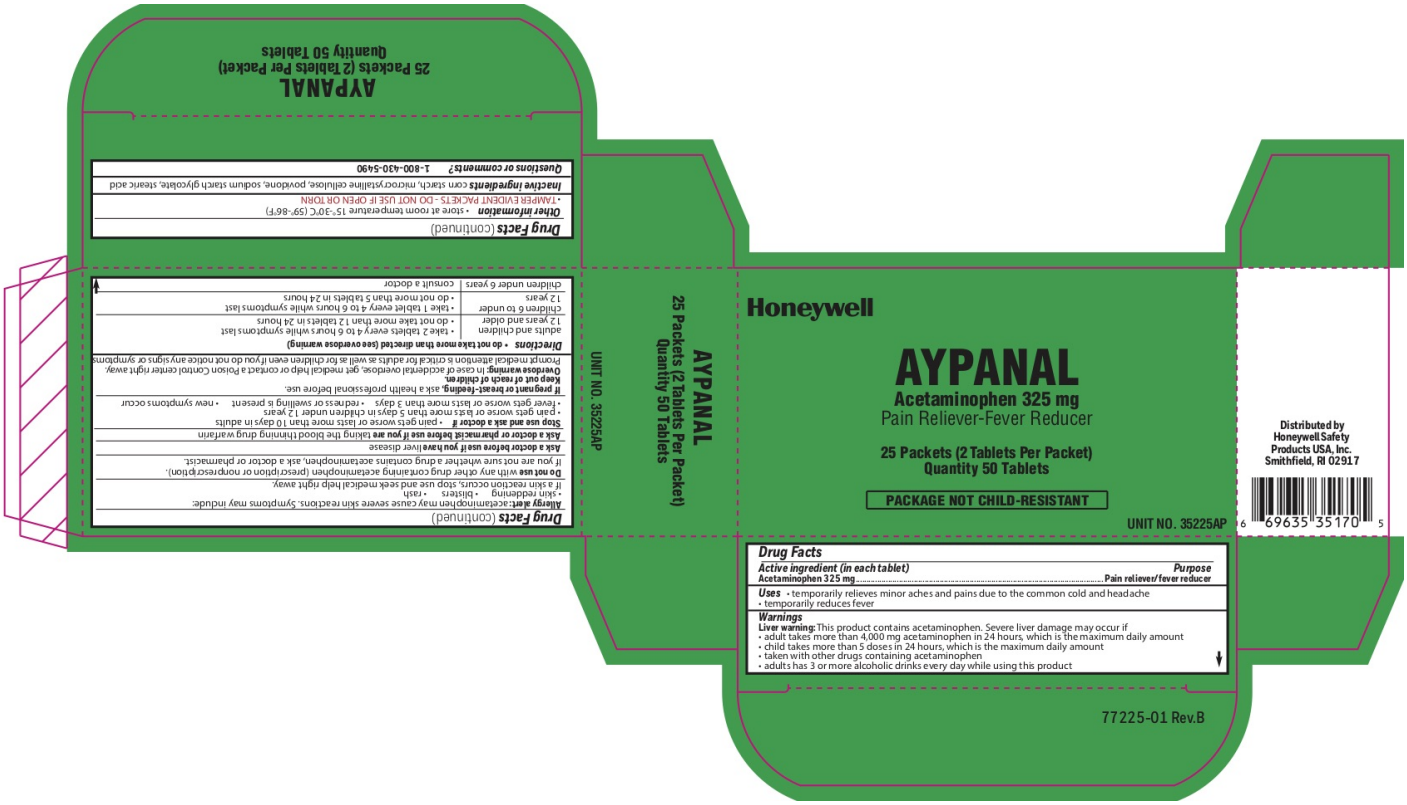
BZK

Principal Display Panel



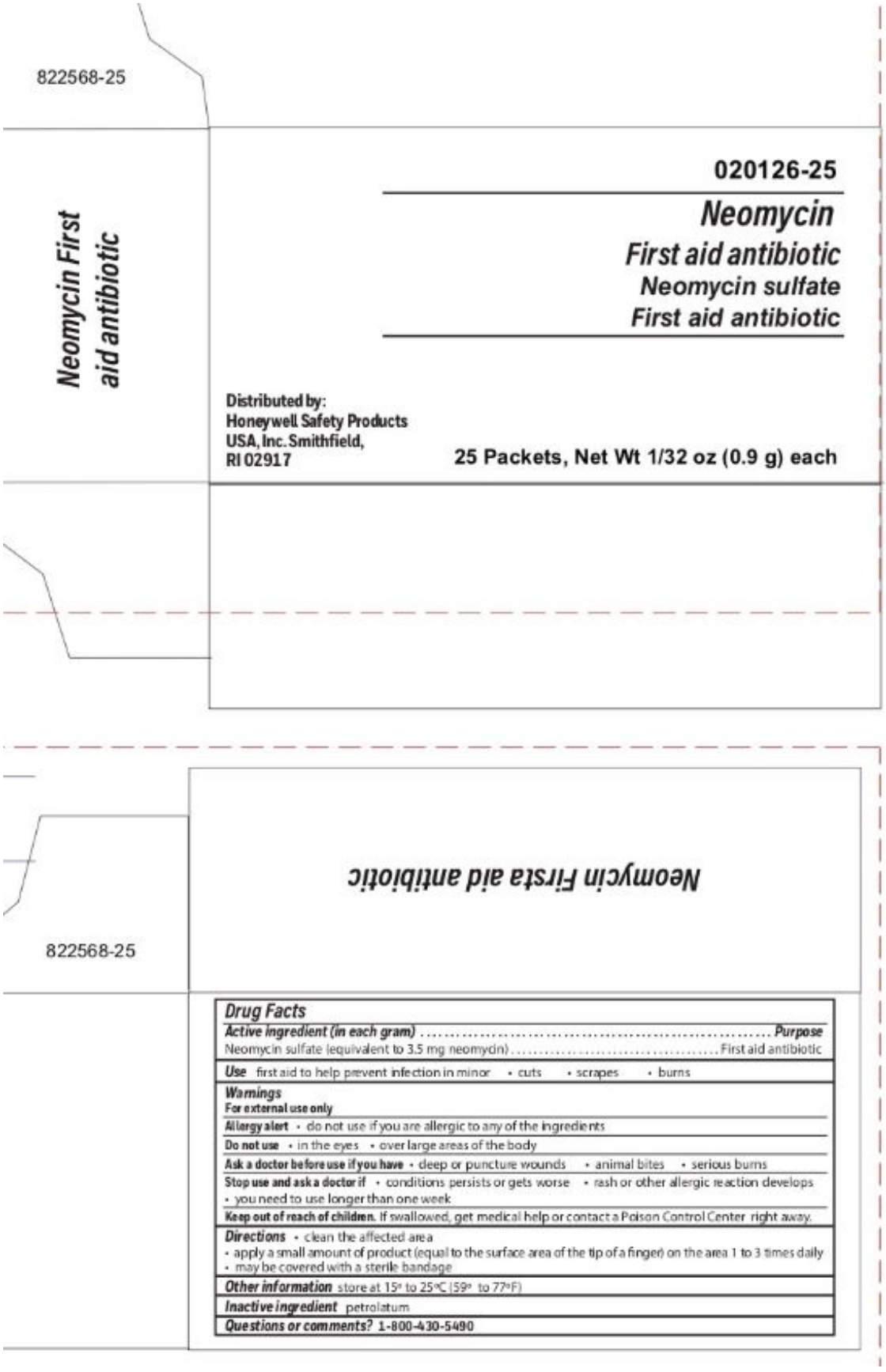


Aypanal
Principal Display Panel



Sting Relief
Principal Display Panel

Neomycin Antibiotic Ointment
Principal Display Panel



4278 Kit Label
SF00002177

46001363 Rev.C
Prints 3 colors
Black, Red (PMS 186) and Blue (PMS 072)

4278 FIRST AID KIT

4278 first aid kit kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4278

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-4278-01	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	3 PACKET	6
Part 3	6 POUCH	2.4 mL
Part 4	10 PACKET	9 g
Part 5	10 PACKET	9 g
Part 6	10 PACKET	14 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		

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

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 3 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		12/23/2017	

other		12/23/2017	

Part 4 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**
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 5 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 6 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	

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