4218 FIRST AID KIT- 4218 first aid 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.

0498-4218: First Aid Kit (neomycin, EW, Burn Jel, PVP Wipes, Sting Relief) 010424-4224

Active ingredient

Ammonia 15%

Purpose

Respiratory stimulant

Uses

to prevent or treat fainting

Warnings

For external use only

Do not use

• if you have 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.

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

Other information

store at room temperature away from light

Inactive ingredients

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

Questions or Comments

1-800-430-5490

Burn Jel Active ingredient

Lidocaine HCl 2.0%

Burn Jel *Purpose*

External analgesic

Burn Jel

Uses

temporarily relieves pain due to minor burns

Burn Jel *Warnings*

For external use only

Do not use

• on large areas of the body, particularly over raw surfaces or blistered areas

When using this product

avoid contact with eyes

Stop use and ask a doctor if

- the condition gets worse
- symptoms persist 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.

Burn JEI Directions

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

you may report a serious reaction to this product to 800-430-5490

Burn Jel

Other information

• store at room temperature - do not use if opened or torn

Burn Jel *Inactive ingredients*

carbopol 940, carbopol 1342, diazolidinyl urea, glycerin, melaleuca alternifolia (tea tree) leaf oil, methylparaben, octoxynol-9, propylene glycol, propylparaben, trolamine, water

Burn Jel *Questions*

1-800-430-5490

Sting Relief Active ingredient (in each wipe)

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

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 ingredient

benzalkonium chloride, menthol, and purified water

Sting Relief Questions or Comments?

1-800-430-5490

PVP

Active ingredient

Povidone-iodine solution USP, 10% (equivalent to 1% titratable iodine)

PVP Purpose

First aid antisepti

PVP

Uses

• first aid to help prevent the risk of infection in minor cuts, scrapes, and burn

PVP

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body
- on individuals who are allergic or sensitive to iodine

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 wek unless directed by a doctor

Stop use and ask a doctor if

- conditions persists or gets worse
- irritation and redness develops

Keep out of reach of children

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

PVP

Directioons

Reverse cardboard sleeve, then crush at dot between thumb and forefinger. Allow solution to saturate tip and apply solution to injury.

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

PVP

Other information

- store at room temperature away from light
- keep from freezing or excessive heat
- do not use if package is torn or open

PVP

Inactive ingredients

citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

PVP

Questions and Comments?

1-800-430-5490

Neomycin Active ingredient

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

Neomycin *Purpose*

First aid antiseptic

Neomycin *Uses*

First aid antibioticfirst aid to help prevent infection in

- minor cuts
- scrapes
- burns

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

- 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 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 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

4218 010424-4224 Kit Contents

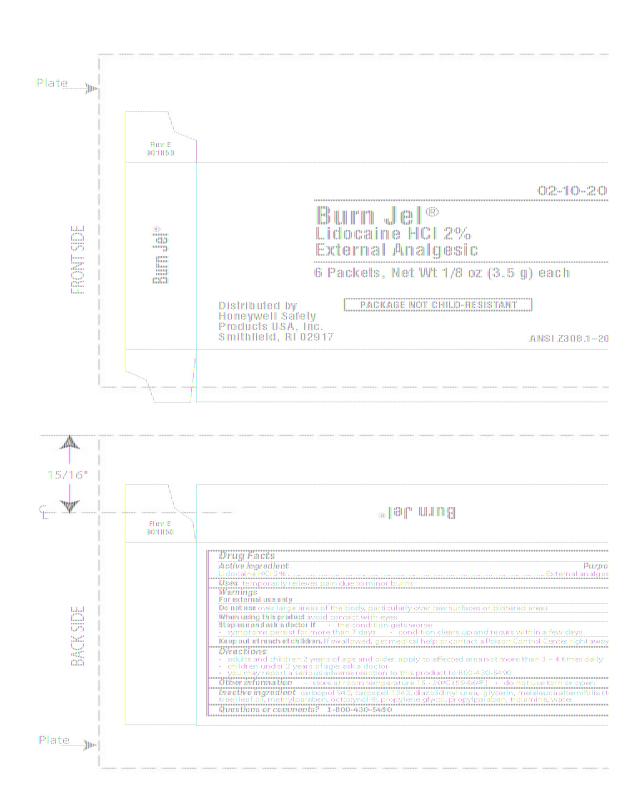
- 2 KNUCKLE BAND 8 PER
- 1 NEOMYCIN ANTIBIOTIC 10 PER
- 1 AMMONIA INHALANTS 10 PER
- 3 GAUZE BANDAGE, 4" X 6 YD
- 1 TOURNIQUET AND FORCEPS 1 EA
- 3 TRIANGULAR BDG, NON-STERILE
- 1 WIRE SPLINT 1 PER
- 1 ADH TAPE, .5" X 2.5 YD, 2 PER
- 3 GAUZE COMPRESS, 1728 SQ IN 1
- 2 GAUZE COMP, 1 SQ YARD, 1 PER
- 3 BANDAGE COMP, 2" OFFSET, 4 PER
- 2 BANDAGE COMP, 4" OFFSET, 1 PER
- 3 ADHESIVE BDG,PLSTIC,1"X3"16PER
- 1 FINGERTIP BANDAGE, 10 PER
- 1 1 OZ EYE WASH W/PADS & STRIPS
- 1 BURN JEL 1/8 OZ, 6 PER
- 1 PVP IODINE WIPES 10 PER
- 1 BIOHAZARD BAG/SCRAPER BBP
- 1 CPR MICROSHIELD W/2 PR LTX GLV
- 1 SCISSOR BDGE 4" RED PLS HDL
- LBL STOCK 6-3/8"X4"
- 1 LBL STOCK 3"x1-7/8"
- 1 LBL CONTS 8"X8", CUSTOM ID B

- 1 KIT STL 36 UN WHT 01 HOR SHELF
- 1 LABL INSTR 24 & 36 UNIT KITS
- 1 STING RELIEF 10

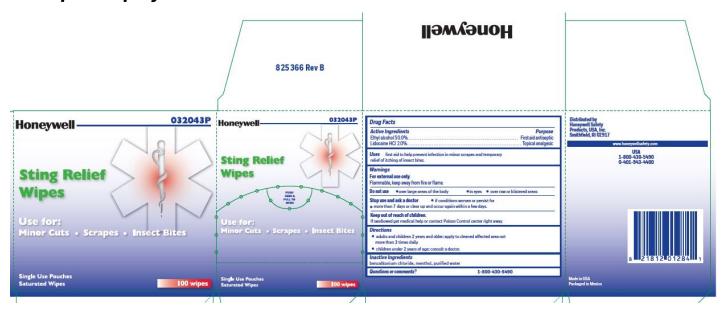
Principal Display Panel

796006 Rev. E (page 3 of 3)

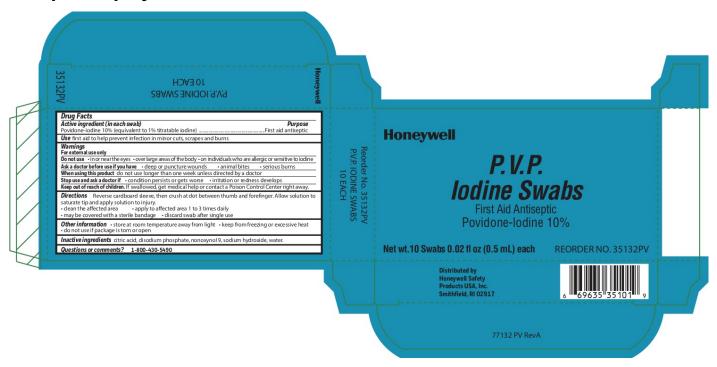
796353 Rev. E Unit Carton Printing Plate for "B" size cartor



Sting Relief Principal Display Panel

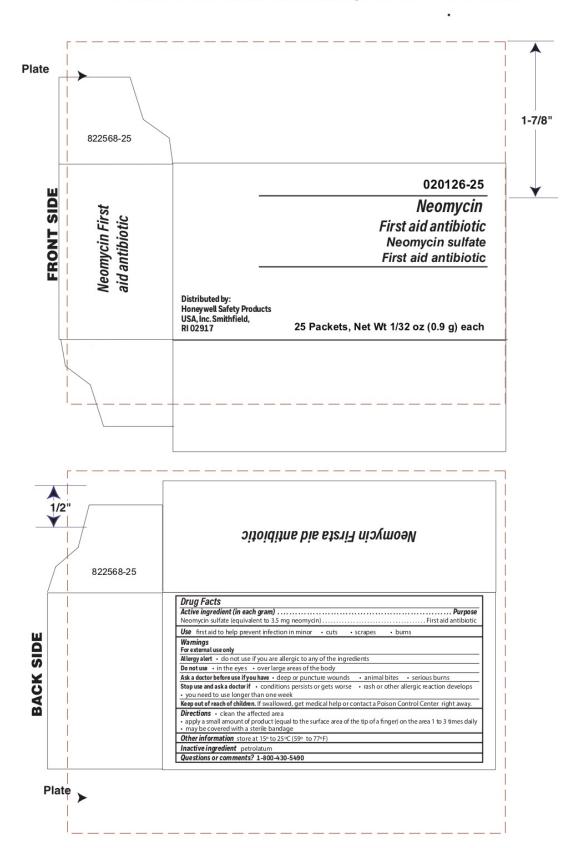


PVP Principal Display Panel



Neomycin Principal Display Panel

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



46001404 Rev. C prints 2 colors black and red (pms 485)

Refill Information

US Poison Control 1-800-222-1222

46001404 Rev. C

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

4218 FIRST AID KIT

4218 first aid kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4218

Packaging

Quantity of Darto

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0498-4218-01	1 in 1 KIT	09/13/2018	

Quant	quantity of Parts		
Part #	Package Quantity	Total Product Quantity	
Part 1	10 AMPULE	3 mL	
Part 2	6 PACKET	21 g	
Part 3	10 POUCH	4 mL	
Part 4	10 POUCH	3 mL	
Part 5	1 BOTTLE	30 mL	

9 g

Part 1 of 6

Part 6 10 PACKET

AMMONIA INHALENT

ammonia inhalent inhalant

Product Information

tem Code	(Source)	NDC:0498-3334

Route of Administration RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety Ingredient Name

- 1			
ı	AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

Basis of Strength

Strength

Inactive Ingredients

Strength

ALCOHOL (UNII: 3K9958V90M)

	ka		

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

Marketing In	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		09/18/2018	12/01/2022		

Part 2 of 6

BURN JEL

gel for burns gel

Product Information

Item Code (Source) NDC:0498-0203

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
TEA TREE OIL (UNII: VIF565UC2G)		
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
GLYCERIN (UNII: PDC6A3C0OX)		
TROLAMINE (UNII: 903K93S3TK)		
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)		
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
OCTOXYNOL-9 (UNII: 7JPC6Y25QS)		
DIPROPYLENE GLYCOL (UNII: E107L85C40)		

1	Pa	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	

		3.5 g in 1 PACKET; Type 0: Not a Combination
-	00	Product

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

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	NDC:0498-0733- 00	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing In	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug		12/22/2017			

Part 4 of 6

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

Item Code (Source) NDC:0498-0121

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Strength Ingredient Name

POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4) IODINE 10 mg in 1 mL

Inactive Ingredients

Ingredient Name Strength

NONOXYNOL-9 (UNII: 48Q180SH9T)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0498-0121- 00	0.3 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

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

Part 5 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		
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)		
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		

l	P	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 6 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:116QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g		

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

ı	P	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	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		09/13/2018		

Labeler - Honeywell Safety Products USA, Inc. (118768815)

Revised: 1/2024 Honeywell Safety Products USA, Inc.