4189 FIRST AID KIT- 4189 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.

0498-4189: First Aid Kit (FABC, Amm Inh, EW, alcohol wipes- 010742-3236)

First Aid Burn Cream Active ingredient

Benzalkonium chloride o.13%

Lidocaine HCl 0.5%

First Aid Burn Cream *Purpose*

First aid antiseptic

External analgesic

First Aid Burn Cream *Uses*

- prevent skin infection
- for temporary relief of pain associated with minor burns

First Aid Burn Cream Warnings

For external use only

Do not use

- in or near the eyes
- if you are allergic to any of the ingredients
- in large areas of the body, particularly over raw surfaces or blistered areas
- for more than 10 days

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

 symptoms persist for more than 7 days or clear up and occurs again within a few days

First Aid Burn Cream Directions

- adults and children 2 years of age and older:
- clean the affected area
- apply a small amount of this product (equal to the surface area of the tip of a finger) onto affected area 1 to 3 times daily
- may be covered with a sterile bandage
- children under 2 years of age: consult a doctor

First Aid Burn Cream Other information

- tamper evident sealed packets
- do not use if packet is opened or torn

First Aid Burn Cream Inactive ingredients

aloe barbadensis juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, trolamine, water

First Aid Burn Cream *Questions*

1-800-430-5490

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 0 to 25 0 C (59 0 to 77 0 F)

Alcohol Wipe Inactive ingredient

water

Alcohol Wipe Questions

1-800-430-5490

Ammonia Inhalent Active ingredient (in each ampule)

Ammonia 15%

Ammonia Inhalent *Purpose*

Respiratory stimulant

Ammonia Inhalent *Uses*

to prevent or treat fainting

Ammonia Inhalent *Warnings*

For external use only

Do not use

• if you have breathing problems such as 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.

Ammonia Inhalent 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

Ammonia Inhalent Other information

store at room temperature away from light

Ammonia Inhalent Inactive ingredients

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

Ammonia Inhalent Questions

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

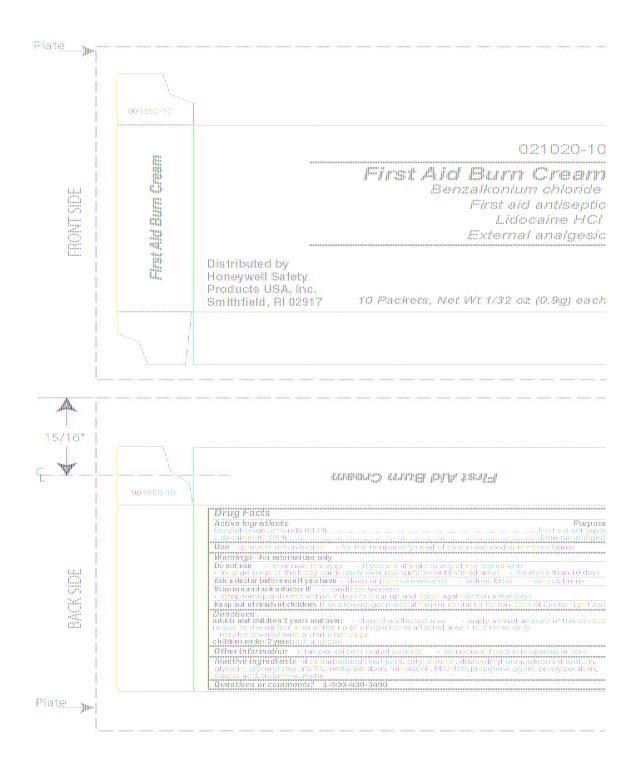
Eyewash Questions

1-800-430-5490

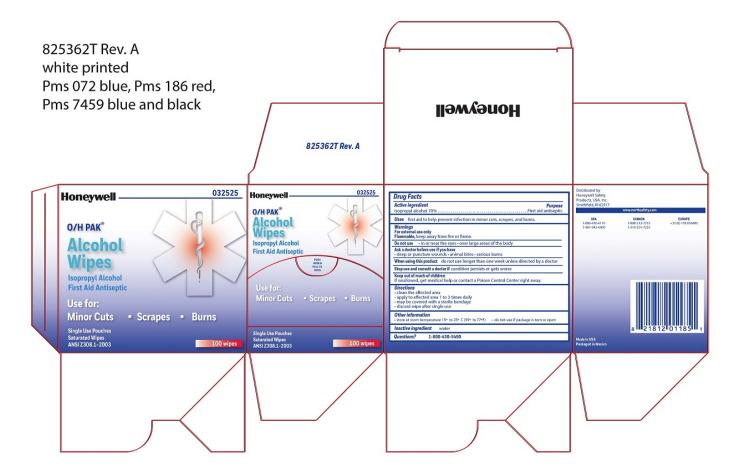
4189 010742-3236 Kit Contents

- 1 FIRST AID BURN CREAM 6 PER
- 2 AMMONIA INHALANTS 10 PER
- 1 TRIANGULAR BDG, NON-STERILE
- 2 GAUZE PADS, 3" X 3", 4 PER
- 1 FORCEPS & SCISSORS, 1 EA
- 1 GAUZE BANDAGE, 2" X 6 YD,2 PER
- 1 BUFFERED EYE WASH 1 OZ BTL
- 4 ADH BDG,PLSTIC,3/4"X3",16 PER
- 1 ALCOHOL PREP PADS 10P
- 1 ADHESIVE TAPE W/P 1/2"X 5 YD
- LBL STOCK 6-3/8"X4"
- 1 LBL STOCK 3"x1-7/8"
- 1 KIT STL 16 UN (VERTICAL)
- 1 LABL INSTR FA REV A

First Aid Burn Cream Principal Display Panel



Principal Display Panel



Ammonia Inahalent *Principal Display Panel*

Principal Display Panel

Isotónico Estéril



Isotonic Solution

16 fl. oz. (473 mL)

Isotonique Stérile

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

when the condition worsens or persists was allowed, 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 32-004510 Rev. J 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

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

4189 Kit Label 010742-3236



First Aid Kit

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

4189 FIRST AID KIT

4189 first aid kit kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4189

Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
		NDC:0498-4189-	1 in 1 KIT; Type 0: Not a Combination	10/18/2018		

Quantity of Parts				
Part #	Package Quantity	Total Product Quantity		
Part 1	6 PACKET	5.4 g		
Part 2	1 BOTTLE	30 mL		
Part 3	10 POUCH	4 mL		

Part 1 of 3

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Item Code (Source) NDC:0498-0903

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	0.13 g in 100 g	
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
WATER (UNII: 059QF0KO0R)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)		
PEG-100 STEARATE (UNII: YD01N1999R)		
LIGHT MINERAL OIL (UNII: N6K5787QVP)		

EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 903K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0903- 34	10 in 1 BOTTLE, UNIT-DOSE		
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		12/20/2017		

Part 2 of 3

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
l		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 3 of 3

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 Na	ne	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			
Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	09/18/2018				
		Citation Date			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/18/2018	

Labeler - Honeywell Safety Products USA, INC (118768815)

Revised: 1/2024 Honeywell Safety Products USA, INC