

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 Safety 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 ° to 25 ° C (59 ° to 77 °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 ° to 25 ° C (59 ° to 77 ° 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 °-30 ° C (59 ° -86 ° 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 °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 ° to 25 ° C (59 ° to 77 ° 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

Honeywell

TAMPER-EVIDENT CAP.
TAPA CON SELLO DE SEGURIDAD.
BOUCHON INDICATEUR D'INFRACTION.

eyesaline®

LAVAOJOS
EYESALINE

Solución
Isotónico Estéril

EYESALINE
EYEWASH

Sterile
Isotonic Solution

LAVAGE
OCULAIRE
EYESALINE

La Solution
Isotonique Stérile

16 fl. oz. (473 mL)

NPN: 80057528



Drug Facts (for USA only)

Active ingredient	Purpose
Sterile water 99%	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 432-004510 Rev. J REORDER / NUEVO PEDIDO / RÉAPPROVISIONNEMENT 032-000454-0000

space for lot code and supplier part number

PEEL / PELAR / PELER

Datos de medicamento (Para EE.UU. solamente)

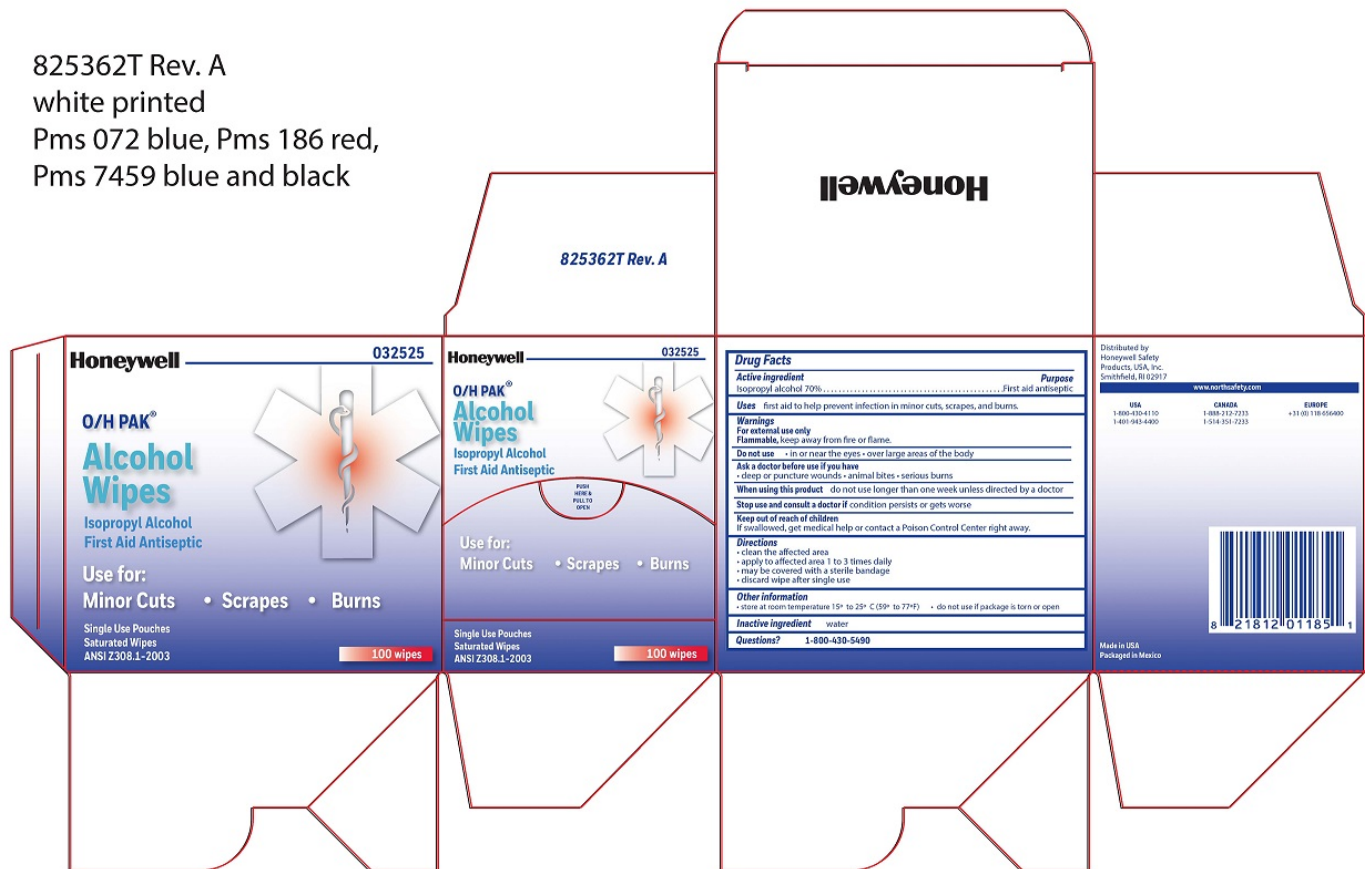
Ingrediente Activo	Propósito
Agua estéril 99%	Lavaojos
Usos para el lavado de ojo para quitar las partículas sueltas y extrañas, los contaminantes aeros, o agua de cloruro.	
Advertencias Para el uso externo sólo - Obtenga tratamiento médico inmediato para todas las heridas abiertas en o cerca de los ojos. Para evitar la contaminación, no toque la punta del envase con ninguna superficie. No vuelva a usar. Vez abierto, descarte. No se use • si la solución se enturbia o cambia de color • 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 • quítense 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 enjuagado con un lavaojos 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 solution a changé de couleur ou si elle est brouillée • si vous avez des plaies ouvertes aux yeux ou à proximité, consultez immédiatement un médecin Cesser 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 les verres de contact 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 une solution de rinçage oculaire d'urgence ou une douche
Ingédients eau stérile, chlorure de sodium, phosphate dibasique de sodium, phosphate monobasique de sodium
Des questions? Faites le 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

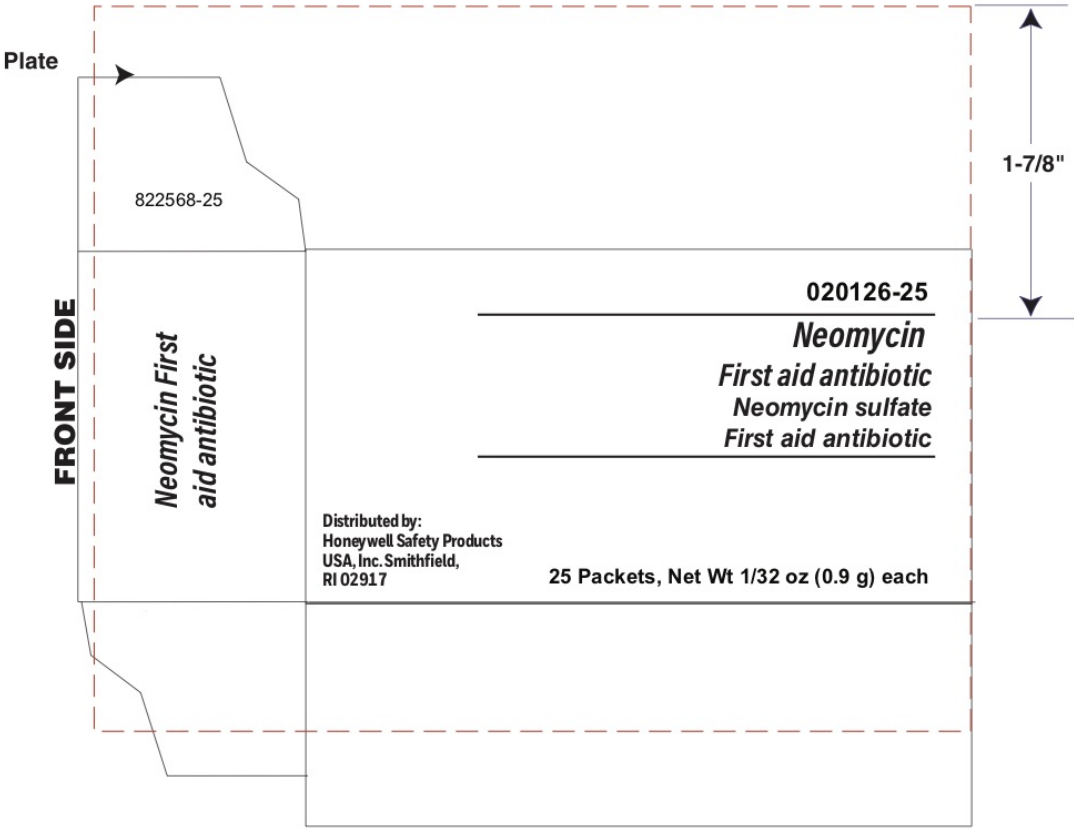
Alcohol Wipe
Principal Display Panel

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.



Principal Display Panel

Honeywell ANTISEPTIC SPRAY <i>Benzalkonium chloride</i> First Aid Antiseptic Net contents 2 fl oz (59 mL) ANSI Z308.1-2003  8 21812 01498 2	032203 																					
	Drug Facts <table><tr><td>Active ingredient Benzalkonium chloride 0.13%</td><td>Purpose First aid antiseptic</td></tr><tr><td colspan="2">Uses first aid to help prevent infection in minor cuts, scrapes and burns</td></tr><tr><td colspan="2">Warnings For external use only Do not use • in or near the eyes • over large areas of the body</td></tr><tr><td colspan="2">Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns</td></tr><tr><td colspan="2">When using this product do not use longer than one week unless directed by a doctor</td></tr><tr><td colspan="2">Stop use and ask a doctor if condition persists or gets worse</td></tr><tr><td colspan="2">Keep out of reach of children. If swallowed get medical help or contact a Poison Control Center right away.</td></tr><tr><td colspan="2">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</td></tr><tr><td colspan="2">Other information • shake well • store at room temperature, 15° to 30°C (59° to 86°F)</td></tr><tr><td colspan="2">Inactive ingredients diazolidinyl urea, edetate disodium, glycerin, hypromellose, methylparaben, octoxynol 9, propylene glycol, propylparaben, triethylamine, water</td></tr><tr><td colspan="2">Questions or comments? 1-800-430-5490</td></tr></table>	Active ingredient Benzalkonium chloride 0.13%	Purpose 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, triethylamine, water		Questions or comments? 1-800-430-5490
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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.)	Cat. No. 201005																			
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47.0306	 6 69635 20032 4 Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917 Honeywell																			

Hand sanitizer
Principal Display Panel



INSTANT

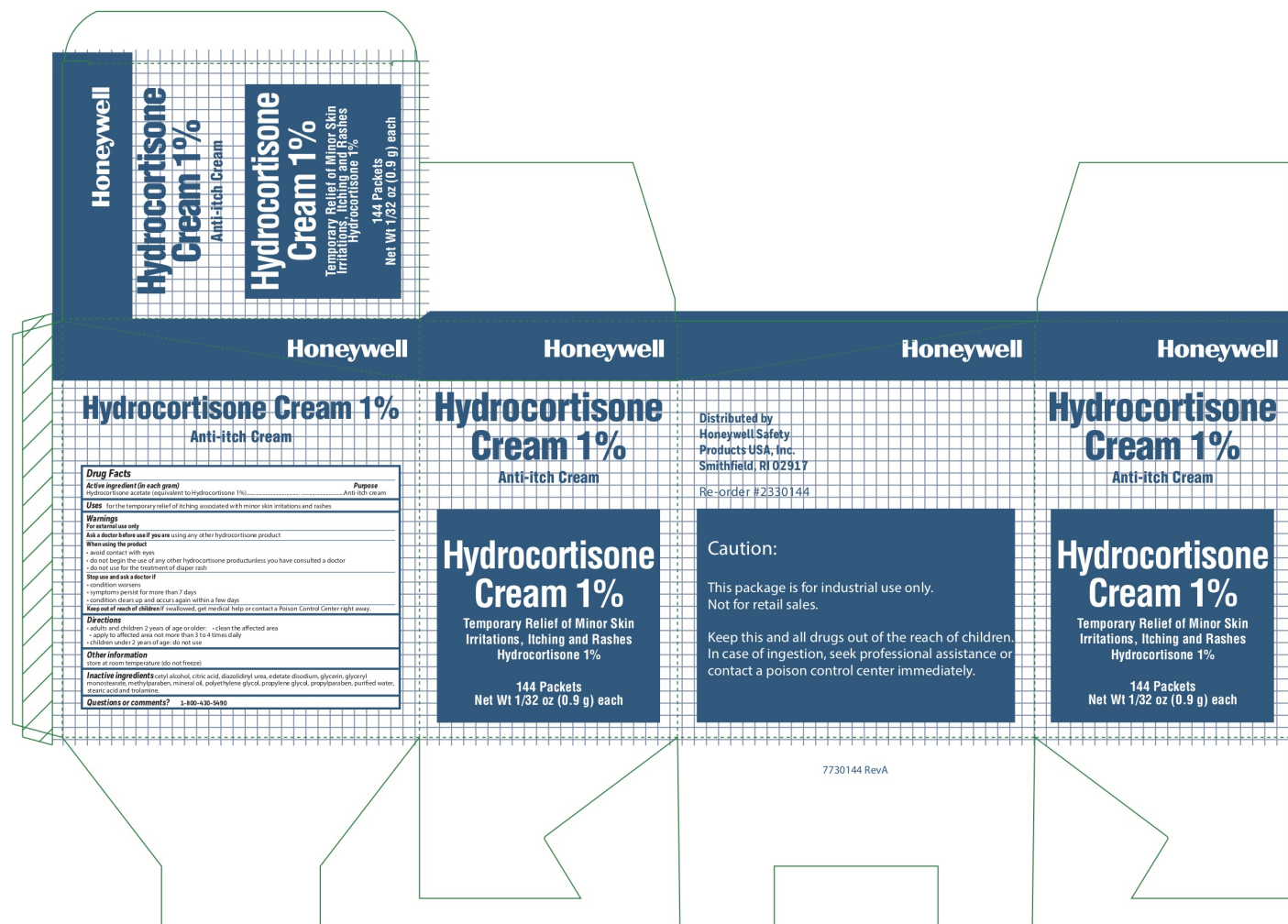
Hand Sanitizer

Antiseptic Gel
With Vitamin E & Aloe

Kills 99.9% of Germs Without Water

240mL - (8 fl oz)

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

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, 1.0%, 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
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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
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1	NDC:0498-0143-04	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		
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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: 3WJQ0SDW1A)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

		Marketing Start	Marketing End
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#	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: V13007Z 41A) (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		
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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	

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