4220 FIRST AID KIT- 4220 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.

0498-4220: First Aid Kit (Neomycin, EW, Burn Jel, PVP Wipes, Sting Relief, alcohol) 019643-4367

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 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 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

Alcohol Active ingredient

Alcohol *Purpose*

First aid antiseptic

Alcohol *Uses*

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

Keep out of the reach of children

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

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

 $\bullet\,$ do not use longer than 1 week unless directed by a doctor

Stop use and consult a doctor if

• condition persists or gets worse

Alcohol Directioons

- 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 o to 25 o C (59 o to 77 oF)
- do not use if packet is torn or opened

Alcohol Inactive ingredient

water

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

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

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.

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

Eyewash *Questions?*

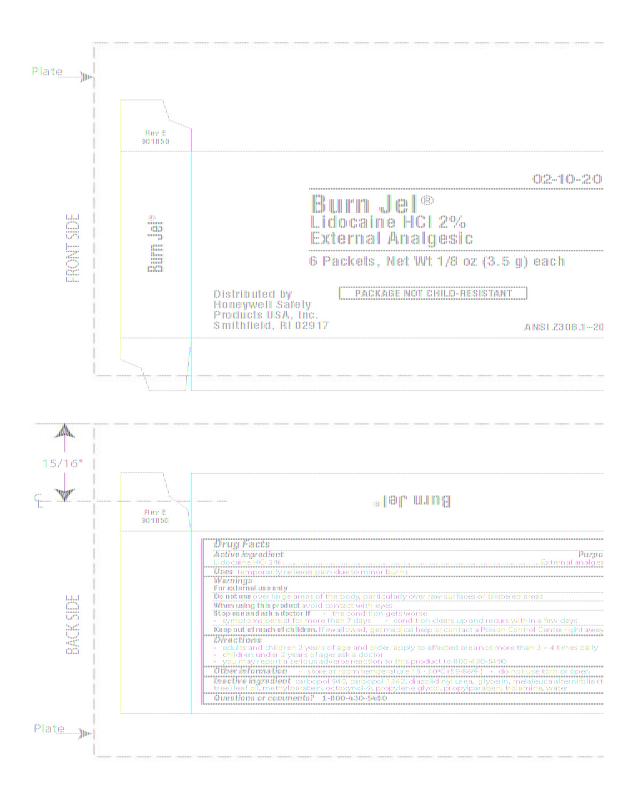
1-800-430-5490

4220 019643-4367 Kit Contents

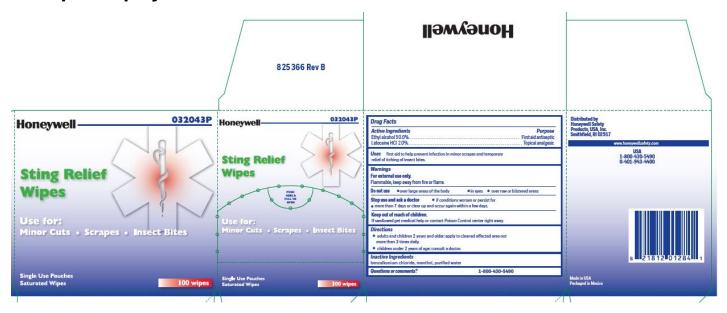
- 1 KNUCKLE BAND 8 PER
- 1 NEOMYCIN ANTIBIOTIC 10 PER
- 1 GAUZE BANDAGE, 4" X 6 YD
- 2 TRIANGULAR BDG, NON-STERILE
- 1 GAUZE PADS, 3" X 3", 4 PER

- 1 ADHESIVE TPE 1"X2-1/2 YD 2 PER
- 2 INSTANT COLD PACK 4" X 6"
- 2 BANDAGE COMP, 2" OFFSET, 4 PER
- 2 BANDAGE COMP, 4" OFFSET, 1 PER
- 2 ADH BAND, EXTRA LARGE, 6 PER
- 2 1 OZ EYE WASH W/PADS & STRIPS
- 1 BURN JEL 1/8 OZ, 6 PER
- 5 WATER JEL DRESSING,2" X 6"
- 1 PVP IODINE WIPES 10 PER
- 2 NITRILE GLOVES 2PR BBP
- 2 ADH BDG, CLOTH, 1"X3", 16 PER
- 1 FIRST AID GUIDE ASHI
- 2 ABD COMBINE PAD 5" X 9"
- 1 MICROSHIELD W/VNL GLV/ALCL
- 1 SCISSOR BDGE 4" RED PLS HDL
- LBL STOCK 6-3/8"X4"
- LBL STOCK 4"X2-7/8"
- 1 LBL STOCK 3"x1-7/8"
- 1 STOCK LABEL 1 7/8" X 1/2"
- 1 LBL CONTS 8"X8", CUSTOM ID B
- 1 KIT 36 UNIT PLASTIC
- 1 STING Relief WIPES 10

Burn Jel *Principal Display Panel*



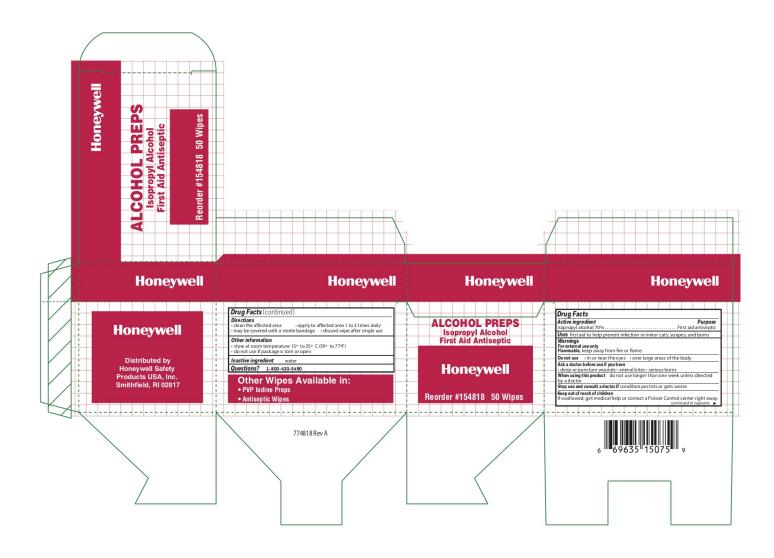
Principal Display Panel



PVP Principal Display Panel

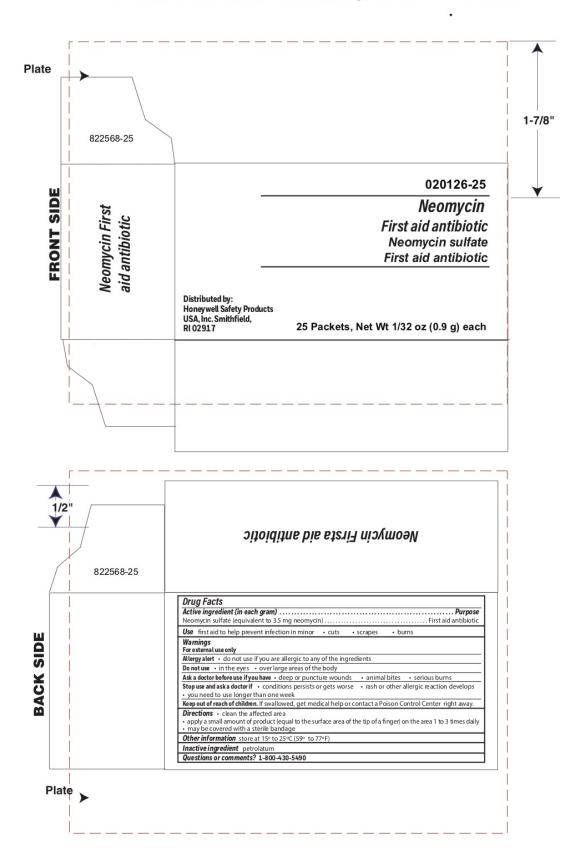


Alcohol Principal Display Panel



Neomycin Principal Display Panel

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



Principal Display Panel



Solución Isotónico Estéril

La Solution Isotonique Stérile

Isotonic Solution 16 fl. oz. (473 mL)

Sterile



Drug Facts (for USA only) Active ingredient Purpose Evewash for flushing the eye to remove loose foreign material, air pollutants, 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: so you experience eye pain of hanges in vision confined reference or pain of the eye condition worsens or irritation of the eye condition worsens or persists

We pout 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

odium chloride, sodium phosphate dibasic, sodium phosphate monobasic

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

Datos de medicamento (Para EE.UU. solamente) Usos para el lavado de ojo para quitar las particulas 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
inguna superficie. No uvelva 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 immediato
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

- quitese los lentes de contacto antes de usar la solución

- tuerza la tapa para quitar

- en juaque 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 monobásico ¿Preguntas? Llame al 1-800-430-5490

#32-000454-0000

RÉAPPROVISIONNEMENT

PEDIDO /

32-004510 Rev. J

Information

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

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

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 ouverles aux yeux ou à proximité,

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 contenar et si nécessaire, confuner de rincre avec unesolution de rinçage oculaire 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

4220 Kit Label 019643-4367

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

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4220

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4220-01	1 in 1 KIT	09/13/2018	

Quant	Quantity of Parts				
Part #	Package Quantity	Total Product Quantity			
Part 1	6 PACKET	21 g			
Part 2	10 POUCH	4 mL			
Part 3	10 POUCH	3 mL			
Part 4	2 BOTTLE	60 mL			
Part 5	10 PACKET	9 g			
Part 6	1 POUCH	0.4 mL			

Part 1 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: Z8IX2SC10H)	
OCTOXYNOL-9 (UNII: 7JPC6Y25QS)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

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

Part 3 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		
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: 059QF0KO0R)		

l	P	ackaging			
	#	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

Part 4 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 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- 01	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 5 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	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 6 of 6

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 I	Name	Strength
WATER (UNII: 059QF0KO0R)		

l	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				
Marketin Categor		ber or Monograph ation	Marketing Start Date	Marketing End Date
unapproved dru other	3		09/18/2018	

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

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

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