4370 FIRST AID KIT- 4370 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.

0498-4370: First Aid Kit (Hand Sanitizer, BZK wipe, FABC, Neomycin, Sting rel-SF00004664/M)

First Aid Burn Cream Active ingredient

Benzalkonium chloride o.13%

Lidocaine HCI 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

Other information

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

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

Questions

1-800-430-5490

BZK Antiseptic Wipe Active ingredient

Benzalkonium chloride 0.13%

BZK *Purpose*

First aid antiseptic

BZK *Uses*

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

BZK Warnings

For external use only

BZK

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 ° to 30 ° C (59 ° 86 °F)
- do not reuse towelette

BZK Inactive ingredients

water

BZK Questions

1-800-430-5490

Sting Relief Active ingredient

Ethyl alcohol 50.0%

Lidocaine HCI 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 ingredients

benzalkonium chloride, menthol, and purified water

Sting relief Questions or Comments

1-800-430-5490

Neomycin Antibiotic Ointment Active ingredient

Neomycin Antibiotic Ointment *Purpose*

First aid antibiotic

Neomycin Antibiotic Ointment *Uses*

first aid to help prevent infection in - minor cuts - scrapes - burns

Neomycin Antibiotic Ointment *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 Antibiotic Ointment *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 Antibiotic Ointment *Other information*

store at 15 ° to 25 °C (59 ° to 77 °F)

Neomycin Antibiotic Ointment *Inactive ingredient*

petrolatum

Neomycin Antibiotic Ointment *Questions*

1-800-430-5490

Hand Sanitizer Active ingredient

Ethyl alcohol 62%

Hand Sanitizer Purpose

Antiseptic handwash

Hand Sanitizer Uses

- for hand washing to decrease bacteria on skin
- recommended for repeated use

Hand Sanitizer Warnings

For external use only

Flammable, keep away from fire or flame

When using this product

- do not use in the eyes
- discontinue use if irritation and redness develops
- If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children

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

Hand Sanitizer Directions

• wet hands thoroughly with product and allow to dry without wiping

Hand Santitizer

Other information

- place a quarter size amount into one hand, spread over both hands to wrist and rub into skin until dry
- store at 15 ° to 25 ° C (59 ° to 77 ° F)

Hand Sanitizer Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, dl-alpha tocopheryl acetate, fragrance, PEG-60 almond glycerides, propylene glycol, purified water .

Hand Sanitizer Questions or Comments

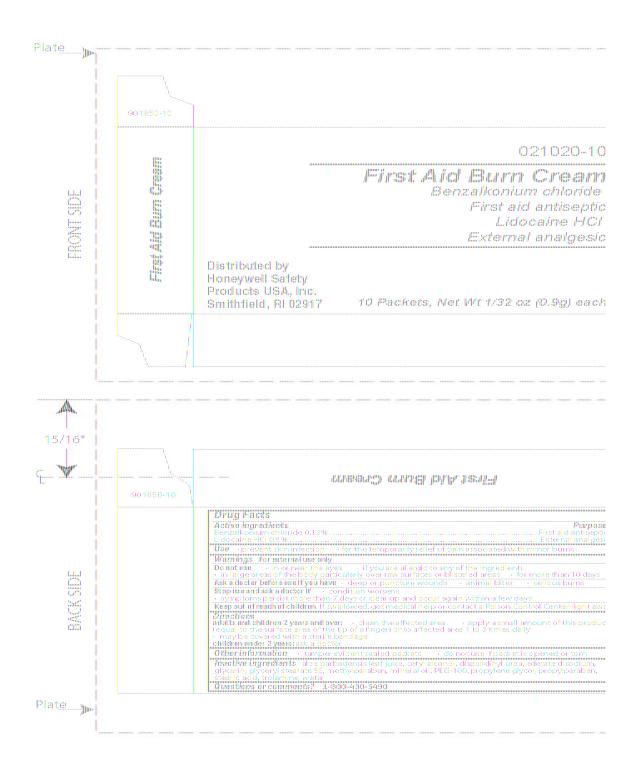
1-800-275-3433 info@waterjel.com

4370 SF00004664/M Kit Contents

- 1 1X3 PLASTIC 100/BOX
- 1 EYE DRESS PKT W/4 ADH STRIPS
- 1 TWEEZER PLASTICS 4"
- 1 FIRST AID GUIDE ASHI
- 10 HAND SANITIZER 0.9G WJ BULK
- 3 GAUZE CLEAN-WRAP BDGE N/S 2"
- 2 GAUZE CLEAN-WRAP BDGE N/S 3"
- 2 ABD COMBINE PAD 5" X 9"
- 1 GZE PADS STERILE 3"X 3" 10'S
- 1 CO-FLEX BANDAGE 2"X 5YDS TAN
- 1 CPR FILTERSHIELD 77-100
- 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"
- 10 BZK ANTISEPTIC WIPE, BULK
- 4 PR LRG NITRILE GLVES ZIP BAG
- 10 FIRST AID BURN CREAM 1.0GR PKT EACH
- 2 TAPE ADHESIVE 1/2 X 2.5 125133

- 10 POUCH NEOMYCIN ANTIBIOTIC .9 G
- 1 WATER-JEL BURN DRESSING 4 X 4
- 2 ADH BNDG PLASTIC EX-LG 4"X 2"
- 1 KIT PP 24 UNIT FA
- 1 LBL CONTENTS ANSI 2015 CL A
- 1 BAG ZIPPER POLY 6 X 6 2 MIL
- 6 SAFETEC STING RELIEF WIPES BULK
- 1 TRI BNDG NON WOVEN 40"X40"X56"
- 1 COLD PACK UNIT 4"X6" BULK
- 4 WOVEN FINGERTIP BANDAGE 2"
- 3 WOVEN KNUCKLE BANDAGE

First Aid Burn Cream Principal Display Panel

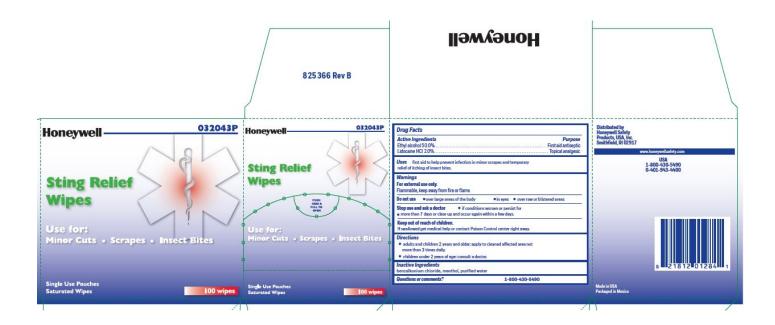


Principal Display Panel



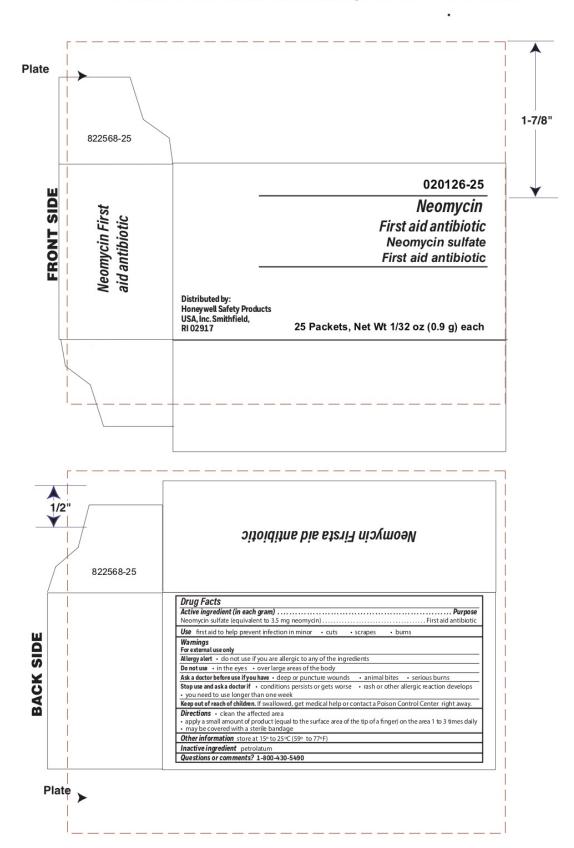
47001083 Rev B	Antiseptic Towelettes	Honeywell	
	Drug Facts	\neg	
	manufacturing and options to the control of the con	urp ose ntiseptic	
	Uses • antiseptic cleaning of face, hands and body without soap and water. • air dries in seconds		
	Warnings For external use only		
	When using this product • do 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, or serious burns Stop use and consult a doctor if irritation, redness or other symptoms develop condition persists or gets worse		
	Do not use ● longer than I week unless directed by doctor	10	
	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	- 12	
	Other Information • store at room temperature 15° -30° C(59° -86° F) • do not reuse towelette		
	Inactive ingredient water		

Sting Relief Principal Display Panel



Neomycin Antibiotic Ointment Principal Display Panel

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





Hand Sanitizer

Antiseptic Gel
With Vitamin E & Aloe

Kills 99.9% of Germs Without Water

240mL - (8 fl oz)

4370 Kit Label SF00004664/M



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

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4370

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0498-4370- 01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quant	Quantity of Parts			
Part #	Package Quantity	Total Product Quantity		
Part 1	6 POUCH	2.4 mL		
Part 2	10 PACKET	9 g		
Part 3	10 PACKET	9 g		
Part 4	10 PACKET	14 mL		
Part 5	10 PACKET	9 mL		

Part 1 of 5

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				
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1		0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

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

Part 2 of 5

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 HYDROCHI ORIDE	0.5 g in 100 a		

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: Z8IX2SC10H)	
DIAZOLIDINYL UREA (UNII: H5RIZ 3MPW4)	

	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		12/20/2017		

Part 3 of 5

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)				

	Packaging				
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
:	NDC:0498-0730-	0.9 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		03/31/2010	

Part 4 of 5

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source) NDC:0498-0501

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)

Basis of Strength

BENZALKONIUM - BENZALKONIUM - CHLORIDE

1.3 mg
in 1 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R)

Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing E Category Citation Date Date				
unapproved drug other		12/21/2017		

Part 5 of 5

INSTANT HAND SANITIZER

alcohol liquid

Product Information

Item Code (Source) NDC:59898-420

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (LINII: 3K9958V90M) (ALCOHOL - LINII: 3K9958V90M)	ALCOHOL	62 ml in 100 ml

Inactive Ingredients

mactive ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
.ALPHATOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)		
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)		
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)		
WATER (UNII: 059QF0KO0R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		

Packaging

ı	_				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:59898-420- 36	0.9 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/15/2011	

Marketing Information

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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)