

**4239 FIRST AID KIT- 4239 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.

0498-4239: First Aid Kit (Triple, EW, Burn Jel, PVP Ampule, BZK wipe, alcohol wipe, sting relief- 013111-4536)

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 Jel
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

Triple

Active ingredient

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 ° to 25 ° C (59 ° to 77 ° F)
- tamper evident sealed packets
- do not use if packet is torn or opened

Triple**Inactive ingredient**

petrolatum

Triple**Questions?**

1-800-430-5490

BZK Wipe

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK Wipe

Purpose

First aid antiseptic

BzK Wipe

Uses

Antiseptic cleansing of face, hands, and body without soap and water

BZK Wipe

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 Wipe

Directions

tear open packet and use as a washcloth

BZK Wipe

Other information

- store at room temperature 15⁰ to 30⁰ C (59⁰ - 86⁰ F)
- do not reuse towelette

BZK Wipe
Inactive ingredient

water

BZK Wipe
Questions

1-800-430-5490

Sting Relief
Active ingredient (in each wipe)

Ethyl alcohol 50.0%

Lidocaine HCl 2.0%

Sting Relief
Purposes

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

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 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

Directions

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

Alcohol

Other information

- store at room temperature 15⁰ to 25⁰ C (59⁰ to 77⁰ F)
- do not use if packet is torn or

Alcohol

Questions

1-800-430-5490

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.

Eyeash *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

Eyeash *Questions*

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

PVP Wipe *Active ingredient*

Povidone-iodine 10% (equivalent to 1% titratable iodine)

PVP Wipe

Purpose

First aid antiseptic

PVP Wipe

Uses

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

PVP Wipe

Warnings

For external use only.

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 or persists for more than 72 hours
- irritation and redness develops

Keep out of reach of children.

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

PVP Wipe

Directions

- clean the affected area
- apply 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first
- discard wipe after single use

PVP Wipe

Other information

- do not use on individuals who are allergic or sensitive to iodine
- store at controlled temperature 59-86°F (15-30°C)
- do not use if pouch is open or torn

PVP Wipe

Inactive ingredients

nonoxynol 9, water

**PVP Wipe
Questions**

1-800-430-5490

4239

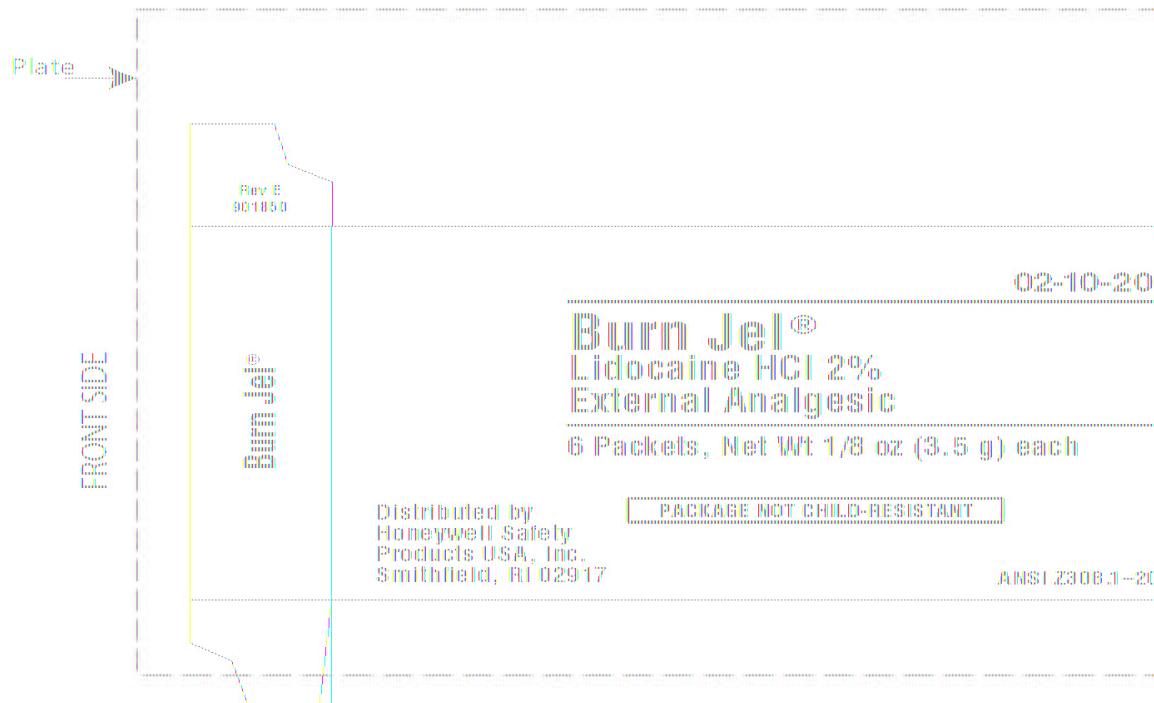
013111-4536 kit contents

1 KNUCKLE BAND 8 PER
1 TRIPLE ANTIBIOTIC 10 PER
1 EYE DRESS PKT W/4 ADH STRIPS
1 TRIANGULAR BDG, NON-STERILE
1 ADH TAPE, .5" X 2.5 YD, 2 PER
1 GAUZE PADS, 4" X 4", 2 PER
1 GAUZE COMP, 1 SQ YARD, 1 PER
1 BUFFERED EYE WASH 1 OZ BTL
1 BANDAGE COMP, 2" OFFSET, 4 PER
1 BANDAGE COMP, 4" OFFSET, 1 PER
1 ADHESIVE BDG,PLSTIC,1"X3"16PER
1 ADH BAND, EXTRA LARGE, 6 PER
1 BURN JEL 1/8 OZ, 6 PER
1 WATER JEL DRESSING 4" X 4"
1 PVP IODINE WIPES 10 PER
1 BIOHAZARD BAG,WIPES/TOWELS BBP
1 FACE MASK W/SHIELD 1 PER BBP
1 NITRILE GLOVES 2PR BBP
1 FIRST AID GUIDE ASHI
1 MICROSHIELD W/VNL GLV/ALCL
LBL STOCK 6-3/8"X4"
LBL STOCK 4"X2-7/8"
1 LBL STOCK 3"x1-7/8"
1 LABEL NORTH CONTENTS 8X8 ID B
1 KIT PP 24 UNIT FA
1 STING Relief WIPES 10

Burn Jel

Principal Display Panel

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



Triple Principal Display Panel



BZK Wipe Principal Display Panel

Antiseptic Towelettes

Honeywell

02-16-35MD

Antiseptic Towelettes

Benzalkonium chloride
First aid antiseptic

Six-Saturated Towelettes

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

Antiseptic Towelettes

Honeywell

Drug Facts

Active Ingredient

Benzalkonium chloride 0.133% w/v **Purpose** First aid antiseptic

Uses

• antiseptic cleaning of face, hands and body without soap and water. • air dries in seconds

Warnings

For external use only

When using this product • do not use in the eyes or apply over large areas of the body

Ask a doctor before use • in case of deep or puncture wounds, animal bites, or serious burns

Stop use and consult a doctor if

• irritation, redness or other symptoms develop • condition persists or gets worse

Do not use • longer than 1 week unless directed by doctor

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

Directions • tear open packet, unfold and use as washcloth

Other Information

• store at room temperature 15° -30° C (59° -86° F) • do not reuse towelette

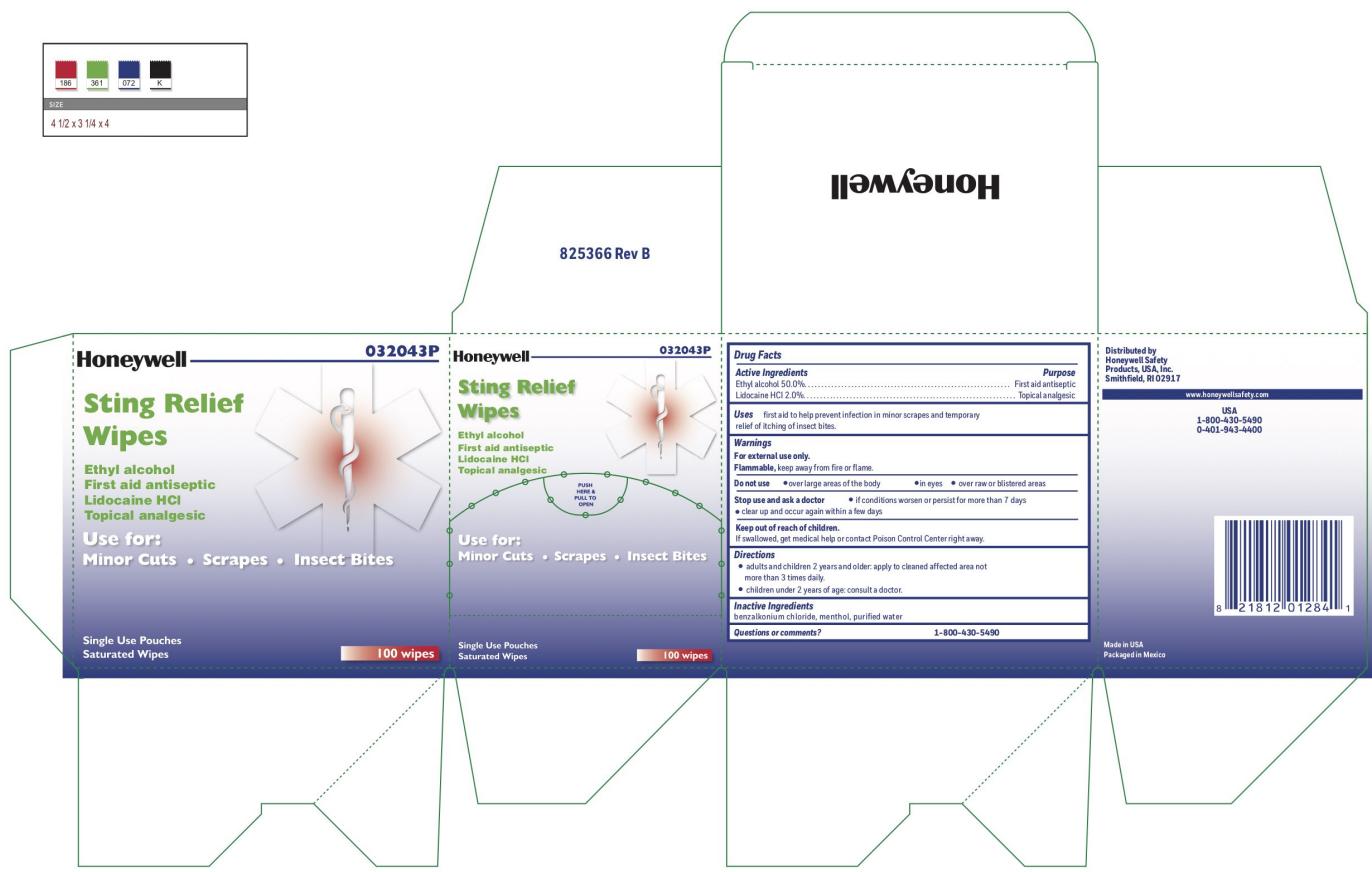
Inactive ingredient

water

Questions or comments

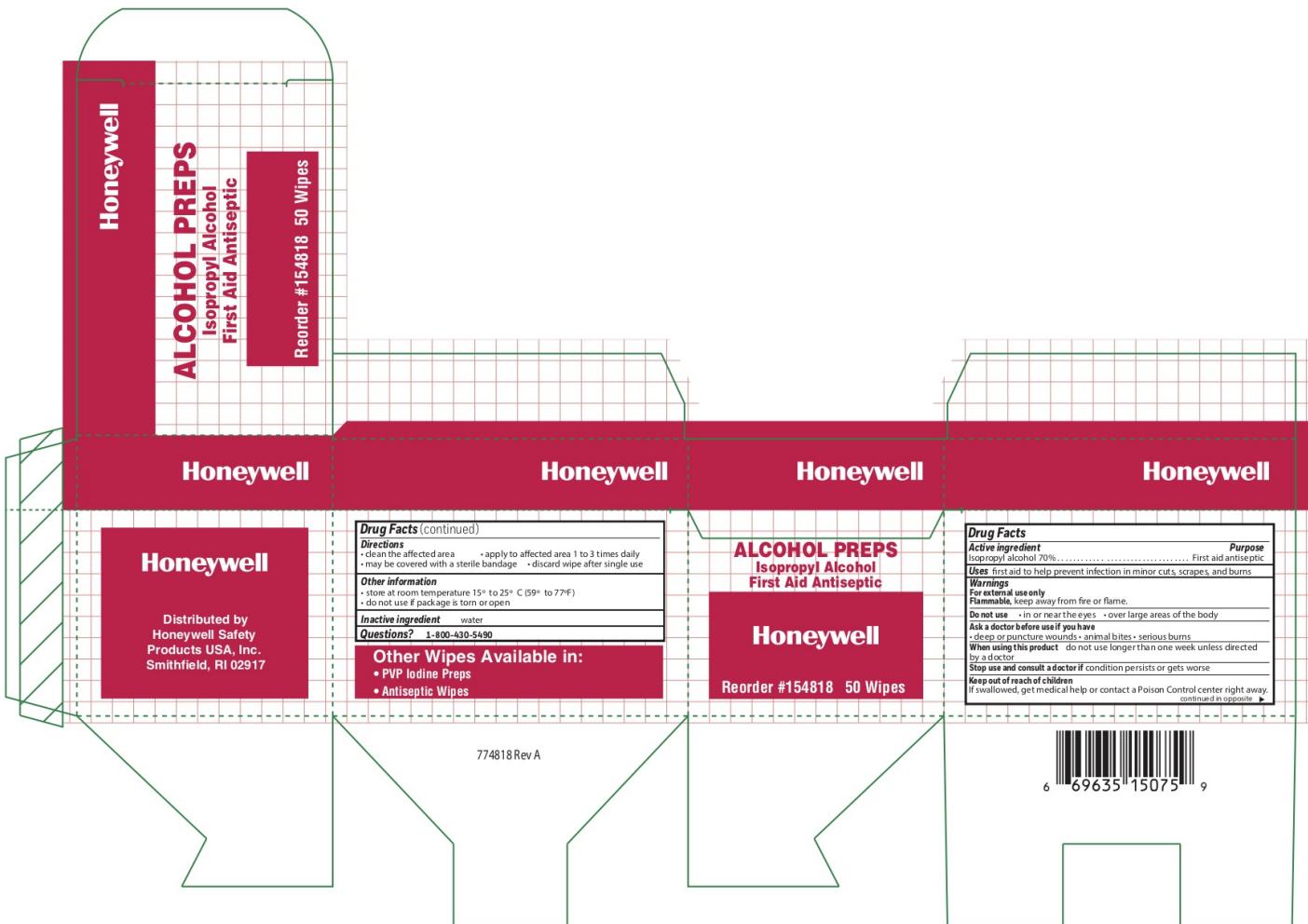
1-800-430-5490

Sting Relief
Principal Display Panel



Alcohol

Principal Display Panel



Eyewash Principal Display Panel

HoneywellTAMPER-EVIDENT CAP.
TAPA CON SELLO DE SEGURIDAD.
BOUCHON INDICATEUR D'INFRACTION.**eyesaline®****LAVAOJOS
EYESALINE****EYESALINE
EYEWASH****LAVAGE
OCULAIRE
EYESALINE**Solución
Isotónico EstérilSterile
Isotonic SolutionLa Solution
Isotonique Stérile**16 fl. oz. (473 mL)****Drug Facts (for USA only)****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 consult 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.**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

LABEL #22-204510 Rev. J
REORDER / NUEVO / PEDIDO / REAPROVISIONAMIENTO #22-200945-0000

space for lot code and supplier part number

PEEL / PELAR / PEELER

Datos de medicamento (Para EE.UU. solamente)**Ingrediente Activo** Agua estéril 99%**Propósito** Lavados**Usos**

para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aéros. o agua de cloruro

Advertencias

Para uso externo sólo - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con ninguna superficie. No vuelva a usar. Vez abierto, descarte.

No se usa

- si tiene heridas abiertas en o cerca del ojo, obtenga ayuda médica de inmediato.

Deje de usar y consulte a un médico si:

- experimenta dolor de ojo
- cambio de visión
- rojez continuo o irritación del ojo
- la condición empeora o persiste

Manténgase fuera del alcance de los niños.

En caso de ingestión accidental, obtenga atención médica o llame a un Centro de control de envenenamiento inmediatamente.

Instrucciones:

- quite los lentes de contacto antes de usar la solución
- tuerza la tapa para quitar
- enjuague el área afectada según se necesite
- controle el chorro haciendo presión en la botella
- si es necesario, sigue enjuagando con un lavajos o ducha de emergencia

Ingredientes inactivos

cloruro de sodio, fosfato de sodio dibásico, fosfato de sodio monobásico

¿Preguntas? Llame al 1-800-430-5490

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

Information**Usages**

Pour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmosphériques ou de l'eau chlorée.

Advertissements

Pour usage externe seulement - Obtenir immédiatement des soins médicaux pour toutes les plaies ouvertes dans ou près des yeux. Pour éviter toute contamination, ne pas toucher la pointe du récipient à n'importe quelle surface. Ne pas réutiliser. Une fois ouvert, jetez-les.

Ne pas utiliser

- si la solución a cambiado de color o si ella es brouillée
- si vous avez des plaies ouvertes aux yeux ou à proximité, consultez immédiatement un médecin

Cessez d'utiliser la solution et consulter un médecin

- vous ressentez une douleur oculaire
- si votre vision change
- rougeur ou irritation persistante des yeux
- condition empire ou persiste

Garder hors de la portée des enfants.

En cas d'ingestion, communiquer immédiatement avec un médecin ou avec un centre antipoison.

Mode d'emploi

- enlever los lentes de contacto avant l'utilisation
- dévisser le bouchon pour l'enlever
- rincer la zone touchée selon les besoins
- ajuster le débit d'écoulement de la solution en augmentant ou en réduisant la pression exercée sur le contenant
- si nécessaire, continuer de rincer avec unesolution de rinçage oculaire d'urgence ou una douche

Ingredíents agua estéril, cloruro de sodio, fosfato dibásico de sodio, fosfato monobásico de sodio**Des questions?** Faites le 1-800-430-5490

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

**PVP Wipe
Principal Display Panel**

822569 X
Rev. ***PVP Iodine Wipes****02-12-01X****PVP Iodine Wipes**

*Povidone-Iodine 10%
First Aid Antiseptic
10 Saturated Wipes
ANSI Z308.1-2009*

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

822569 X
Rev. ***PVP Iodine Wipes****Drug Facts****Active ingredient**

Povidone-Iodine 10% (equivalent to 1% titrable iodine).....

Purpose

First aid antiseptic

Use

first aid to help prevent the risk of 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 • 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 week unless directed by a doctor**Stop use and ask a doctor if** • condition persists or gets worse • irritation or redness develops**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away**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

Other information • store at room temperature: 15-30° C (59-86° F)

• do not use if package is torn or open • do not use on individuals who are allergic or sensitive to iodine

Inactive ingredients nonoxynol-9, water**Questions or comments?** 1-800-430-5490

4239 Kit Label
013111-4536

46001365 Rev. C
Prints 3 colors
Black, Red (PMS 186) and Blue (PMS 072)

Refill Information

US Poison Control 1-800-222-1222



Contact your authorized Honeywell Safety Products
Distributor with your refill orders.

Honeywell

www.honeywellsafety.com

USA
1-800-430-5490

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

46001365 Rev. C

4239 first aid kit

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:0498-4239

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4239-01	1 in 1 KIT	09/13/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	21 g
Part 2	10 POUCH	3 mL
Part 3	1 BOTTLE	30 mL
Part 4	10 PACKET	9 g
Part 5	2 PACKET	2.8 mL
Part 6	4 POUCH	1.6 mL
Part 7	10 POUCH	4 mL

Part 1 of 7

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)	
EDETA TE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C00X)	

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)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0203-00	3.5 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 2 of 7

PVP IODINE WIPE

povidone-iodine 10% swab

Product Information

Item Code (Source)	NDC:0498-0121		
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
NONOXYNOL-9 (UNII: 48Q180SH9T)	
WATER (UNII: 059QF0KOOR)	

Packaging

	Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0121-00	0.3 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

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: 059QF0KOOR) (WATER - UNII:059QF0KOOR)	WATER	98.6 mL in 100 mL

Inactive Ingredients

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

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 4 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:J2VZ 07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	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:0498-0750-35	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		09/19/2018	

Part 5 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: 059QF0KOOR)	

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	

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

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:98P1200987)	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: 059QF0KOOR)

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 Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/23/2017	

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.