

**4128 FIRST AID KIT- 4128 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.

0498-4128: First Aid Kit (BZK wipes, FABC- SF00004238)

First Aid Burn Cream
Active ingredient

Benzalkonium chloride 0.13%

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

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

Keep out of reach of children

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

Stop use and ask a doctor if

- irritation, redness or other symptoms develop
- the condition persists or gets worse

BZK

Directions

- tear open packet and use as a washcloth

BZK

Other information

- store at room temperature 15⁰ to 30⁰ C (59⁰ - 86⁰ F)
- do not reuse towelette

BZK

Inactive ingredients

water

BZK

Questions

1-800-430-5490

4128

SF00004238 Kit Contents

LBL STOCK 6-3/8"X4"

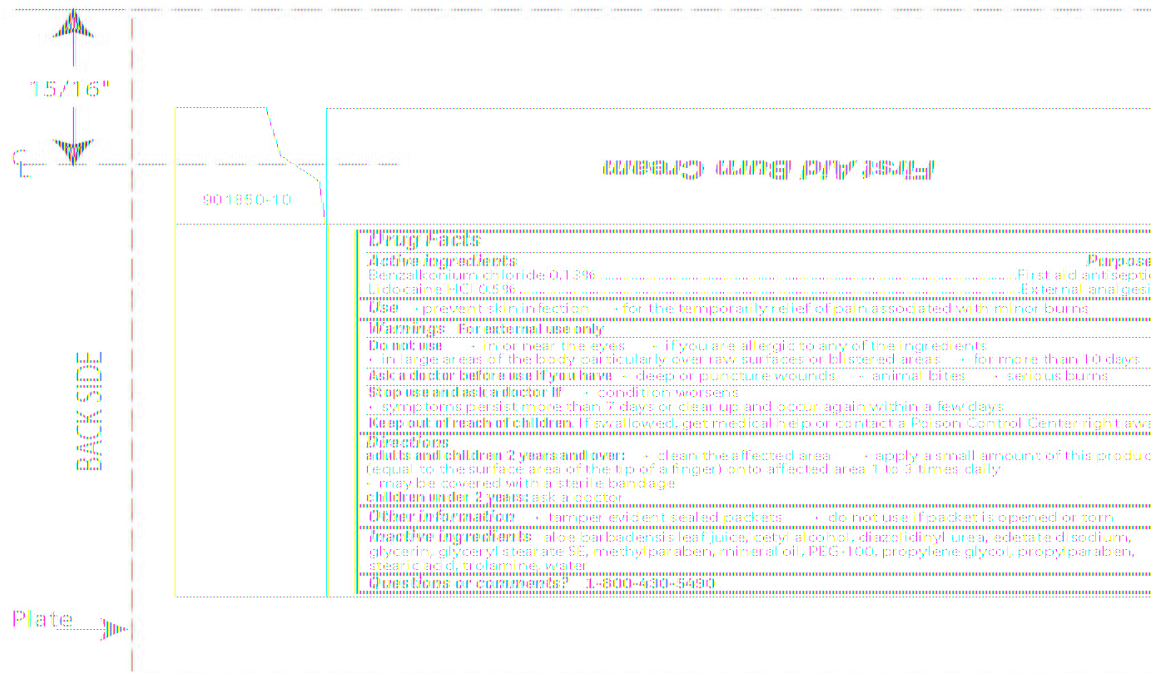
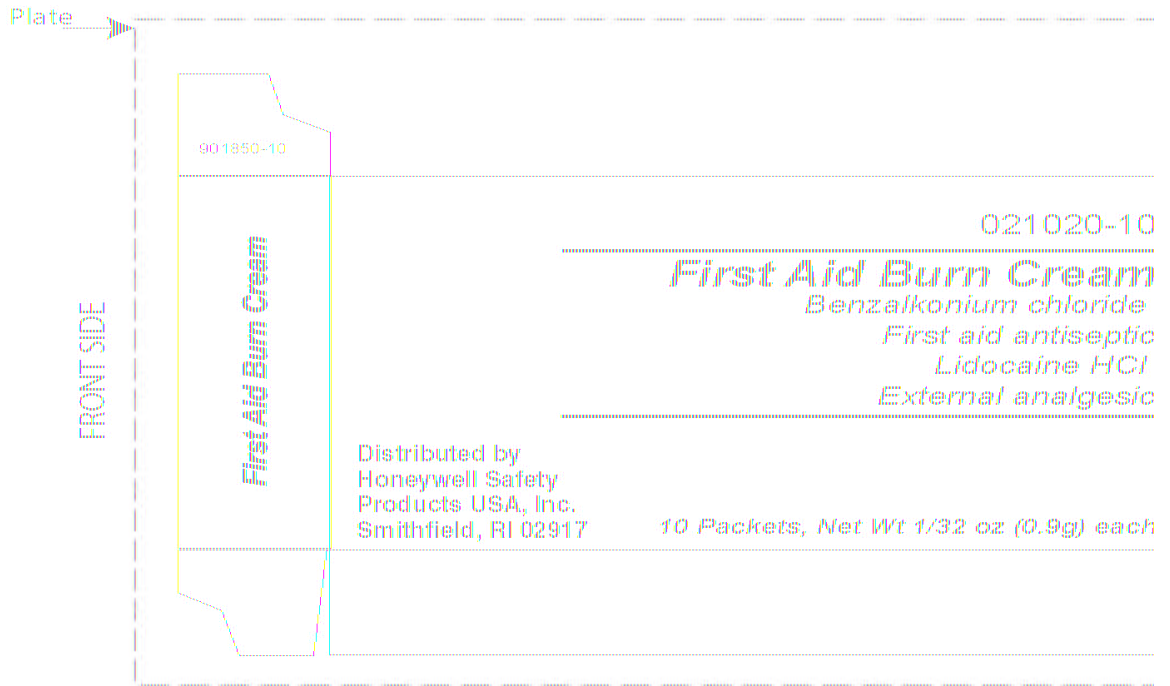
1 LBL STOCK 3"x1-7/8"

4 BZK ANTISEPTIC WIPE, BULK

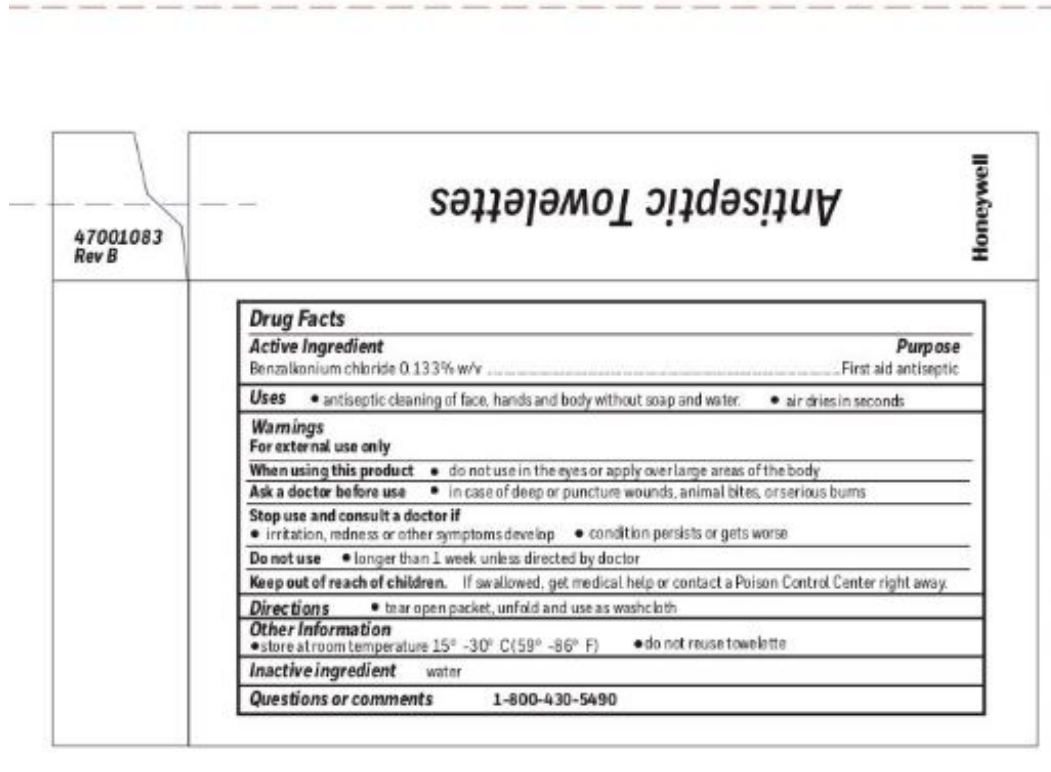
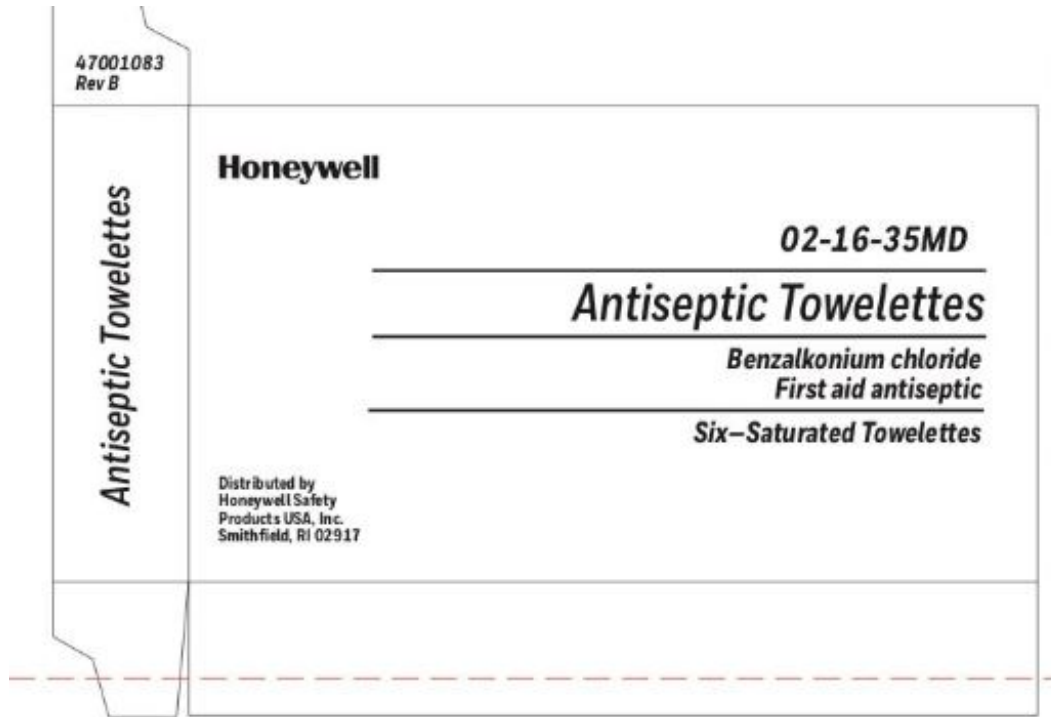
4 FIRST AID CREAM 1.0GR PKT EACH

1 KIT PP PROMOTIONAL FA KIT
10 ADH BANDAGE 3/8" X 1 1/2" DNX
8 PLASTIC BANDAGE 3/4" X 3"
2 WOVEN KNUCKLE BANDAGE
2 HEAVY FLEX LARGE PATCH 2" X 3"

**First Aid Burn Cream
Principal Display Panel**



Principal Display Panel



4128 Kit Label
SF00004238

Honeywell

Compact First Aid Kit

Distributed By



www.wisesafetyenv.com

4128 FIRST AID KIT

4128 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4128
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4128-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 PACKET	3.6 g
Part 2	4 PACKET	5.4 mL

Part 1 of 2

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
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: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

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 2 of 2

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source) NDC:0498-0501

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0501-00	1.4 mL 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		09/18/2018	

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)