4141 FIRST AID KIT- 4141 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.

0498-4141: First Aid Kit (Eye Wash, Hand Sanitizer, FABC, Neomycin, alcohol wipe- FAKREFU-B)

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

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

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 Santitizer
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 º to 25 º C (59 º to 77 º 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

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

Flammable, keep away from fire and flame

Do not use
- in or near 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 1 week unless directed by a doctor

Stop use and consult a doctor if
- condition persists or gets worse

Keep out of the reach of children

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

Alcohol Wipe
Directions

- store at room temperature 15 o to 25 o C (59 o to 77 oF)
- do not use if packet is torn or opened

Alcohol Wipe
Inactive ingredient

water

Alcohol Wipe
Questions

1-800-430-5490

4141
FAKREFU-B Kit Contents

1 HAND SANITIZER 10/PER

1 NEOMYCIN OINT 0.9 GM , UNTZD 25/BX

1 EYE DRESS PKT W/4 ADH STRIPS

1 GAUZE BANDAGE, 4" X 6 YD

1 TOURNIQUET

2 TRIANGULAR BDG, NON-STERILE

1 WIRE SPLINT 1 PER

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

1 ADH TAPE, .5" X 2.5 YD, 2 PER

2 BANDAGE COMP 24" X 72", UNTZD 2/BX

1 FORCEPS & SCISSORS, 1 EA

1 GAUZE BANDAGE, 2" X 6 YD,2 PER

2 INSTANT COLD PACK 4" X 6"

1 1" X 3" PLAS STRIP BAN, UNITZD 50/BX

1 BURN CREAM POUCHES 25/EA

2 BURN-STOP BURN DRESSING 4 X 4

1 ALCOHOL WIPES, UNITIZED 50/BX

2 NITRILE GLOVES 2PR BBP

1 FIRST AID GUIDE ASHI

1 CPR MICROSHIELD DOUBLE UNIT

1 4OZ BFS EYEWASH TRILINGUAL BOTTLE

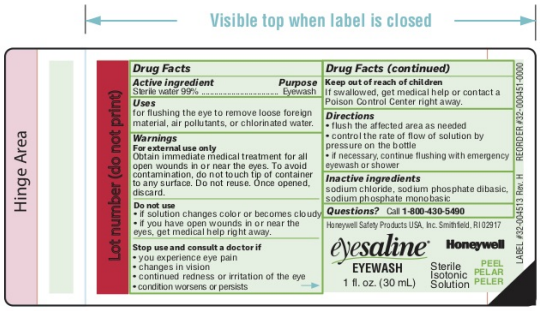
LBL STOCK 6-3/8"X4"

1 LBL STOCK 3"x1-7/8"

1 ZIP LOCK BAG 14 X 20 1.5 MIL

Eye Wash Package label

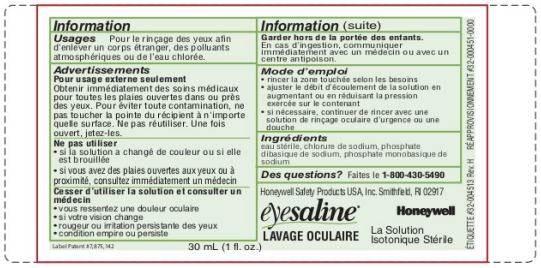
#32-004513 Rev. H



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)

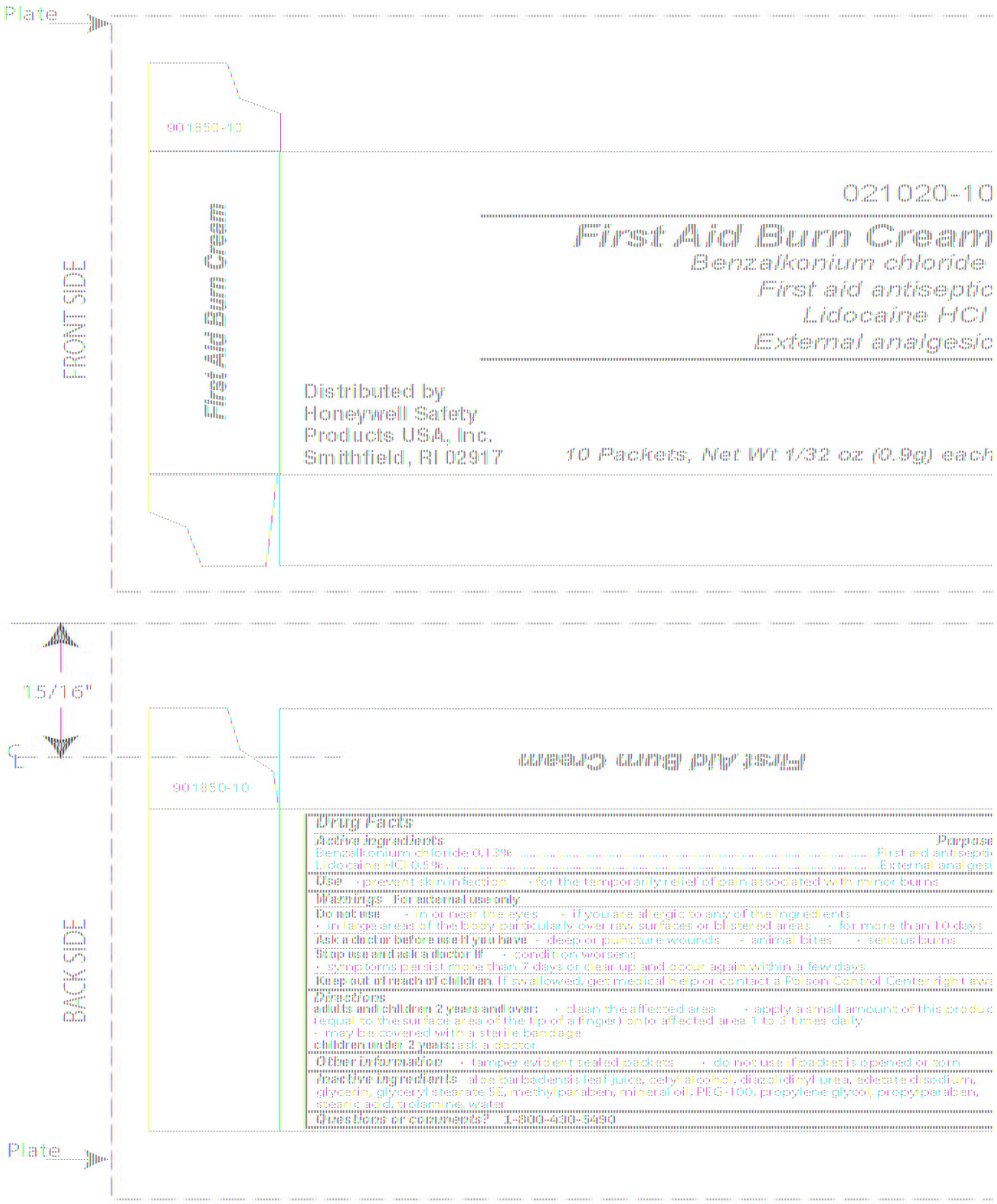


Base
3/32″ from die edges

Printable Text Area

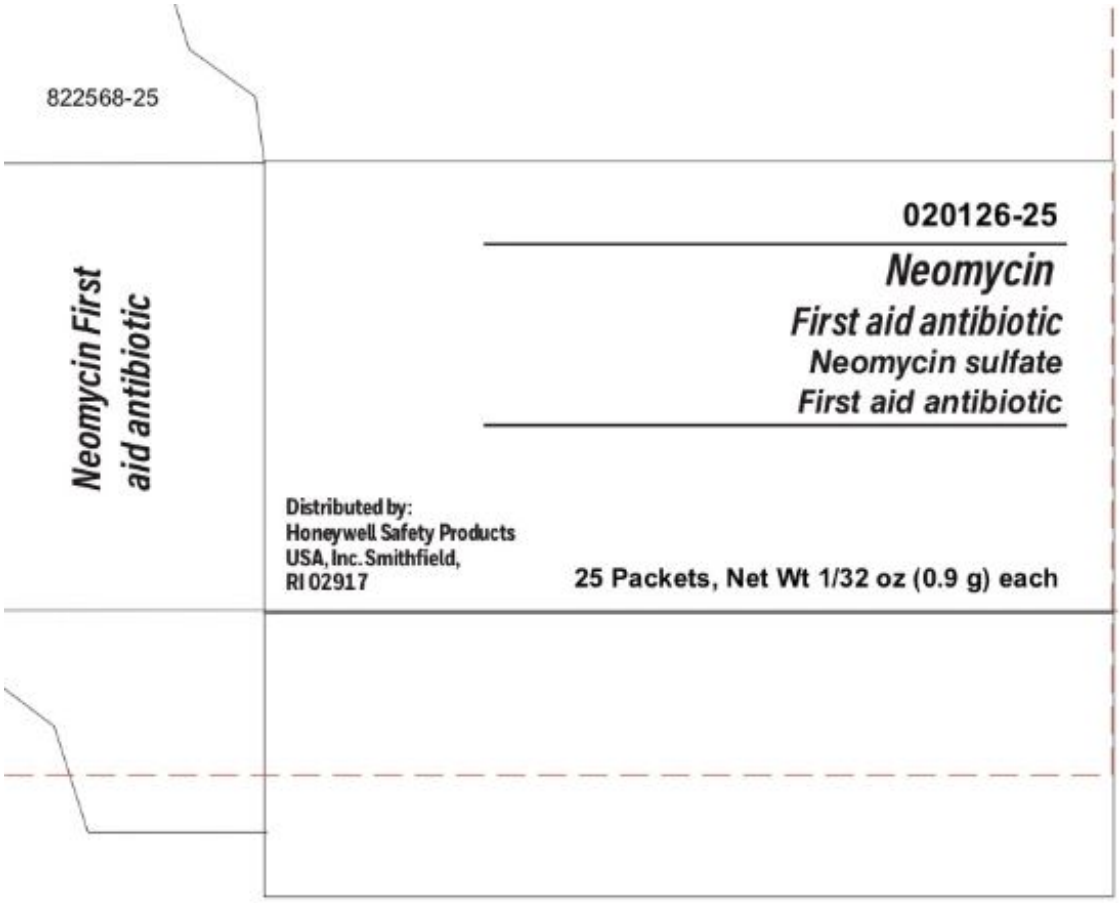
First Aid Burn Cream
Principal Display Panel

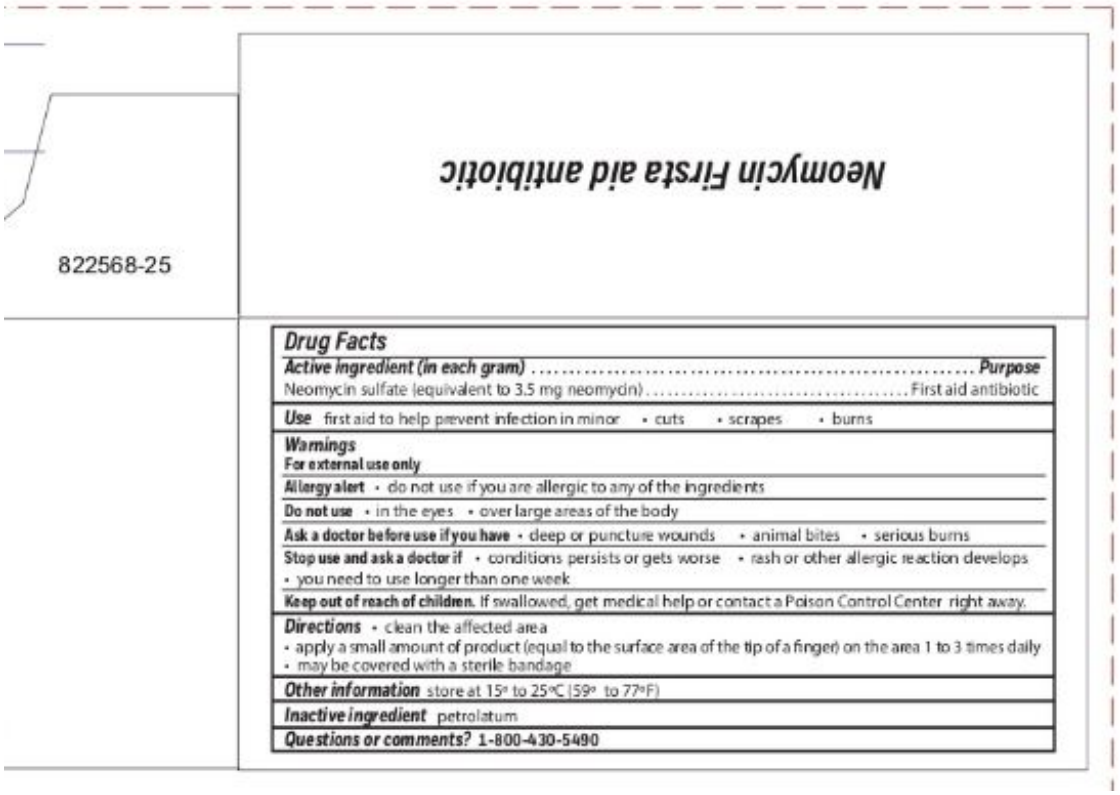
796353-10 Rev. A Unit Carton Printing Plate for "B" size



Neomycin Antibiotic Ointment

Principal Display Panel





Hand Sanitizer
Principal Display Panel

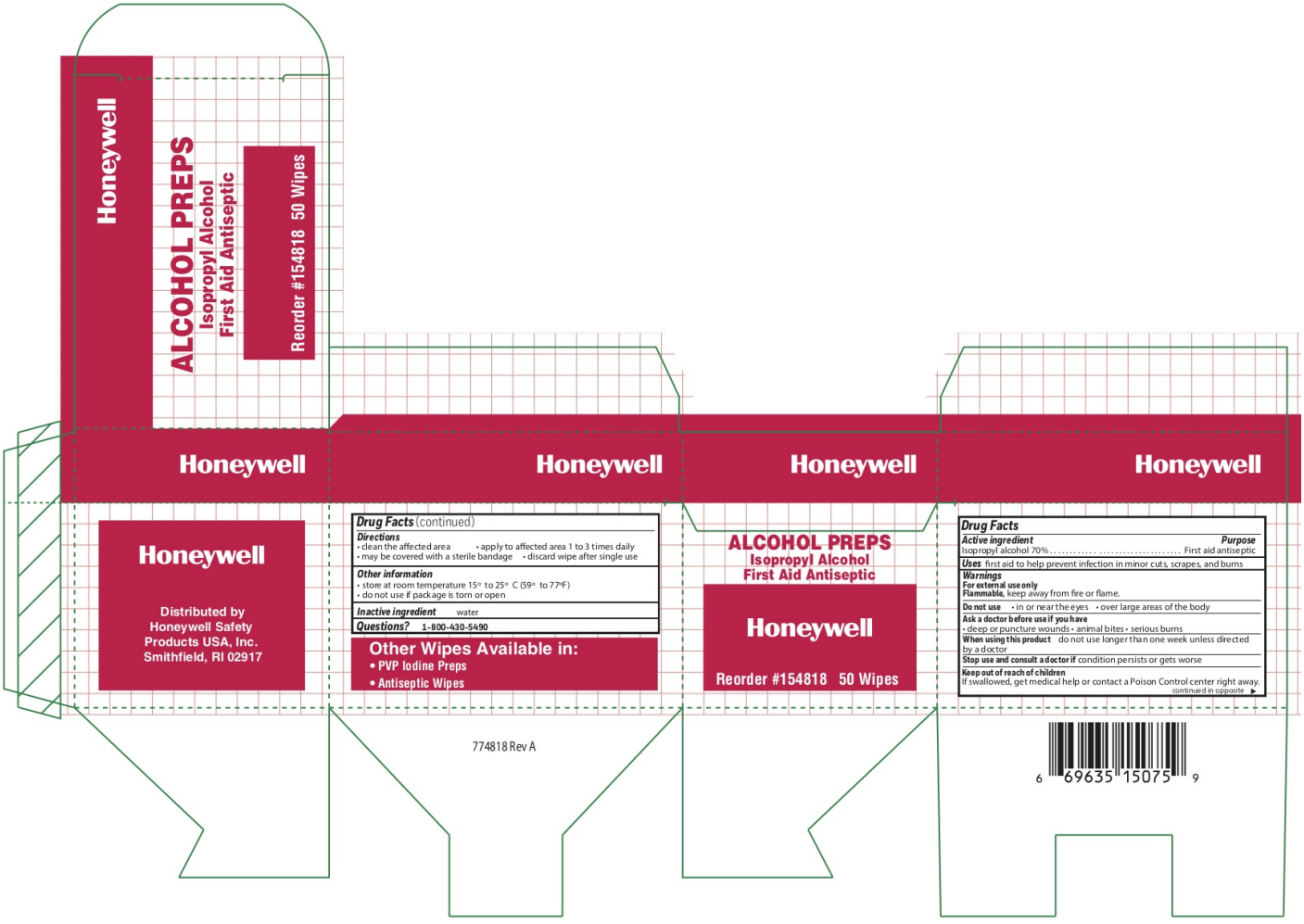
WATER-JEL®

INSTANT

Hand Sanitizer

Antiseptic Gel
With Vitamin E & Aloe

Alcohol Wipe
Principal Display Panel



4141 Kit Label

FAKREFU-B

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

4141 FIRST AID KIT

4141 first aid kit kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4141

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-4141-01	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	50 POUCH	20 mL
Part 3	25 PACKET	22.5 g
Part 4	25 PACKET	22.5 g
Part 5	10 PACKET	9 mL

Part 1 of 5

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 5

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:0498-0143-00	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		09/11/2018	

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

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: 059QF0KO0R)	
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**
unapproved drug other		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 (118768815)

Revised: 1/2024 Honeywell Safety Products USA, INC