4360 FIRST AID KIT- 4360 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.

0498-4360: First Aid Kit (Neomycin, EW, alcohol wipes, HC cr, Burn Sray, Antiseptic Spray, Hand Sanitizer- FRKSOFTPAK-CLSB)

Eyesaline
Active ingredient

Sterile Water 99%

Eyesaline
Purpose

Eyewash

Eyesaline
Uses

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

Eyesaline
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.

Eyesaline
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

Eyesaline
Inactive ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Eyesaline
Questions

1-800-430-5490 Honeywell Sadety Products USA, Inc. Smithfield, RI 02917

Alcohol Wipe
Active ingredient

Isopropyl alcohol 70%

Alcohol Wipe
Purpose

First aid antiseptic

Alcohol Wipe
Uses

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

Alcohol Wipe
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 burn

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 Wipe
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 Wipe
Other information

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

Alcohol Wipe
Inactive ingredient

water

Alcohol Wipe
Questions

1-800-430-5490

Neomycin
Active ingredient

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

Neomycin
Purpose

First aid antibiotic

Neomycin
Uses

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

Do not use

- in the eyes
- over large areas of the body

Neomycin
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

- a rash or other allergic reaction develops
- you need to use longer than 1 week

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
Direction

- 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
Other information

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

Neomycin
Inactive ingredient

petrolatum

Neomycin
Questions?

1-800-430-5490

Antiseptic Spray
Active ingredient

Benzalkonium chloride 0.13%

Antiseptic Spray
Purpose

First aid antiseptic

Antiseptic Spray
Uses

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

Antiseptic Spray
Warnings

For external use only

Do not use
- in or near 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 ask a doctor if

- 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

Antiseptic Spray
Directions

- clean the affected area
- spray a small amount of this product on the area 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first

Antiseptic Spray
Other information

- shake well
- store at room temperature 15 ^{0}-30 ^{0} C (59 ^{0} -86 ^{0} F)

Antiseptic Spray
Inactive ingredients

diazolidinyl urea, edetate disodium, glycerin, hypromellose, methylparaben, octoxynol 9, propylene glycol, propylparaben, trolamine, water

Antiseptic Spray
Questions

1-800-430-5490

Burn Spray
Active ingredient

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

Burn Spray
Purpose

Benzethonium chloride 0.2% w/w

Benzocaine 10% w/w

Menthol 0.33% w/w

Burn Spray
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 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 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 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

Burn Spray
Other information

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

harmful or fatal

Burn Spray
Inactive ingredients

dipropylene glycol, isobutane, n-butane, propane

Hans Sanitizer
Active ingredient

Ethyl alcohol 62%

Hand Santizer
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

store at 15 ^{0} to 25 ^{0} C (59 ^{0} to 77 ^{0} 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, triisopropanolamine

Hand Sanitizer
Questions or Comments?

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

Hydrocortisone
Active ingredient (in each gram)

Hydrocortisone acetate (equivalent to Hydrocortisone 1%)

Hydrocortisone
Purpose

Anti-itch cream

Hydrocortisone
Uses

- for the temporary relief of itching associated with minor skin irritations and rashes

Hydrocortisone
Warnings

For external use ;only

Ask a doctor before use if

- if you are using any other hydrocortisone product

When using the product

- avoid contact with eyes
- do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash

Stop use and ask a doctor if

- condition worsens
- condition persists for more than 7 days
- condition clears up and recurs within a few days

Keep out of reach of children

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

Hydrocortisone
Directions

- adults and children 2 years and older:
- clean the affected area
- apply to the area not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor

Hydrocortisone
Other information

- store at room temperature (do not freeze)

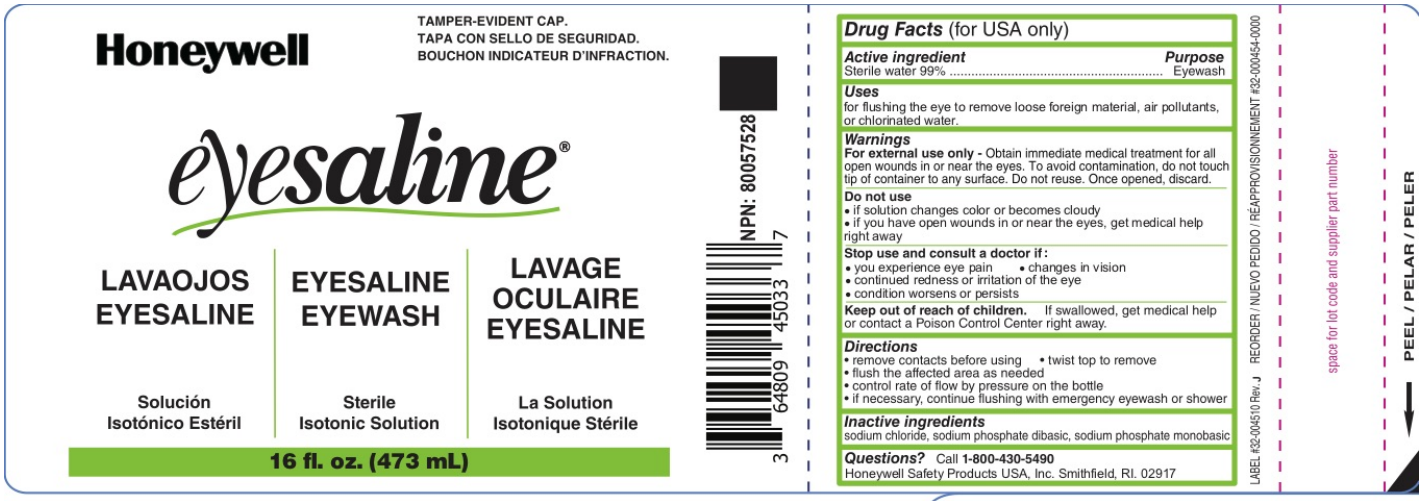
Hydrocortisone
Inactive ingredients

cetyl alcohol, citric acid, diazolidinyl urea, edetate disodium, glycerin, glyceryl monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol, propylparaben, purified water, stearic acid, trolamine

Hydrocortisone
Questions or Comments?

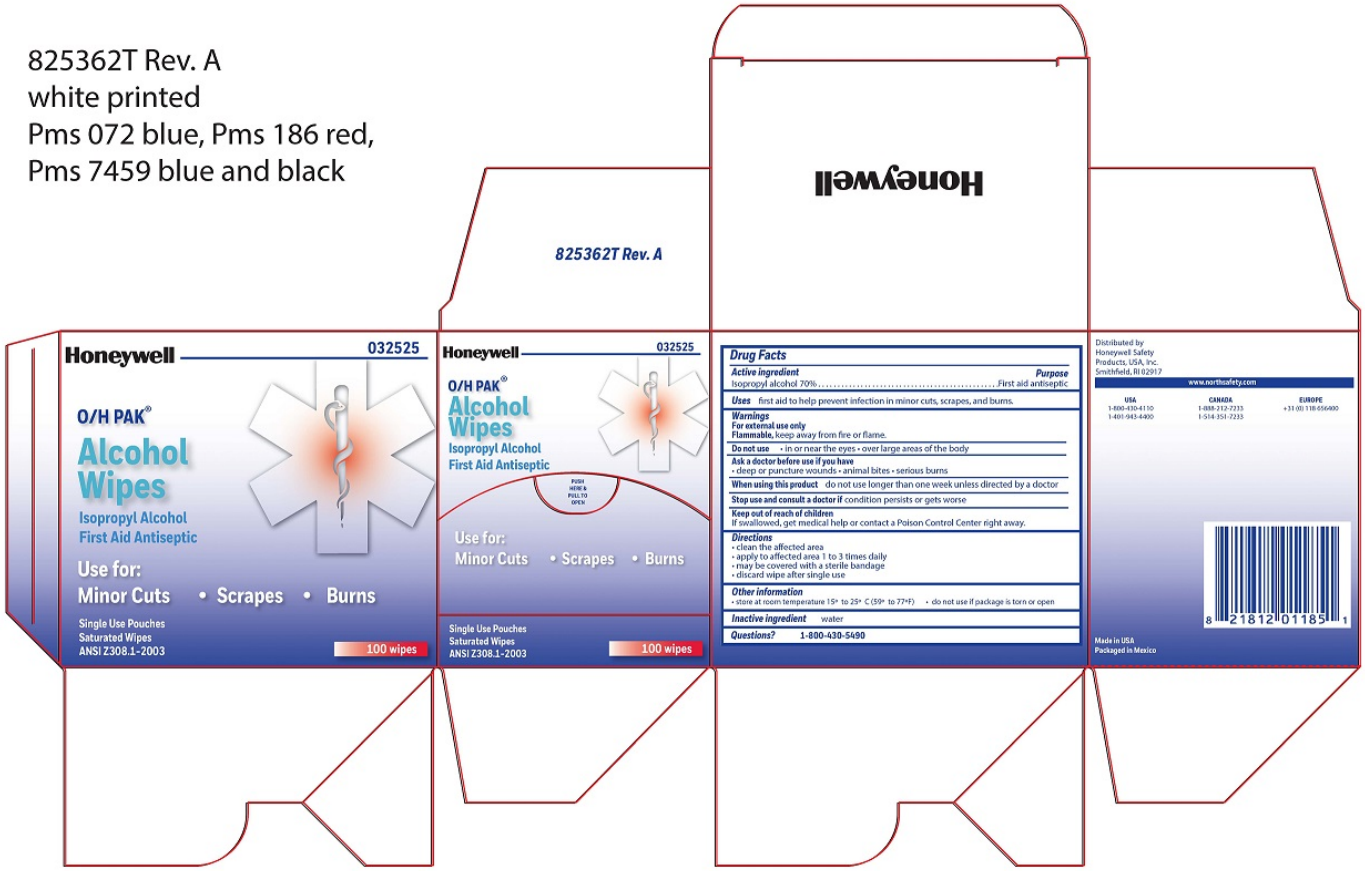
1-800-430-5490

Eyesaline
Principal Display Panel



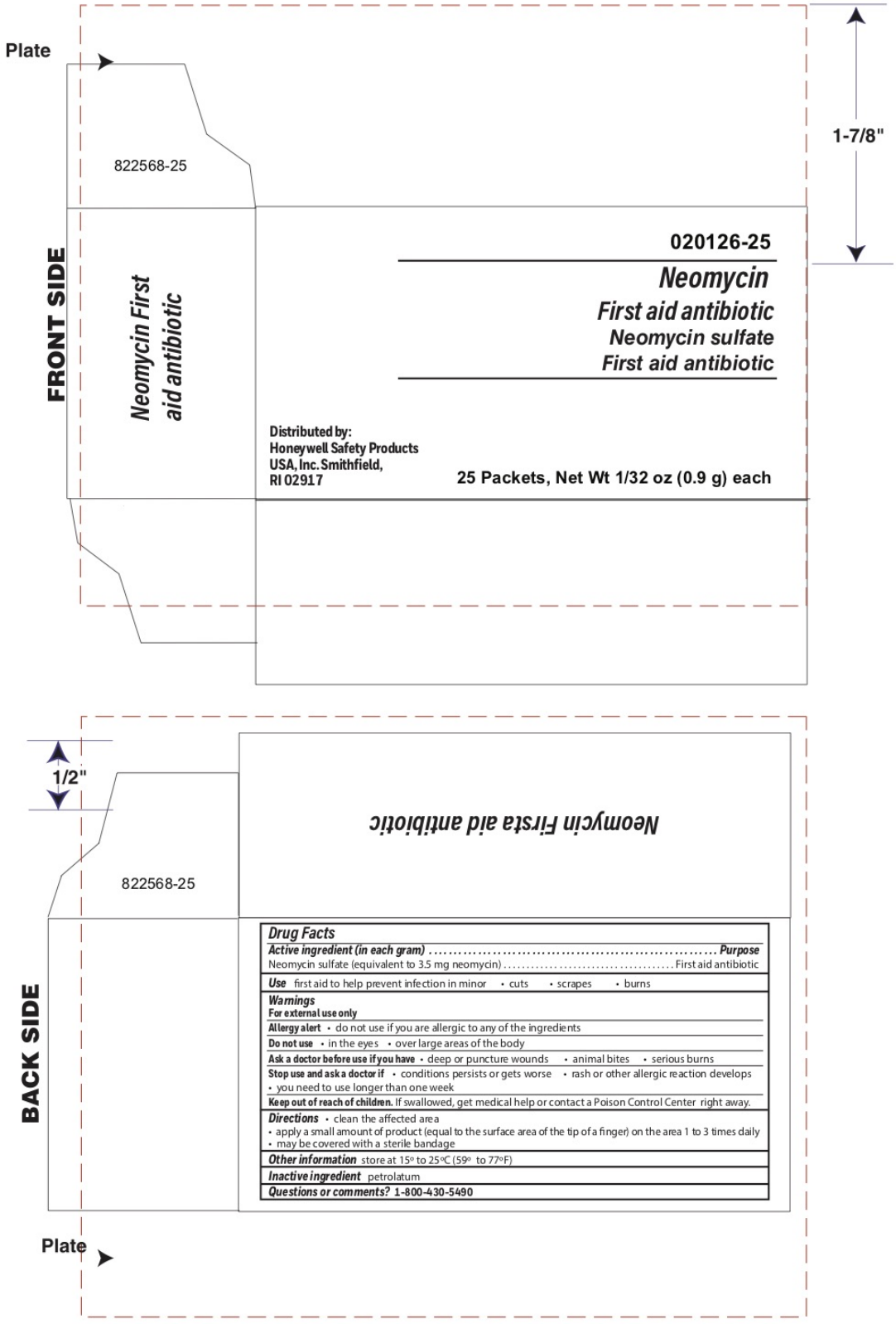
Alcohol Wipe
Principal Display Panel

825362T Rev. A
white printed
Pms 072 blue, Pms 186 red,
Pms 7459 blue and black



Neomycin
Principal Display Panel

796041-25 Rev A Unit Carton Printing Plate for "**C**" size carton.



Antiseptic Spray

Principal Display Panel

Honeywell 032203

ANTISEPTIC SPRAY

Benzalkonium chloride

First Aid Antiseptic

Net contents 2 fl oz (59 mL)

ANSI Z308.1-2003

8 21812 01498 2

Drug Facts

Active ingredient — **Purpose**
Benzalkonium chloride 0.13% ..... First aid antiseptic

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

Warnings
For external use only

Do not use • in or near 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 ask 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.

Directions • clean the affected area
• spray a small amount of this product on the area 1 to 3 times daily
• may be covered with a sterile bandage • if bandaged, let dry first

Other information • shake well
• store at room temperature, 15° to 30°C (59° to 86°F)

Inactive ingredients diazolidinyl urea, edetate disodium, glycerin, hypromellose, methylparaben, octoxynol 9, propylene glycol, propylparaben, trolamine, water

Questions or comments? 1-800-430-5490

Mfg. for: Honeywell Safety Products USA, Inc. Smithfield, RI 02917 032203 Rev. G

Burn Spray
Principal Display Panel

SHAKE WELL BEFORE USING

Honeywell

BURN SPRAY

Water soluble
Benzethonium chloride
Topical antiseptic
Benzocaine
Topical anesthetic
Menthol
Topical anesthetic

Provides antiseptic treatment and helps relieve the pain of minor burns and sunburn.

CAUTION: FLAMMABLE
Contents under pressure
Read warning on back panel.

NET WT. 3 OZ (85gm.)

47.0306

Cat. No. 201005

DRUG FACTS

Active ingredients — **Purpose**
Benzethonium chloride 0.2% w/w ..... Topical antiseptic
Benzocaine 10% w/w ..... Topical anesthetic
Menthol 3.3% ..... Topical anesthetic

Uses • for the temporary relief of pain and itching and helps to protect against infection in • minor cuts and scrapes • burns • sunburn • insect bites • minor skin irritations

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°F

Do not use • in or near eyes or other mucus membranes • in case of serious burns • in case of deep or puncture wounds • for a prolonged period of time • on large portion of the body

Stop use and ask a doctor if:
• conditions 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 reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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

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

Inactive ingredients dipropylene glycol, isobutane, n-butane, propane

Questions or comments? 1-800-430-5490

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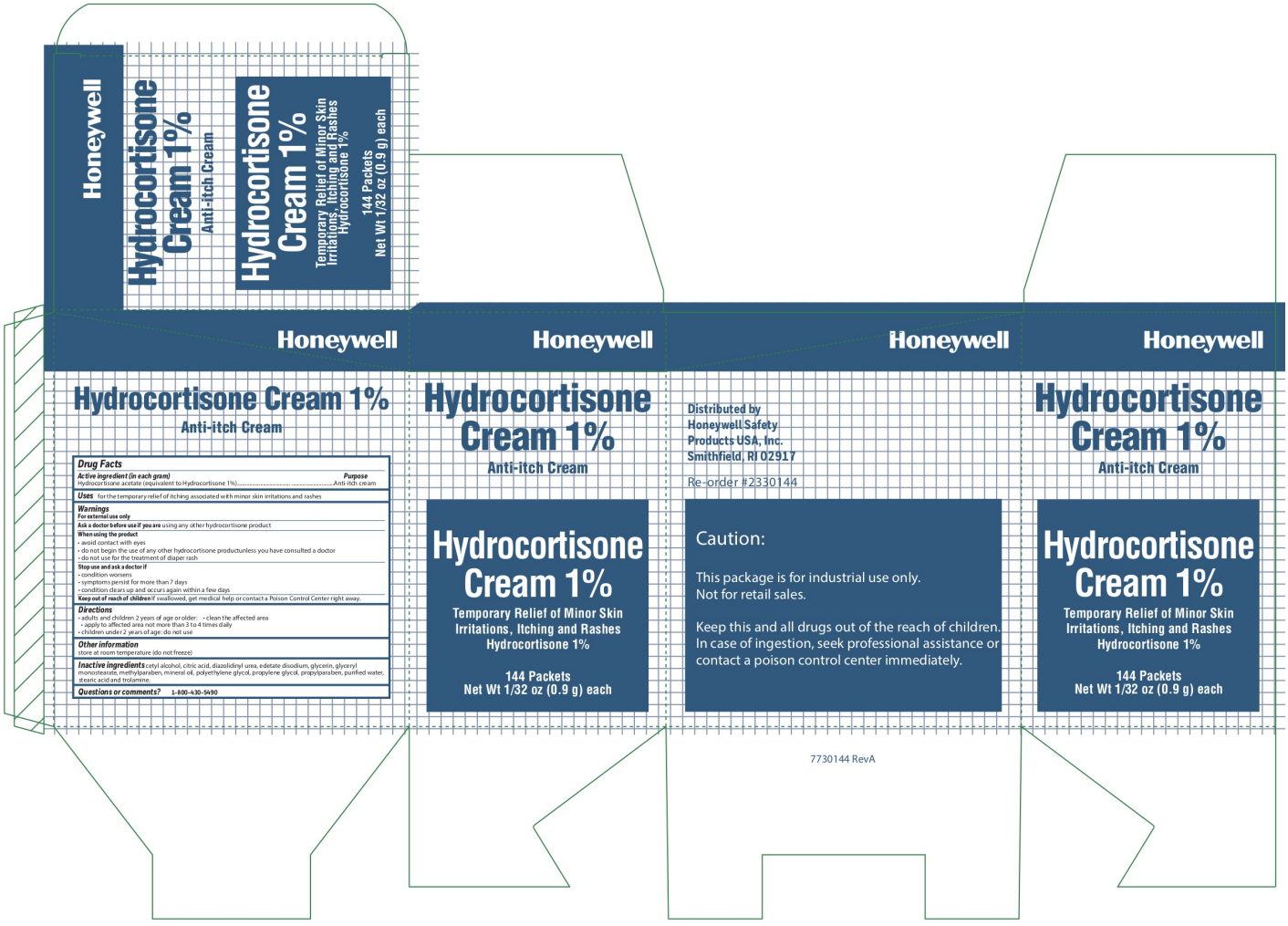
Honeywell Safety
Products USA, Inc.
Smithfield, RI 02917

Honeywell

Hand sanitizer
Principal Display Panel

Hydrocortisone
Principal Display Panel



4360 Kit Label
FRKSOFTPAK-CLSB

Honeywell

First Aid Kit

www.honeywellsafety.com

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

754000 Rev. E

4360 Kit Contnets

FRKSOFTPAK-CLSB

1 1 X 3 WOVEN 100/BOX

3 NEOMYCIN ANTIBIOTIC 10 PER

2 EYE DRESS PKT W/4 ADH STRIPS

1 TOURNIQUET, 1 PER

1 WIRE SPLINT 1 PER

1 RESCUE BLANKET 1 PER

1 GAUZE COMP, 18" X 36", 2 PER

1 BANDAGE COMP, 2" OFFSET, 4 PER

1 ALCOHOL PREP PADS 10P

1 HYDROCORTISON,1.O%,1/32 OZ,10P

1 O/H PAK,ADH BDG 2"X4", X-LG,10 PER

1 O/H PUMP ANTISEPTIC 2 OZ ID F

1 O/H PUMP BURN RELIEF 2 OZ ID G

1 FIRST AID GUIDE ASHI

1 TAPE ADHESIVE 1"X 5 YD PLSTC

10 HAND SANITIZER 0.9G WJ BULK

2 GAUZE CLEAN-WRAP BDGE N/S 2"

1 GAUZE CLEAN-WRAP BDGE N/S 4"

4 BLOODSTOPPER

1 GZE PADS STERILE 3"X 3" 10'S

1 ELASTIC BANDAGE 3" X 4.5YD

1 CPR FILTERSHIELD 77-100

1 4OZ BFS EYEWASH TRILINGUAL BOTTLE

1 SCISSOR BDGE 4" RED PLS HDL

1 KIT TWEEZER 3 1/2" SLANTED

LBL STOCK 6-3/8"X4"

LBL STOCK 4"X2-7/8"

1 LBL STOCK 3"x1-7/8"

4 PR LRG NITRILE GLVES

2 WATER-JEL BURN DRESSING 4 X 4

1 KIT BAG SOFT PACK LARGE

1 LBL CONTENTS ANSI 2015 CL B

2 TRI BNDG NON WOVEN 40"X40"X56"

2 COLD PACK UNIT 4"X6" BULK

4 WOVEN FINGERTIP BANDAGE 2"

6 WOVEN KNUCKLE BANDAGE

4360 FIRST AID KIT

4360 first aid kit kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4360

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-4360-01	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 POUCH	4 mL
Part 3	10 PACKET	9 g
Part 4	1 BOTTLE, SPRAY	59 mL
Part 5	1 BOTTLE, SPRAY	59 mL
Part 6	30 PACKET	27 g
Part 7	10 PACKAGE	9 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 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:0498-0100-02	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

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 3 of 7

HYDROCORTISONE

anti-itch cream

Product Information

Item Code (Source)	NDC:0498-0801
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0801-35	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		10/15/2019	

Part 4 of 7

BURN RELIEF

lidocaine hydrochloride spray

Product Information

Item Code (Source)	NDC:0498-0221
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	24.64 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
OCTOXYNOL-9 (UNII: 7JPC6Y25QS)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TEA TREE OIL (UNII: VIF565UC2G)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:0498-0221-59	59 mL in 1 BOTTLE, SPRAY; 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 5 of 7

ANTISEPTIC

benzalkonium chloride spray

Product Information

Item Code (Source)	NDC:0498-0402
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 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
OCTOXYNOL 9 (UNII: 7JPC6Y25QS)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0402-59	59 mL in 1 BOTTLE, SPRAY; 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 6 of 7

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 7 of 7

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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
WATER (UNII: 059QF0KO0R)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59898-420-36	0.9 mL in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/15/2010	

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