4146 FIRST AID KIT- 4146 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.

0498-4146: First Aid Kit (Burn Jel, BZK wipe, alcohol wipe, sting relief-SF00004042)

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

BZK Wipe Active ingredient

Benzalkonium chloride 0.13% w/v

BZK Wipe Purpose

First aid antiseptic

BzK Wipe *Uses*

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

BZK Wipe 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 Wipe Directions

tear open packet and use as a washcloth

BZK Wipe Other information

- store at room temperature 15 0 to 30 0 C (59 0 86 0 F)
- do not reuse towelette

BZK Wipe Inactive ingredient

water

BZK Wipe Questions

1-800-430-5490

Sting Relief Active ingredient (in each wipe)

Ethyl alcohol 50.0%

Lidocaine HCI 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

Alcohol Active ingredient

Isopropyl alcohol 70%

Alcohol *Purpose*

First aid antiseptic

Alcohol

Uses

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

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

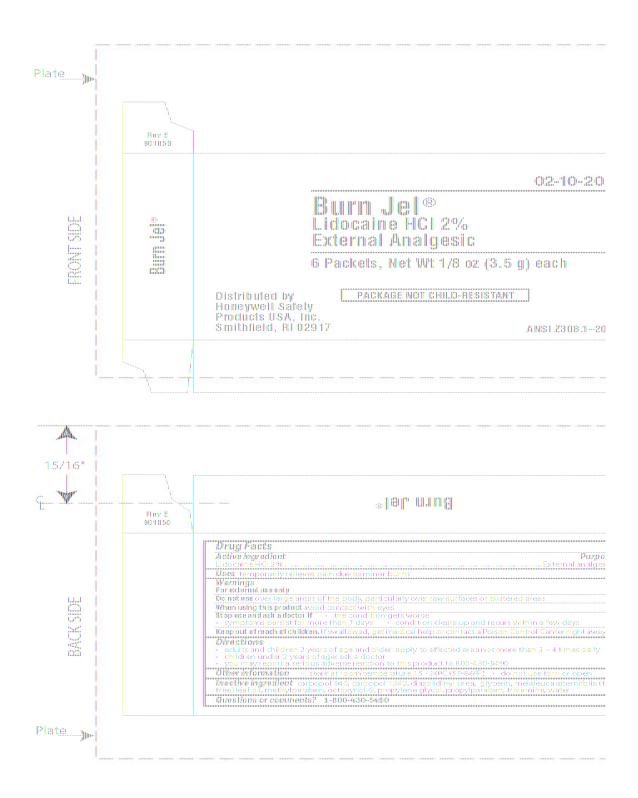
Alcohol Questions

1-800-430-5490

4146 SF00004042 kit contents

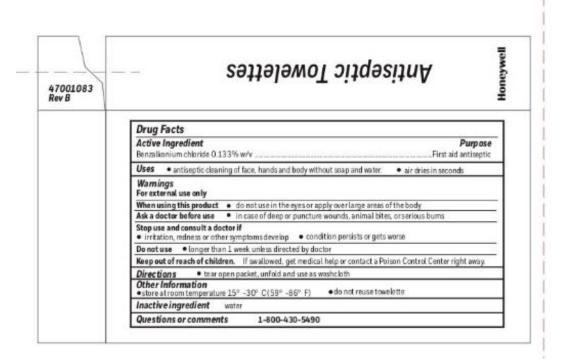
- 1 KNUCKLE BAND 8 PER
- 1 TRIANGULAR BDG, NON-STERILE
- 1 GAUZE PADS, 3" X 3", 4 PER
- 1 ADH TAPE, .5" X 2.5 YD, 2 PER
- 1 GAUZE COMPRESS, 1728 SQ IN 1
- 1 FORCEPS & SCISSORS, 1 EA
- 1 INSTANT COLD PACK 4" X 6"
- 1 ADHESIVE BDG,PLSTIC,1"X3"16PER
- 1 BURN JEL 1/8 OZ, 6 PER
- 1 NITRILE GLOVES 2PR BBP
- 1 ANTIMCRBL ANTSPTC TWLETTS
- 1 MICROSHIELD W/VNL GLV/ALCL
- 1 F. A. INST CHART SM (INDIVIDUAL LBL)
- 1 BANDAGE COMP 4" W/TELFA PAD 1
- LBL STOCK 6-3/8"X4"
- LBL STOCK 4"X2-7/8"
- 1 LBL STOCK 3"x1-7/8"
- 1 KIT STL 16 UN (VERTICAL)
- 1 STING Relief SWAB 10

Burn Jel *Principal Display Panel*

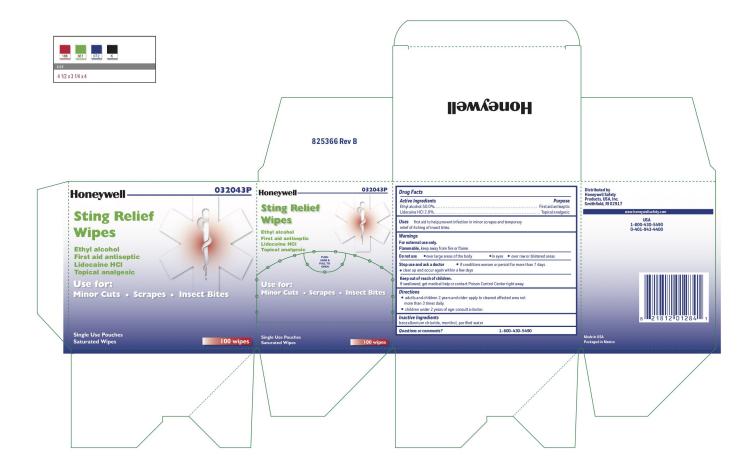


Principal Display Panel

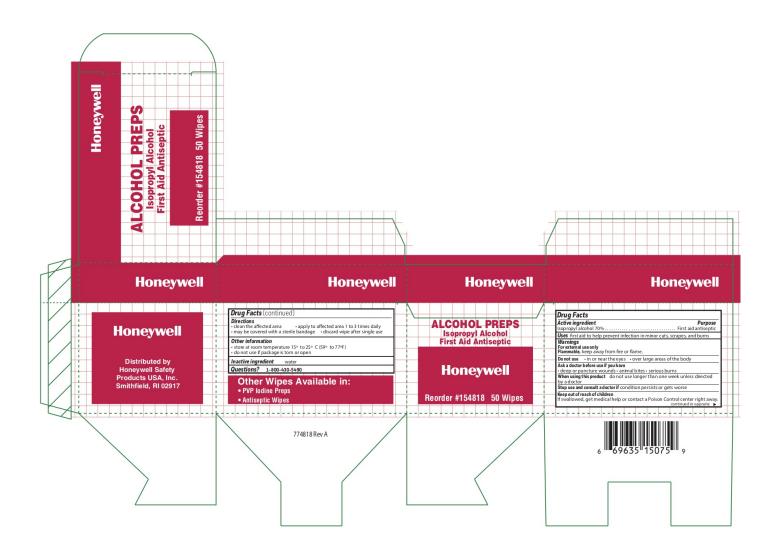




Sting Relief
Principal Display Panel



Alcohol Principal Display Panel



4146 Kit Label SF00004042





CITY OF SEATTLE

FIRST AID KIT

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

4146 FIRST AID KIT

4146 first aid kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4146

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4146-01	1 in 1 KIT	09/13/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	21 g

Part 2	2 PACKET	2.8 mL
Part 3	4 POUCH	1.6 mL
Part 4	10 POUCH	4 mL

Part 1 of 4

BURN JEL

UNII:98PI200987)

gel for burns gel

Product Information

Item Code (Source) NDC:0498-0203

Route of Administration TOPICAL

DIPROPYLENE GLYCOL (UNII: E107L85C40)

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - LIDOCAINE HYDROCHLORIDE 2 g

ANHYDROUS

in 100 g

Ingredient Name

Strength

TEA TREE OIL (UNII: VIF565UC2G)

DIAZOLIDINYL UREA (UNII: H5RIZ 3MPW4)

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)

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:0498-0203-	3.5 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
unapproved drug other		09/19/2018	

Part 2 of 4

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source) NDC:0498-0501

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

- u = u = u = u = u = u = u = u = u = u					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:0498-0501- 00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
unapproved drug other		12/22/2017	

Part 3 of 4

ALCOHOL WIPE

isopropyl alcohol swab

Product Information

NDC:0498-0143 Item Code (Source)

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Basis of Ingredient Name Strength Strength ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -ISOPROPYL 0.7 mL

UNII:ND2M416302)

in 1 mL ALCOHOL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:0498-0143-	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		09/18/2018	

Part 4 of 4

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information

Item Code (Source) NDC:0498-0733

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Active ingredient/Active Molecy		
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					
4	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
:	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				

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)

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