4241 FIRST AID KIT- 4241 first aid kit
4125 FIRST AID KIT- 4125 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.

0498-4125 & 0498-4241: First Aid Kit (BZK wipes, 1st aid Sp, EW, amm. Inh, triple, burn spray WS, alcohol wipe- SF00004255, 68400LNT)

Eyewash
Active ingredient

Sterile Water 99%

Eyewash
Purpose

Eyewash

Eyewash
Uses

- for flushing the eye to remove loose foreign material, air pollutants or chlorinated water

Eyewash
Warnings

For external use only Obtain immediate medical treatment for all open wounds in or near 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.

Eyewash
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

Eyewash
Inactive ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Eyewash
Questions

1-800-430-5490

Ammonia
Active ingredient

Ammonia 15%

Ammonia
Purpose

Respiratory stimulant

Ammonia
Uses

- to prevent or treat fainting

Ammonia
Warnings

For external use only

Do not use

- if you have breathing problems such as asthma or emphysema

Stop use and ask a doctor if

- condition persists

Keep out of reach of children

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

Ammonia
Directions

- hold inhalant away from face and crush ampoule between thumb and forefinger at position indicated on sleeve.
- hold near nostrils for inhalation of volatile vapor

Ammonia
Other information

- store at room temperature away from light

Ammonia
Inactive ingredient

alcohol USP, FD&C red #40, lavender oil, lemon oil fcc, nutmeg oil, purified water

Ammonia
Questions or Comments?

1-800-430-5490

Water Soluble 1st Aid Spray
Active ingredient

Benzethonium chloride 0.2% w/w - Benzocaine 10% w/w

Water Soluble 1st Aid Spray
Purpose

Topical antiseptic

Topical anesthetic

Water Soluble 1st Aid Spray
Uses

for temporary relief of pain and itching and helps protect against infection in

- minor cuts and scrapes
- insect bites
- minor skin irritations

Water Soluble 1st Aid Spray
Warnings

For external use only

Flammable

- keep away from fire or flame
- contents under pressure
- do not puncture or incinerate container
- do not expose to temperature above 120 0 F

Do not use
- in the eyes or other mucous membranes
- in cases of serious burns
- in case of deep orpuncture wounds
- for a prolonged period of time
- on large portion of the body

Stop use and ask a doctor if
- condition worsens or symptoms persist for more than 7 days
- condition clears up and occurs again within a few days
- redness, swelling, or irritation occurs

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

Water Soluble 1st Aid Spray
Directions

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage. If bandaged, let dry first
- for adult institutional use only
- not intended for use on children

Water Soluble 1st Aid pray
Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal

Water Soluble 1st Aid pray
Inactive ingredients

dipropylene glycol, isobutane, N-butane, propane

Burn Relief Water Soluble
Active ingredients

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

Burn Relief Water Soluble
Purpose

Topical antiseptic

Topical anesthetic

Topical anesthetic

Burn Relief Water Soluble
Uses

for the temporary relief of pain and itching and helps protect against infection in:

- minor cuts and scrapes
- burns
- sunburn
- insect bites
- minor skin irritations

Burn Relief Water Soluble
Warnings

For external use only

Flammable keep away from fire or flame

- contents under pressure
- do not puncture or incinerate container
- do not expose to temperatures above 120 0 F

Do not use

- in or near the eyes or other mucous membranes
- in case of serious burns
- in case of deep or puncture wounds
- for prolonged period of time
- on large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- condition clears up and recurs within a few days
- redness, swelling, or irritation occurs

Keep out of the reach of children

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

Burn Relief Water Soluble
Directions

- clean the affected area
- shake can well before using
- hold 4 - 6 inches from surface and spray area until wet
- may be covered with a sterile bandage, if bandaged let dry first
- for adult institutional use only
- not intended for use on children

Burn Relief Water Soluble
Other information

- avoid inhaling
- use only as directed
- intentional misuse by deliberately concentrating or inhaling the contents may be harmful or fatal

Burn Relief Water Soluble
Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Triple
Active ingredients

Bacitracin zinc 400 units

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

Polymyxin B sulfate 5000 units

Triple
Purpose

First aid antibiotic

First aid antibiotic

First aid antibiotic

Triple
Uses

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

Triple
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
- a 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 the reach of children

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

Triple
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

Triple
Other information

- store at 15 ^{0} to 25 ^{0} C (59 ^{0} to 77 ^{0} F)
- tamper evident sealed packets
- do not use if packet is torn or opened

Triple
Inactive ingredient

petrolatum

Alcohol
Active ingredient

Isopropyl alcohol 70%

Alcohol
Purpose

First aid antiseptic

Alcohol
Uses

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

Alcohol
Warnings

For external use only

Flammable, keep away from fire or flame.

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

When using this product

- do not use longer than one week unless directed by a doctor

Stop use and consult a doctor

- if condition persists or gets worse

Keep out of reach of children

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

Alcohol
Directions

- clean the affected area
- apply wipe to affected area 1 to 3 times daily
- may be covered with a sterile bandage
- discard wipe after single use

Alcohol

Other information

store at room temperature 15 ^{0} to 25 ^{0} C (59 ^{0} to 77 ^{0} F)

Alcohol
Inactive ingredient

water

BZK
Active ingredient

Benzalkonium chloride 0.13% w/v

BZK
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
Other information

- store at room temperature 15 ^{0} to 30 ^{0} C (59 ^{0} - 86 ^{0} F)
- do not reuse towelette

BZK
Inactive ingredient

water

BZK
Questions

1-800-430-5490

4125
SF00004255 kit contents

1 3/4 X 3 PLAS 100/BOX

1 1X3 PLASTIC 100/BOX

1 FNGERTP WOVEN REG 40/BOX

1 WOVEN 2" X 3" 25/BOX

1 1X3 WOVEN SING 50/BOX

1 SWIFT KNUCKLE 40/BX

1 AMMONIA INHALANTS 10 PER

1 INSTANT COLD PACK 4" X 6"

1 ELASTIC TAPE 1" X 5YD

1 O/H TAPE ADHESIVE TRI-CUT

1 FIRST AID GUIDE ASHI

6 GAUZE CLEAN-WRAP BDGE N/S 2"

2 BLOODSTOPPER

1 NON ADHERENT PADS 2"X3" 50'S

1 GZE PADS STERILE 2"X 2" 25'S

1 GZE PADS STERILE 4"X 4" 25'S

1 CO-FLEX BANDAGE 2"X 5YDS TAN

1 COTTON TIPS 100 PER VIAL

1 ANTISEPTIC WIPES BZK CHL 20'S

1 FIRST AID SPRAY AEROSOL 3 OZ

1 ALCOHOL WIPES 50'S

1 BURN SPRAY 3 OZ

1 TRIPLE BIOTIC .5 GRAM PKT 20

1 QUICK RELIEF 1 PACK OF 2/BOX

1 COLD PACK 5"X9" BOXED

1 4OZ BFS EYEWASH TRILINGUAL BOTTLE

1 SCISSOR BDGE 4" RED PLS HDL

1 SPLINTER FORCEP 4 1/2"

1 SCISSOR LISTER BDG S/S 5 1/2"

1 400 EMPTY KIT BLANK

1 POCKET INSERT RED #400 KIT 5R

1 TONGUE BLADES SR WRAPPED 6'S

1 ISO-SHIELD CPR ADULT/CHLD 1BG

1 LBL STOCK 6-3/8"X4"

1 LBL STOCK 4"X2-7/8"

1 LBL STOCK 3"x1-7/8"

2 TRI BNDG NON WOVEN 40"X40"X56"

1 RESCUE BLANKET 1EA

1 RED BIO BAGS 2/BX

6 NITRILE GLOVES 1 PR

4241
68400LNT KIT CONTENTS

1 3/4 X 3 PLAS 100/BOX

1 1X3 PLASTIC 100/BOX

1 WOVEN 2" X 3" 25/BOX

1 FINGERTIP "T" WOVEN 40/BOX

1 1X3 WOVEN SING 50/BOX

1 SWIFT KNUCKLE 40/BX

1 AMMONIA INHALANTS 10 PER

1 ELASTIC TAPE 1" X 5YD

1 FORCEPS POINTED METAL

1 O/H TAPE ADHESIVE TRI-CUT

1 FIRST AID GUIDE ASHI

6 GAUZE CLEAN-WRAP BDGE N/S 2"

2 BLOODSTOPPER

1 NON ADHERENT PADS 2"X3" 50'S

2 GZE PADS STERILE 2"X 2" 25'S

1 GZE PADS STERILE 4"X 4" 25'S

1 CO-FLEX BANDAGE 2"X 5YDS TAN

1 COTTON TIPS 100 PER VIAL

1 ANTISEPTIC WIPES BZK CHL 20'S

1 FIRST AID SPRAY AEROSOL 3 OZ

1 ALCOHOL WIPES 50'S

1 BURN SPRAY 3 OZ

1 TRIPLE BIOTIC .5 GRAM PKT 20

1 COLD PACK 5"X9" BOXED

1 4OZ BFS EYEWASH

1 SCISSOR BDGE 4" RED PLS HDL

1 KIT TWEEZER 3 1/2" SLANTED

1 400 EMPTY KIT BLANK

1 POCKET INSERT RED #400 KIT 5R

1 TONGUE BLADES SR WRAPPED 6'S

1 LBL STOCK 6-3/8"X4"

1 LBL STOCK 4"X2-7/8"

1 LBL STOCK 3"x1-7/8"

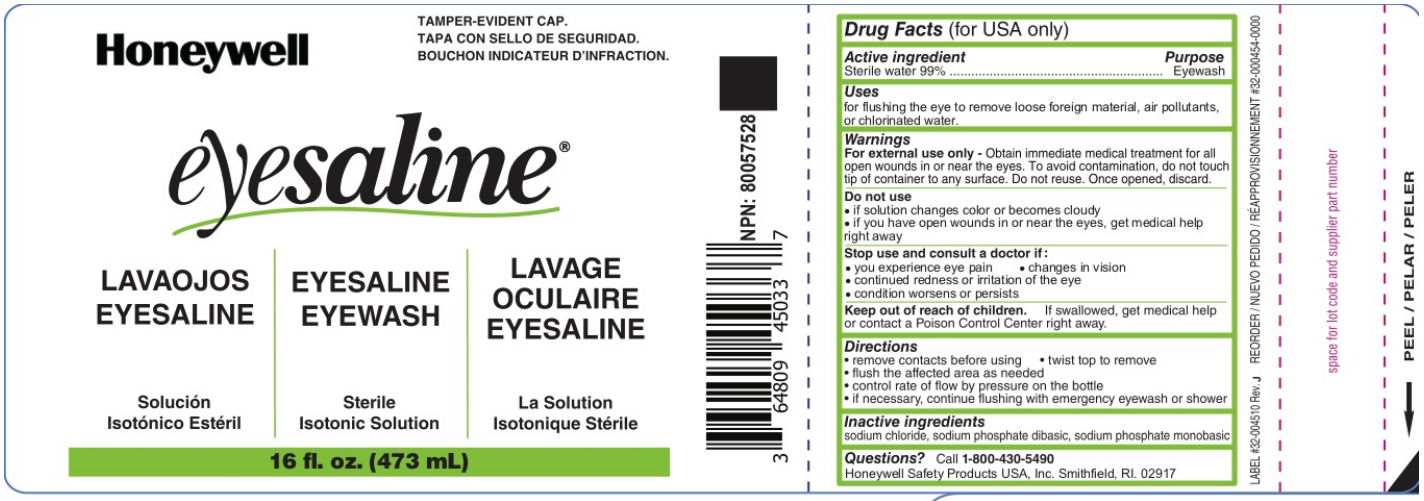
2 PR LRG NITRILE GLVES ZIP BAG

2 ADHES TAPE EYE STRIPS 2'S

2 TRI BNDG NON WOVEN 40"X40"X56"

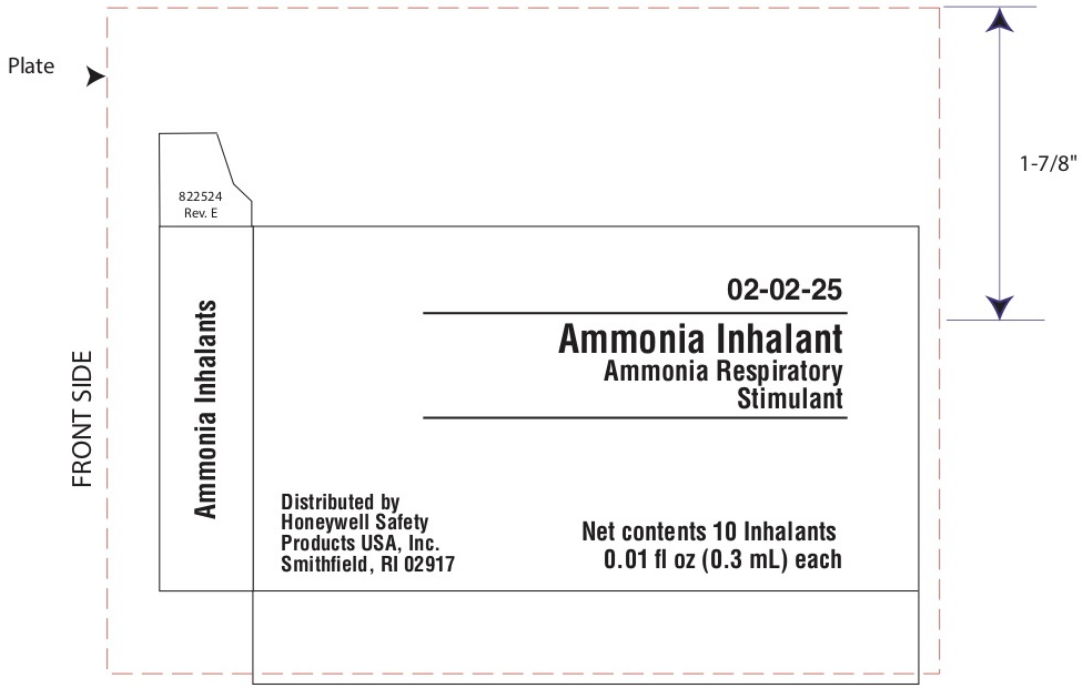
1 RED BIO BAGS 2/BX

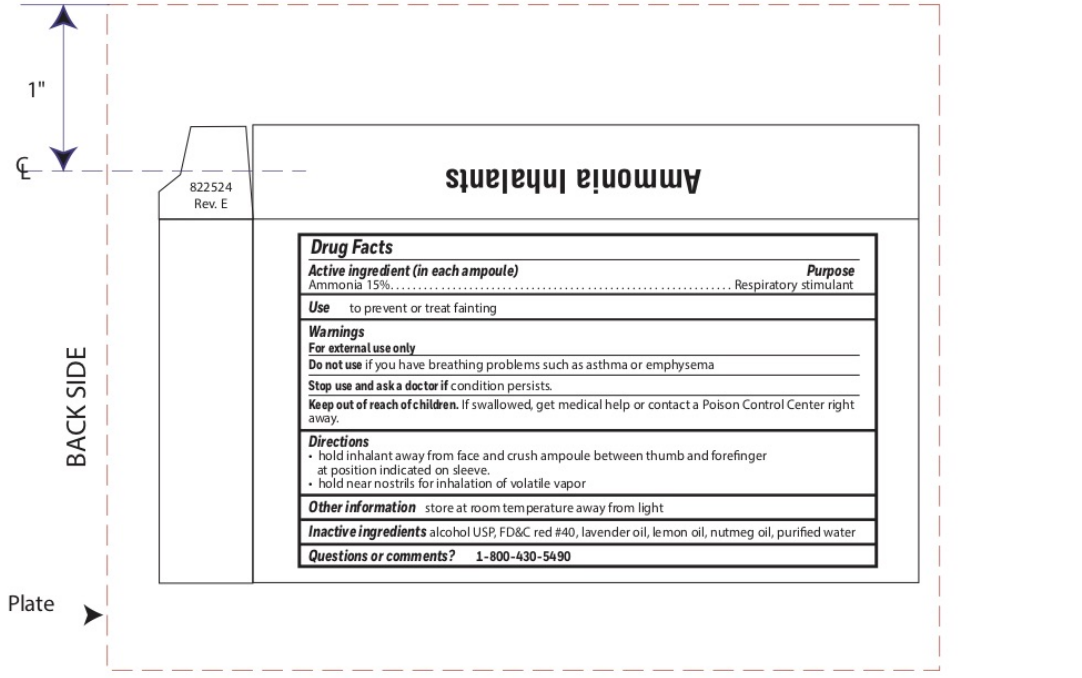
Eyewash
Principal Display Panel



Ammonia
Principal Display Panel

796006 Rev. E Unit Carton Printing Plate for "A" size carton.





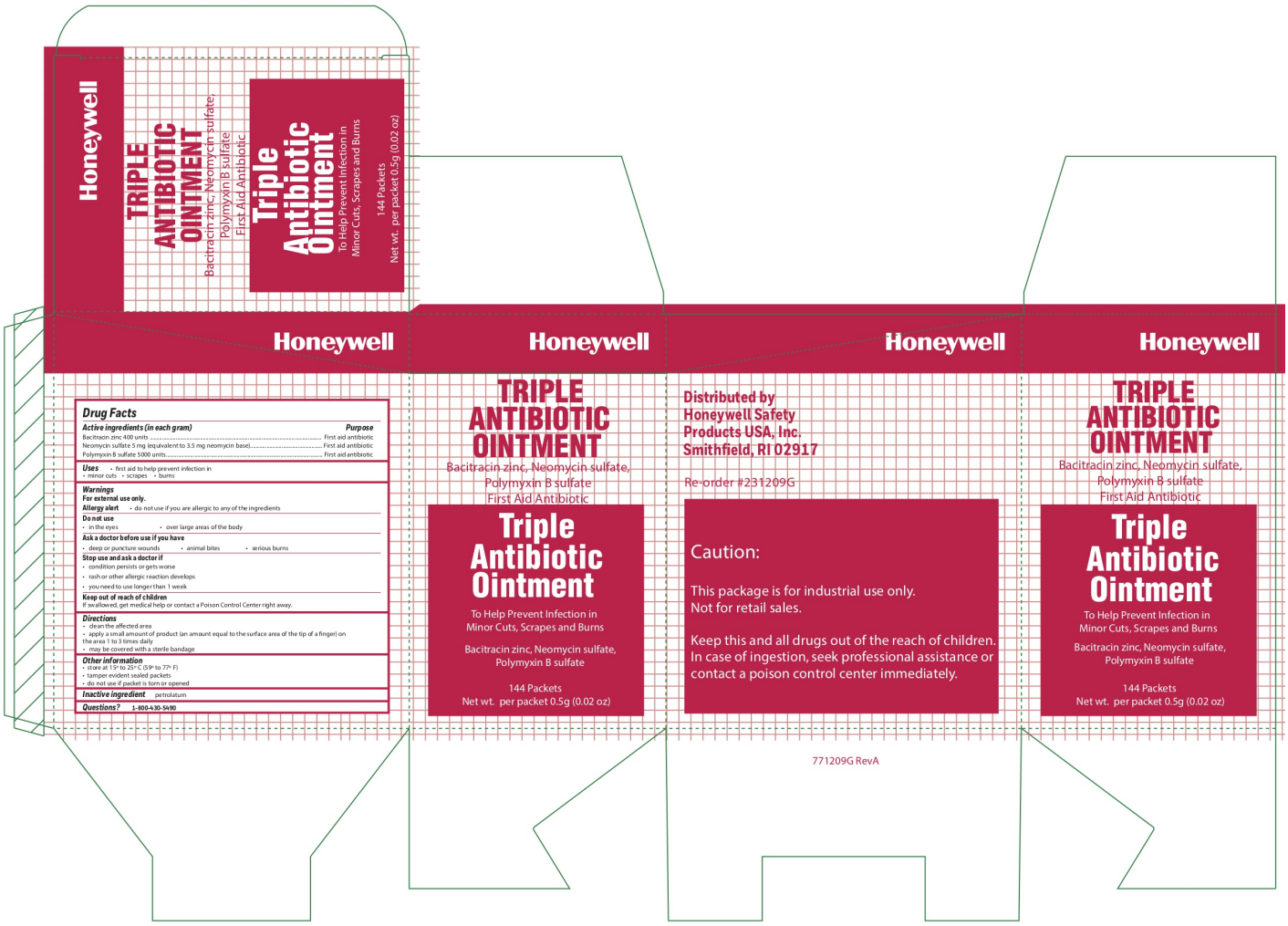
796006 Rev. E (page 3 of 3)

Water Soluble 1st Aid Spray

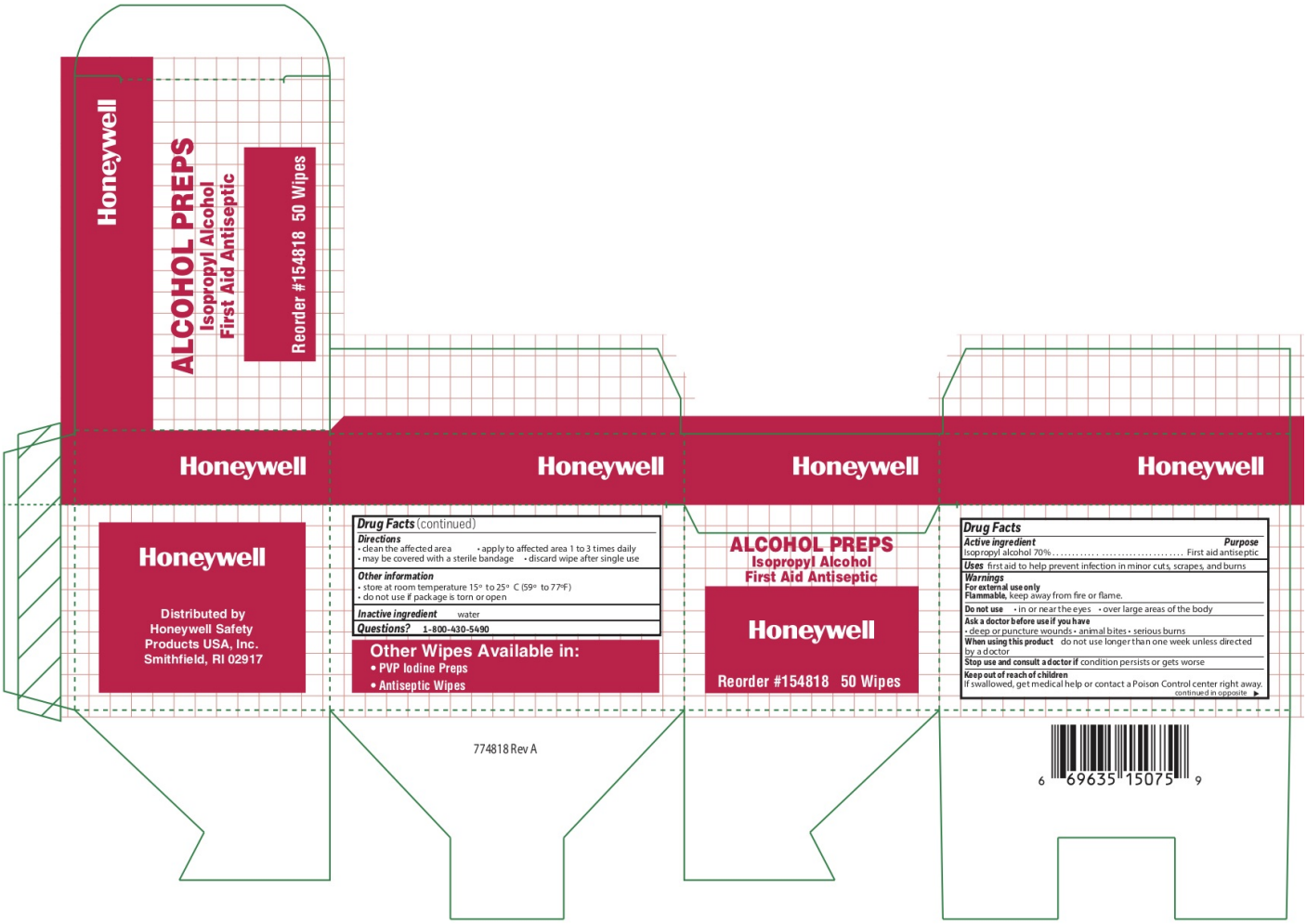
Principal Display Panel

Burn Relief Water Soluble
Principal Display Panel

Triple
Principal Display Panel



Alcohol
Principal Display Panel



BZK
Principal Display Panel

4125 Kit Label
SF00004255

First Aid

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

4241 Kit Label
68400LNT

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

4241 FIRST AID KIT

4241 first aid kit kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4241

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4241-01	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	118 mL
Part 2	10 AMPULE	3 mL
Part 3	1 CAN	85 g
Part 4	1 CAN	85 g
Part 5	20 PACKET	10 g
Part 6	50 POUCH	20 mL
Part 7	20 PACKET	28 mL

Part 1 of 7

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 PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-02	118 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 7

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 7

FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

Product Information	
Item Code (Source)	NDC:0498-0031
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0031-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/19/2018	

Part 4 of 7

BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

Product Information	
Item Code (Source)	NDC:0498-0021
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.33 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0021-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/12/2018	

Part 5 of 7

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information	
Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g

NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g

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

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0750-36	0.5 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		09/19/2018	

Part 6 of 7

ALCOHOL WIPE

isopropyl alcohol swab

Product Information	
Item Code (Source)	NDC:0498-0143
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0143-04	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		09/18/2018	

Part 7 of 7

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
unapproved drug other		12/22/2017	

Marketing Information

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

4125 FIRST AID KIT

4125 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4125

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4125-01	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	118 mL
Part 2	10 AMPULE	3 mL
Part 3	1 CAN	85 g
Part 4	1 CAN	85 g
Part 5	20 PACKET	10 g
Part 6	50 POUCH	20 mL
Part 7	20 PACKET	28 mL

Part 1 of 7

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 PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0100-02	118 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 7

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information	
Item Code (Source)	NDC:0498-3334
Route of Administration	RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Inactive Ingredients	
Ingredient Name	**Strength**

ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-3334-00	0.3 mL in 1 AMPULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/18/2018	

Part 3 of 7

FIRST AID ANTISEPTIC WATER SOLUBLE

benzethonium chloride, benzocaine spray

Product Information	
Item Code (Source)	NDC:0498-0031
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0031-	85 g in 1 CAN; Type 0: Not a Combination		

1	40	Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/19/2018	

Part 4 of 7

BURN WATER SOLUBLE

benzocaine, benzethonium chloride, menthol spray

Product Information

Item Code (Source)	NDC:0498-0021
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.2 g in 100 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	10 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.33 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0021-40	85 g in 1 CAN; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

unapproved drug other		11/12/2018	

Part 5 of 7

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information	
Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g

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

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0750-36	0.5 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		09/19/2018	

Part 6 of 7

ALCOHOL WIPE

isopropyl alcohol swab

Product Information	
Item Code (Source)	NDC:0498-0143
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0143-04	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		09/18/2018	

Part 7 of 7

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
unapproved drug other		12/22/2017	

Marketing Information

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

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