4247 FIRST AID KIT- 4247 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.

4247 First Aid Kit (FABC, Amm Inh, EW, BZK wipes- 685533T13)

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

Ammonia Inhalent

Active ingredient (in each ampule)

Ammonia 15%

Ammonia Inhalent

Purpose

Respiratory stimulant

Ammonia Inhalent

Uses

• to prevent or treat fainting

Ammonia Inhalent *Warning*s

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*

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

BZK

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK

Purpose

First aid antiseptic

BZK

Uses

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

BZK 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

Directions

• tear open packet and use as a washcloth

BZK

Other information

- store at room temperature 15^{0} to 30^{0} C (59⁰ 86⁰ F)
- do not reuse towelette

BZK

Inactive ingredients

water

BZK

Questions

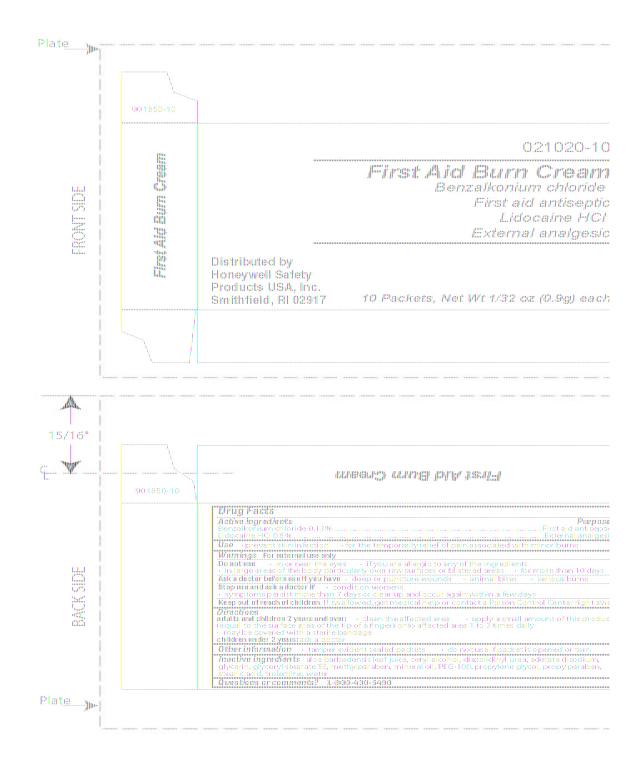
1-800-430-5490

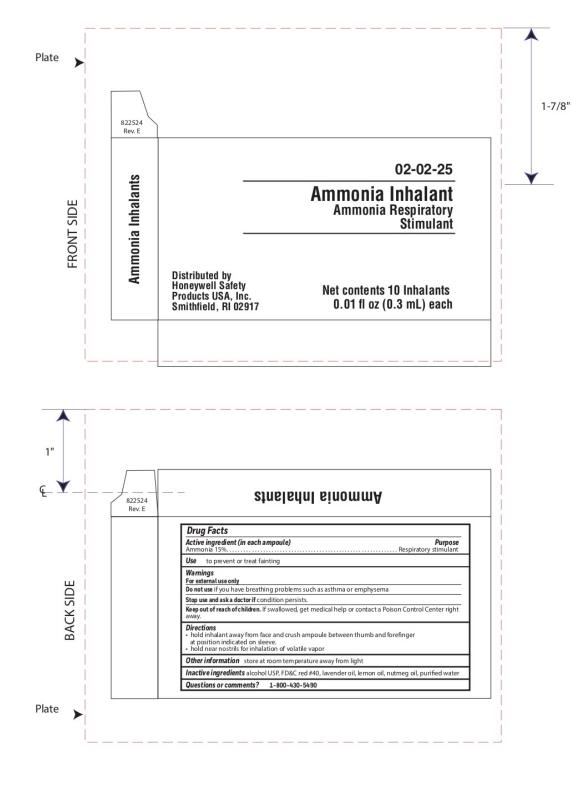
4247

685533T13 Kit Contents 1 3/4 X 3 PLAS 100/BOX 2 INSTANT COLD PACK 4" X 6" 1 ADHESIVE TAPE W/P 1/2"X 5 YD **1 FIRST AID GUIDE ASHI** 4 GAUZE CLEAN-WRAP BDGE N/S 2" 1 GAUZE CLEAN-WRAP BDGE N/S 3" 2 ABD COMBINE PAD 5" X 9" 1 GZE PADS STERILE 2"X 2" 25'S **11OZ, BUFF EYEWASH** 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" 9 BZK ANTISEPTIC WIPE, BULK **1 PR LRG NITRILE GLVES ZIP BAG** 1 FIRST AID CREAM FOIL PKS 27/BG **1 KIT STL BULK MEDIUM** 1 TRI BNDG NON WOVEN 40"X40"X56" **5 EYE PADS STD OVAL STERILE**

15 NON ADHERENT PAD 2" X 3" 10 HEAVY FLEX LARGE PATCH 2" X 3" 3 AMMONIA INHALANT, BULK

First Aid Burn Cream Principal Display Panel

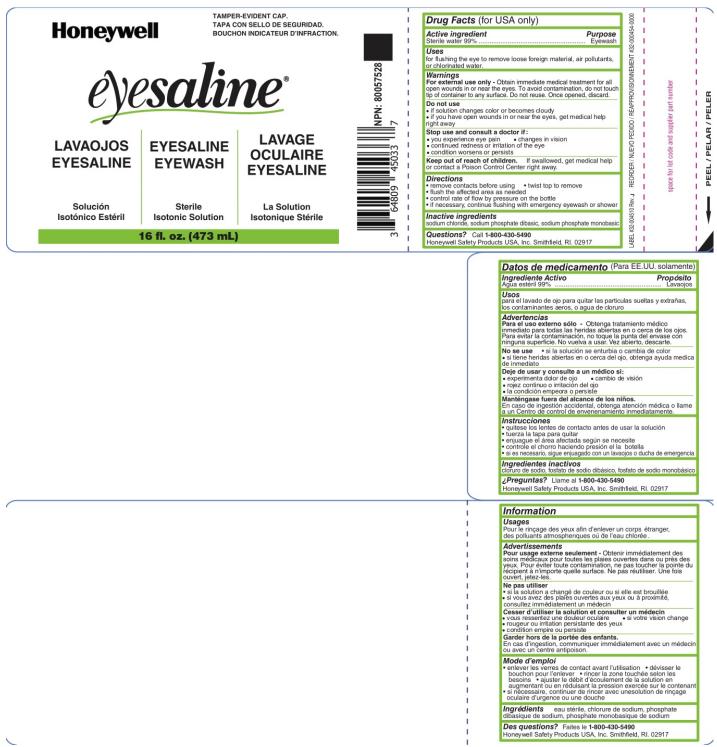


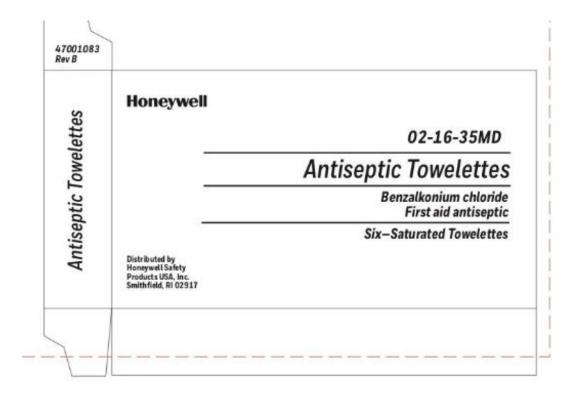


796006 Rev. E Unit Carton Printing Plate for "A" size carton.

796006 Rev. E (page 3 of 3)

Eyewash Principal Display Panel





47001083 Rev B	səttələwoT oitqəsitnA
	Drug Facts
	Active Ingredient Purpose Benzalkonium chloride 0.133% w/v First ald 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 overlarge areas of the body
	Ask a doctor before use In case of deep or puncture wounds, animal bites, orserious 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

4247 Kit Label 685533T13



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

4247 FIRST AI	D KIT			
4247 first aid kit kit				
Product Informati	on			
Product T ype	HUMAN OTC DRUG	HUMAN OTC DRUGItem Code (Source)NDC:0498-424		
Packaging				
# Item Code	Package Description		Marketing Start Date	Marketing End Date
1 NDC:0498-4247-01	1 in 1 KIT; Type 0: Not a Combination F	in 1 KIT; Type 0: Not a Combination Product		

Quantity of Parts					
Part # Packa	nge Quantity	Total Product Quantity			
Part 1 27 PACKET		24.3 g			
Part 2 3 AMPULE		0.9 mL			
Part 3 1BOTTLE		30 mL			
Part 4 9 PACKET		12.6 mL			
Part 1 of 4					
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)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

	Ingredient Name		Strength
PROPYLENE GLYCOL (U	INII: 6DC9Q167V3)		
ALOE VERA LEAF (UNII:	ZY81Z83H0X)		
WATER (UNII: 059QF0KO	0 R)		
STEARIC ACID (UNII: 4EL	V7Z65AP)		
METHYLPARABEN (UNII:	A2I8C7HI9T)		
CETYL ALCOHOL (UNII:	936JST6JCN)		
GLYCERYL MONOSTEA	RATE (UNII: 230OU9XXE4)		
PEG-100 STEARATE (UN	I: YD01N1999R)		
LIGHT MINERAL OIL (UI	NII: N6K5787QVP)		
EDETATE DISODIUM (UN	II: 7FLD91C86K)		
TROLAMINE (UNII: 903K	93S3TK)		
GLYCERIN (UNII: PDC6A3	COOX)		
PROPYLPARABEN (UNII:	Z8IX2SC1OH)		
DIAZOLIDINYL UREA (U	NII: H5RIZ3MPW4)		
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date

1 0.9 g i	in 1 PACKET; Ty	pe 0: Not a Combination Product		
Marketing Info				
Marketing Category		ion Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not fina	al part333A		12/20/2017	
Part 2 of 4				
AMMONIA IN	ΗΔΙ FNT			
ammonia inhalent inh				
	lialain			
Product Information	ion			
Item Code (Source)		NDC:0498-3334		
Route of Administrat	ion	RESPIRATORY (INHALATION)		
Active Ingredient/		•		
	0	ent Name	Basis of Strength	Strength
AMMO NIA (UNII: 51380	219F1X) (AMMC	0NIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL
Inactive Ingredier	nts			
]	Ingredient Name		Strength
ALCOHOL (UNII: 3K99	58V90M)			
Packaging				
# Item Code		Package Description	Marketing Start Date	Marketing End Date
1 NDC:0498-3334-00	0.3 mL in 1 AMP	ULE; Type 0: Not a Combination Product	t	
Marketing Info				
Marketing Category	Applicatio	n Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other			09/18/2018	
Part 3 of 4				
	MERGEN	ICY EYEWASH		
purified water liquid				
purmed water iiquid				

	n					
Item Code (Source)		NDC:0498-0100				
Route of Administration	n	OPHTHALMIC				
Active Ingredient/A	ctive Moi	ety				
	Ingredie	nt Name	Basis	of Strength	St	trength
WATER (UNII: 059QF0KO	00R) (WATER	- UNII:059QF0KO0R)	WATER		98.6 mL	in 100 mL
Inactive Ingredients	5					
		Ingredient Name				Strength
SO DIUM PHO SPHATE, M	IONOBASIC	, MONOHYDRATE (UNII: 593YOG76R)	V)			
SODIUM CHLORIDE (UN	II: 451W47IQ8	3X)				
SODIUM PHO SPHATE, D	IBASIC (UNI	I: GR686LBA74)				
Packaging						
# Item Code		Package Description		ting Start Date	Market	ting End Dat
1 NDC:0498-0100-01 30	mL in 1 BOT	TLE; Type 0: Not a Combination Produc	t			
wial keung inioi	mation					
Marketing Information Marketing Category OTC monograph final		on Number or Monograph Citation	Marke 12/18/201	ting Start Date 18	Market	ting End Date
Marketing Category OTC monograph final p	Applicatio	on Number or Monograph Citation		-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4	Applicatio part349			-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T	Applicatio part349 OWELE			-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4	Applicatio part349 OWELE			-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T(benzalkonium chloride	Applicatio part349 OWELE liquid			-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride	Applicatio part349 OWELE liquid	TTE		-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information Item Code (Source)	Applicatio part349 OWELE liquid	TTE NDC:0498-0501		-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information	Applicatio part349 OWELE liquid	TTE		-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information Item Code (Source) Route of Administration	Applicatio part349 OWELE liquid	TTE NDC:0498-0501 TOPICAL		-	Market	ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information Item Code (Source) Route of Administration	Application part349 OWELE liquid n n ctive Moie	TTE NDC:0498-0501 TOPICAL		-		ting End Date
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information Item Code (Source) Route of Administration	Applicatio Dart349 OWELE liquid n n ctive Moie Ingr	TTE NDC:0498-0501 TOPICAL		18	rength	
Marketing Category OTC monograph final p Part 4 of 4 ANTISEPTIC T benzalkonium chloride Product Information Item Code (Source) Route of Administration Active Ingredient/Addition	Applicatio Dart349 OWELLE liquid n n ctive Moie Ingr RIDE (UNII: F	TTE NDC:0498-0501 TOPICAL ety redient Name		18 Basis of Str BENZALKONIUM	rength	Strength 1.3 mg

keting Start Date Mar	rketing End Date				
keting Start Date Mai	rketing End Date				
-	I Ketting Elite Date				
OTC monograph not final part333E 09/18/2018 Marketing Information					
	rketing End Date				
	keting Start Date Mar				

Labeler - Honeywell Safety Products USA, INC (079287321)

Establishment

Name	Address	ID/FEI	Business Operations
James Alexander		040756421	manufacture(0498-3334)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4247)

Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0903)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc.		167518617	manufacture(0498-0100)

Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0501)

Revised: 4/2019

Honeywell Safety Products USA, INC