4149 FIRST AID KIT- 4149 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.

0498-4149: First Aid Kit (alcohol wipes, HC cr, PVP wipes, sting relief, EW-SF00003660)

Sting Relief Active ingredient (in each wipe)

Ethyl alcohol 50.0%

Lidocaine HCl 2.0%

Sting Relief Purposse

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

Eyewash Active ingredient

Sterile Water 99%

Eyewassh *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.

Eyewash 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

Hydrocortisone Active ingredient (in each gram)

Hydrocortisone acetate (equivalent to Hydrocortisone 1%)

Hydrcortisone *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

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

Alcohol Active ingredient

Isopropyl alcohol 70%

Alcohol *Purpose*

First aid antiseptic

Alcohol Uses

first aid to help prevent infection in

- minor cuts
- scrapes
- 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 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 affeted are 1 to 3 times daily

• discard wipe after single use

Alcohol

Other information

- store at room temperature 15 0 to 25 0 C (59 0 to 77 0 F)
- do not use if packet is torn or opened

Alcohol Inactive ingredient

water

Alcohol Questions

1-800-430-5490

PVP

Active ingredient

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

PVP Purpose

First aid antiseptic

PVP

Uses

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

PVP

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

Directions

- clean the affected area
- apply1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first
- discard wipe after single use

PVP

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

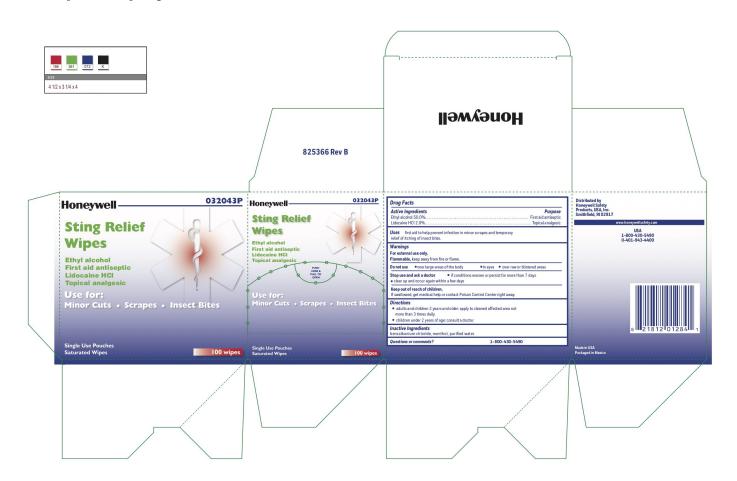
Inactive ingredients nonoxynol 9, water

4149 SF00003660 Kit Contents

- 1 KNUCKLE BAND 8 PER
- 2 TRIANGULAR BDG, NON-STERILE
- 1 ADH TAPE, .5" X 2.5 YD, 2 PER
- 1 FORCEPS & SCISSORS, 1 EA
- 2 GAUZE BANDAGE, 2" X 6 YD,2 PER
- 1 GAUZE COMP, 1 SQ YARD, 1 PER
- 1 INSTANT COLD PACK 4" X 6"
- 1 BANDAGE COMP, 2" OFFSET, 4 PER
- 1 BANDAGE COMP, 4" OFFSET, 1 PER
- 1 FINGERTIP BANDAGE, 10 PER
- 1 1 OZ EYE WASH W/PADS & STRIPS
- 2 WATER JEL DRESSING, 2" X 2"
- 2 WATER JEL DRESSING, 2" X 6"
- 1 ALCOHOL PREP PADS 10P

- 1 HYDROCORTISON, 1.0%, 1/32 OZ, 10P
- 3 PVP IODINE WIPES 10 PER
- 1 IVYX CLEANSER TOWEL 5 PER
- 1 NITRILE GLOVES 2PR BBP
- 2 ADH BDG, CLOTH, 1"X3", 16 PER
- 1 CPR FILTERSHIELD 77-100
- 1 40Z BFS EYEWASH TRILINGUAL BOTTLE
- LBL STOCK 6-3/8"X4"
- LBL STOCK 4"X2-7/8"
- 1 LBL STOCK 3"x1-7/8"
- 1 KIT 36 UNIT PLASTIC
- 1 LBL 36U CVR NORTH ID C.
- 1 STING Relief WIPES 10

Sting Relief Principal Display Panel



Eyewash Principal Display Panel

Isotónico Estéril



Isotonic Solution

16 fl. oz. (473 mL)

Isotonique Stérile

#32-000454-0000 Drug Facts (for USA only) Active ingredient Purpose for flushing the eye to remove loose foreign material, air pollutants, RÉAPPROVISIONNEMENT 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 PEDIDO / Stop use and consult a doctor if: Stop use and consult a doctor in:
you experience eye pain of hanges in vision
condition worsens or praists of the eye
condition worsens or persists

Keep out of reach of children.
Fixed the worse wo REORDER 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 #32-004510 Rev. J sodium phosphate dibasic, sodium phosphate monobasi

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

PEEL / PELAR / PELER

Datos de medicamento (Para EE.UU. solamente) Ingrediente Activo Agua estéril 99% Usos para el lavado de ojo para quitar las particulas sueltas y extrañas, los contaminantes aeros, o agua de cloruro

LABEL

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
inguna 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 medica de inmediato

de inmediato
Deje de usar y consulte a un médico si:

• experimenta dolor de ojo
• cambio de visión
• rojez continuo o rintación 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 |
- quitese los lentes de contacto antes de usar la solución |
- tuerza la tapa para quitar |
- enjuaque el área afectada según se necesite |
- controle el chorro haciendo presión el 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

UsagesPour le rinçage des yeux afin d'enlever un corps étranger, des polluants atmospheriques où de l'eau chlorée.

Advertissements

Pour usage externs 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, jetze-jes.

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

• ou votre vision change

• ougeur ou infallon 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.

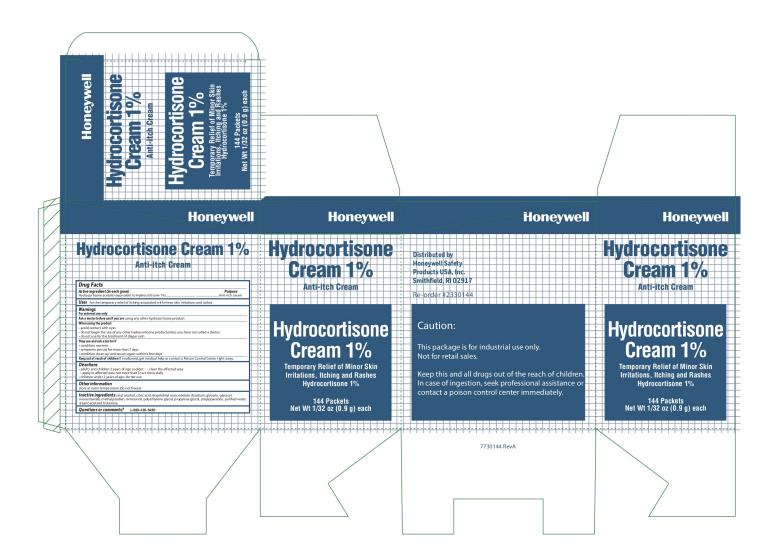
Mode d'emploi

• enlever les verres de contact avant l'utilisation • dévisser le bouchon pour l'enlever • nincer la zone touchée selon les besons • ajuster le débit d'écoulement de la solution en augmentant ou en rédulsant la pression exercée sur le contenant culture de la coultion en contract le contenant en coulaire d'urgence ou une douche

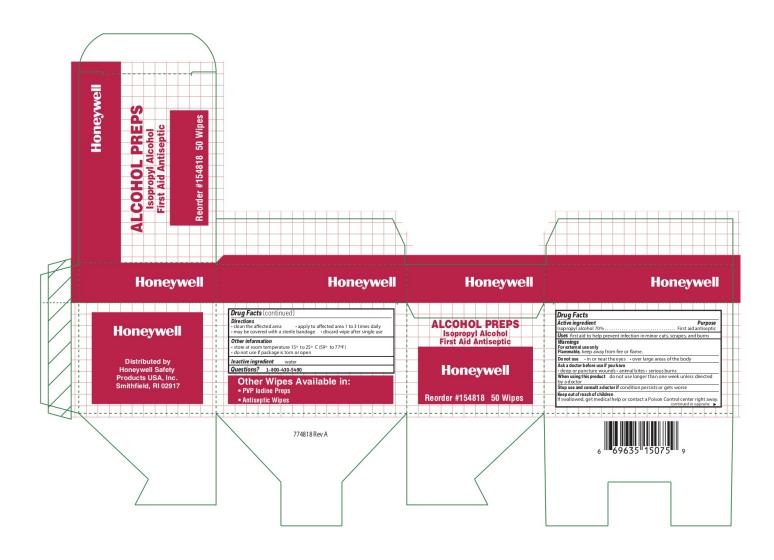
Ingrédients eau stérile, chlorure de sodium, phosphate dibasique de sodium, phosphate monobasique de sodium

Des questions? Faites le 1-800-430-5490 Honeywell Safety Products USA, Inc. Smithfield, RI. 02917

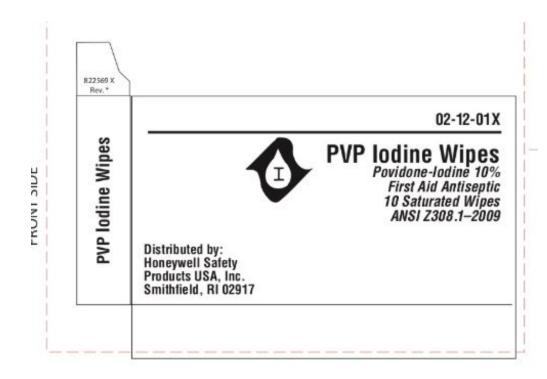
Hydrocortisone Principal Display Panel

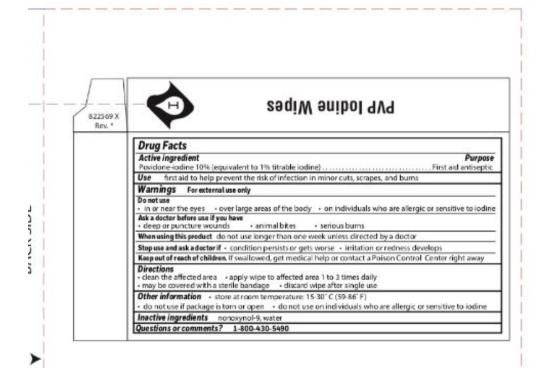


Alcohol Principal Display Panel

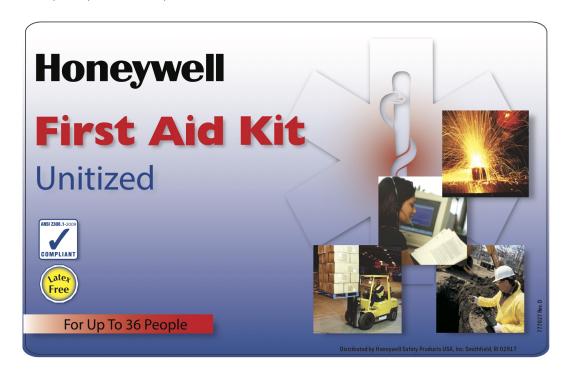


PVP Principal Display Panel





4149 Kit Label SF00003660



4149 FIRST AID KIT

4149 first aid kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4149

Packaging

#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4149-01	1 in 1 KIT	09/13/2018	

Ouantity of Parts

Quant	eductive of a dies		
Part #	Package Quantity	Total Product Quantity	
Part 1	1 BOTTLE	30 mL	
Part 2	10 POUCH	4 mL	
Part 3	10 POUCH	4 mL	
Part 4	10 PACKET	9 g	
Part 5	30 POUCH	9 mL	
Part 6	1 BOTTLE	118 mL	
Part 7	10 PACKET	9 g	

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 PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:0498-0100-	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M018	12/18/2018	

Part 2 of 7

ALCOHOL WIPE

isopropyl alcohol swab

B		
Droduct	Intorm	ation

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)	

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:0498-0143- 04	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Mark	Marketing Information			
	keting egory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapprov other	ed drug		09/18/2018	

Part 3 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:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL	

Inactive Ingredients		
Ingre	dient Name	Strength

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:0498-0733-	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	

Part 4 of 7

HYDROCORTISONE

anti-itch cream ointment

Product Information

Item Code (Source) NDC:0498-0800

Route of Administration TOPICAL

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

Inactive Ingredients	
Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
TROLAMINE (UNII: 903K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

PROPYLPARABEN (UNII: Z8IX2SC10H)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

l	P	ackaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:0498-0800- 34	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
unapproved drug other		03/06/2013	

Part 5 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		
WATER (UNII: 059QF0KO0R)			
NONOXYNOL-9 (UNII: 48Q180SH9T)			

	Packaging				
#	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 6 of 7

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)NDC:0498-0100Route of AdministrationOPHTHALMIC

Active Ingredient/Active Moiety Ingredient Name Basis of Strength WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER (UNII: 059QF0KO0R) (WATER - UNII: 059QF0KO0R)

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	
	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 7 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 HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE UNII: W4X0X7BPJ) HYDROCORTISONE ACETATE 1 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
LIGHT MINERAL OIL (UNII: N6K5787QVP)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)		
TROLAMINE (UNII: 903K93S3TK)		
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)		
PROPYLPARABEN (UNII: Z8IX2SC10H)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		

l	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		03/06/2013		

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.