4295 FIRST AID KIT- 4295 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.

0498-4295: First Aid Kit (Triple,Burn Jel, BZK wipe, antiseptic hand gel- SF00002877)

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 JEl
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

Triple
Active ingredient

Bacitracin zinc 400 units

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Polymyxin B sulfate 5000 units

Triple
Purpose

First aid antibiotic

First aid antibiotic

First aid antibiotic

Triple
Uses

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

Triple
Warnings

For external use only

Allergy alert: **do not use if you are allergic to any of the ingredients**

Do not use

- in the eyes
- over large areas of the body
- Ask a doctor before use if you have
- a 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 the reach of children

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

Triple
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

Triple
Other information

- store at 15 ^{0} to 25 ^{0} C (59 ^{0} to 77 ^{0} F)
- tamper evident sealed packets
- do not use if packet is torn or opened

Triple
Inactive ingredient

petrolatum

Triple
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

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 Sanitizer

Other information

- store at 15 ^{0} to 25 ^{0} C (59 ^{0} to 77 ^{0} 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, triisopropanolamine

Hand Sanitizer
Questions or Comments?

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

4252
68P2CCU Kit Contents

1 TRIPLE ANTIBIOTIC 10 PER

1 INSTANT COLD PACK 4" X 6"

1 BURN JEL 1/8 OZ, 6 PER

1 POR. CLOTH TAPE 2X10Yd

1 pOR. CLOTH TAPE 1/2X10Y

3 GAUZE CLEAN-WRAP BDGE N/S 4"

1 ABD COMBINE PAD 5" X 9"

1 ABD PADS 8"X10" STERILE

1 ELASTIC BANDAGE 3" X 4.5YD

1 CPR FILTERSHIELD 77-100

1 ANTISEPTIC WIPES BZK CHL 20'S

1 SCISSOR UTILITY SHEARS 7-1/4"

LBL STOCK 6-3/8"X4"

LBL STOCK 4"X2-7/8"

1 LBL STOCK 3"x1-7/8"

3 PR LRG NITRILE GLVES ZIP BAG

1 ANTISEPTIC HAND GEL 4OZ

1 WATER-JEL BURN DRESSING 4 X 4

1 KIT PP 24 UNIT FA

2 TRI BNDG NON WOVEN 40"X40"X56"

1 EYE PADS STD OVAL STERILE

1 GAUZE PADS 4"X4" 12PLY

5 WOVEN FINGERTIP BANDAGE 3"

10 HEAVY FLEX BANDAGE 7/8" X 3"

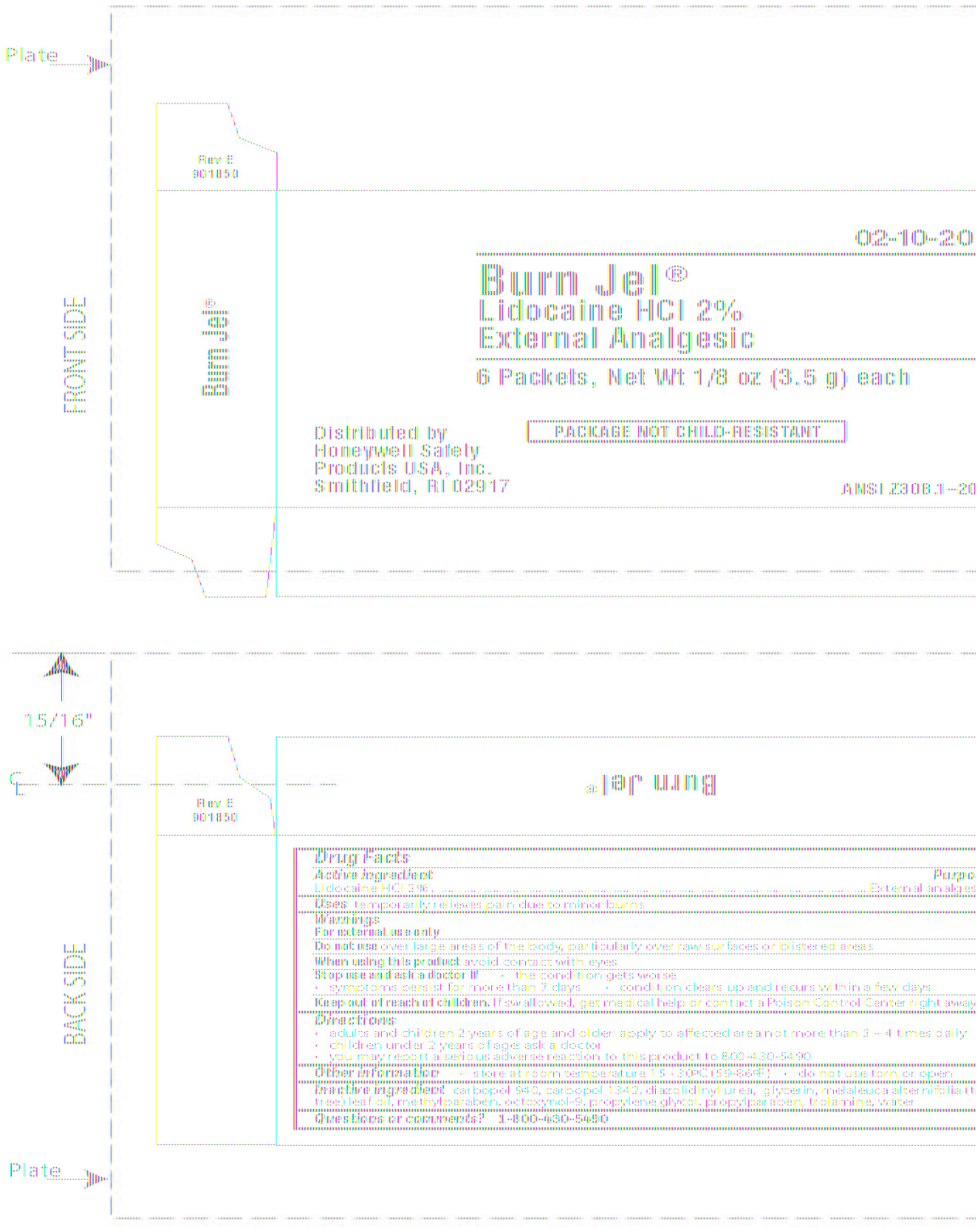
5 HEAVY FLEX KNUCKLE BANDAGE

5 HEAVY FLEX LARGE PATCH 2" X 3"

1 ZIP-LOCK BAG 5" X 5" .002

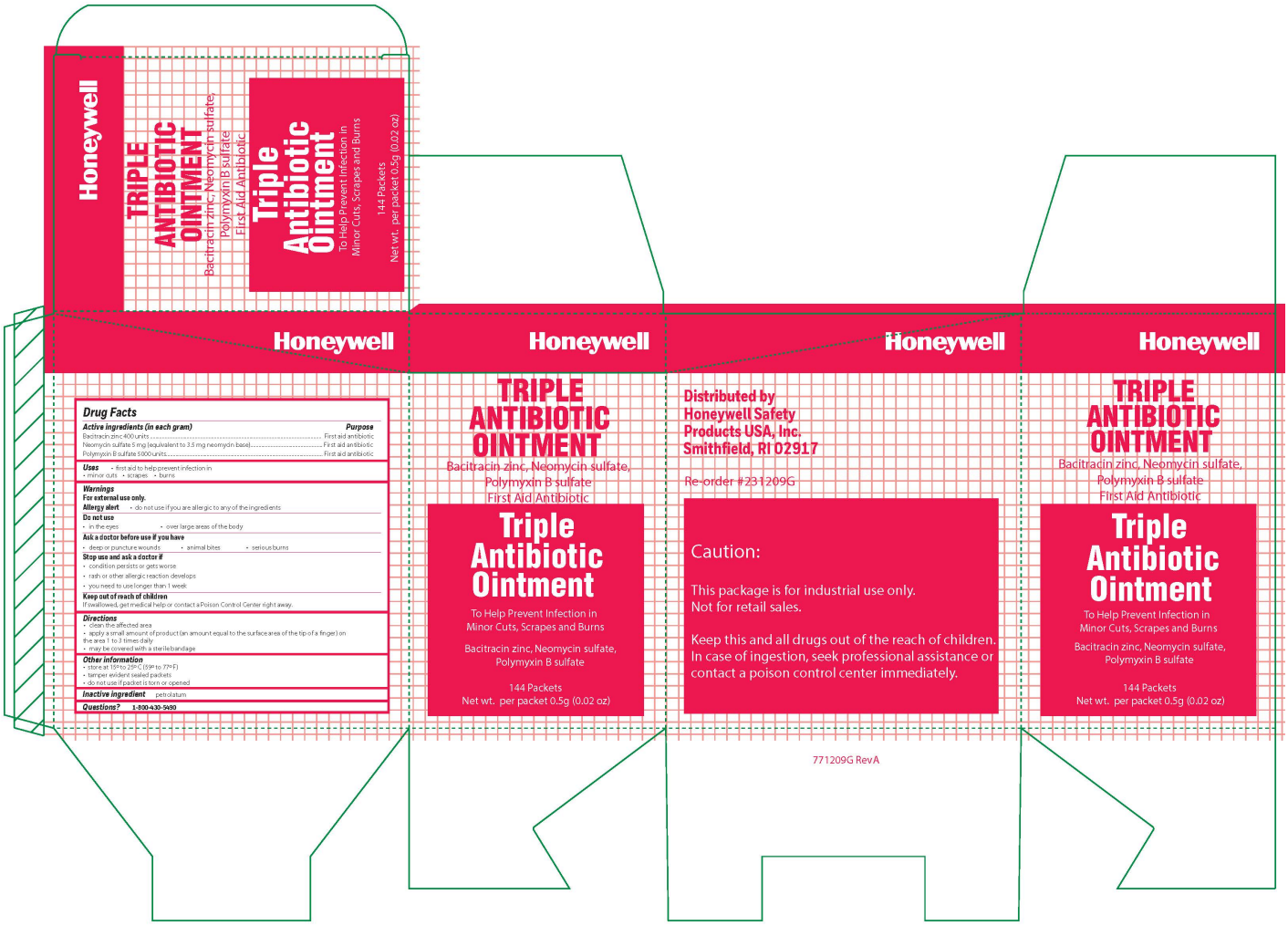
Burn Jel
Principal Display Panel

796353 Rev. E Unit Carton Printing Plate for "B" size cartor



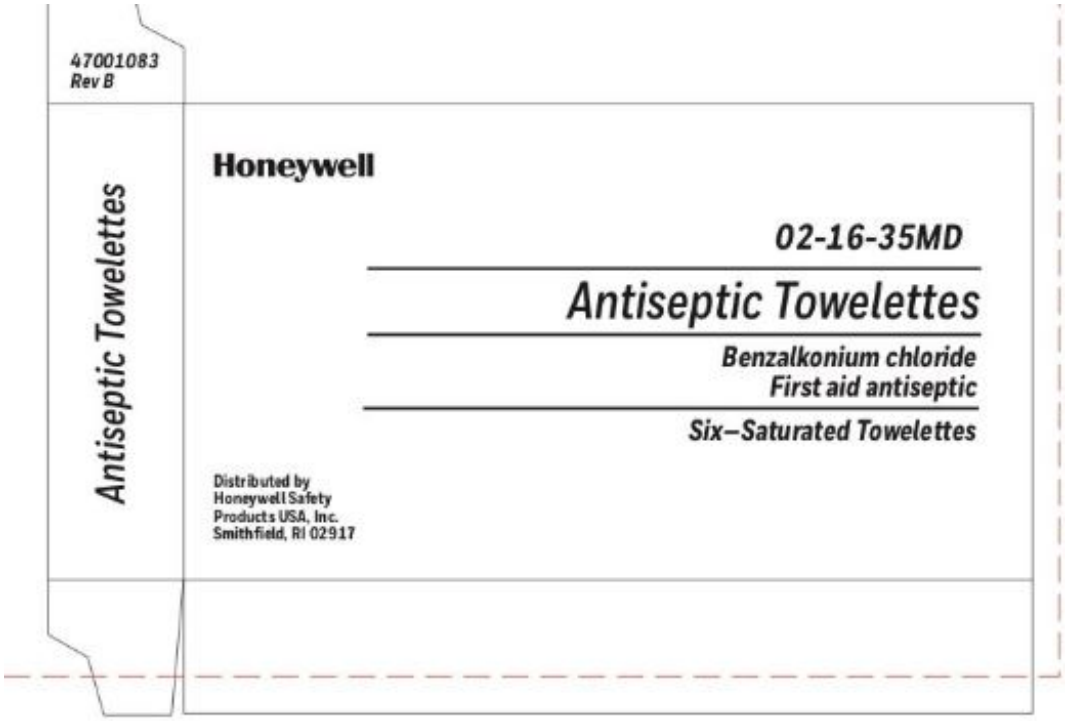
Triple

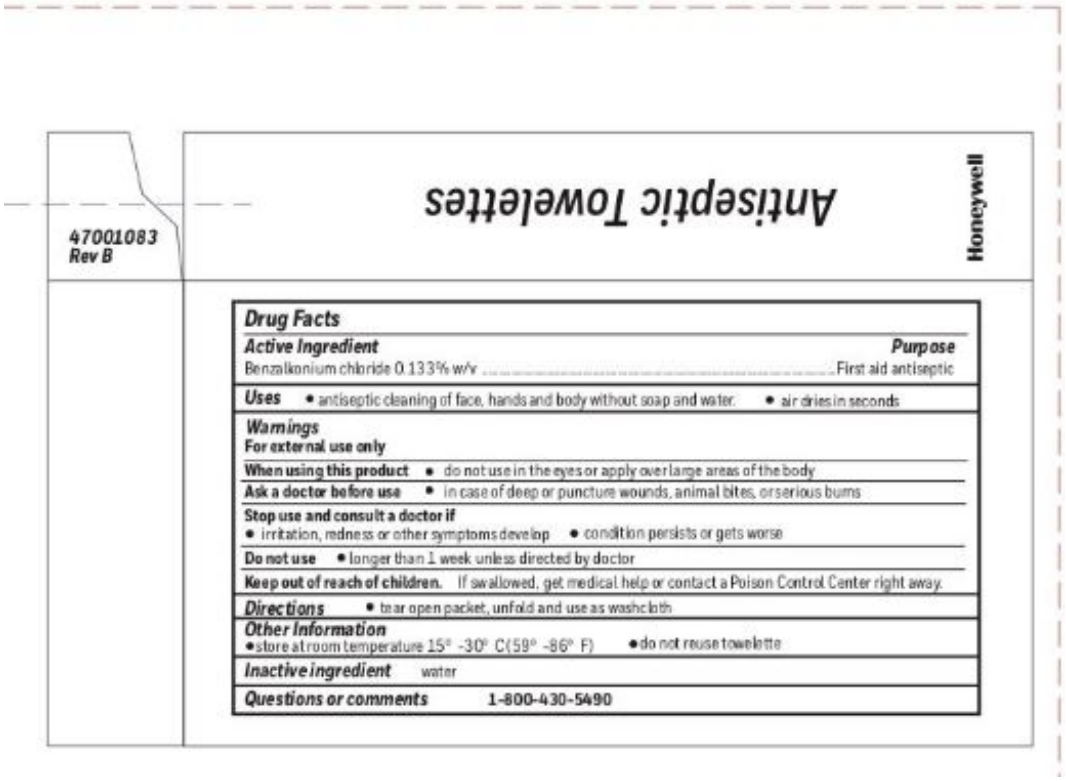
Principal Display Panel



BZK Wipe

Principal Display Panel





Hand Sanitizer
Principal Display Panel

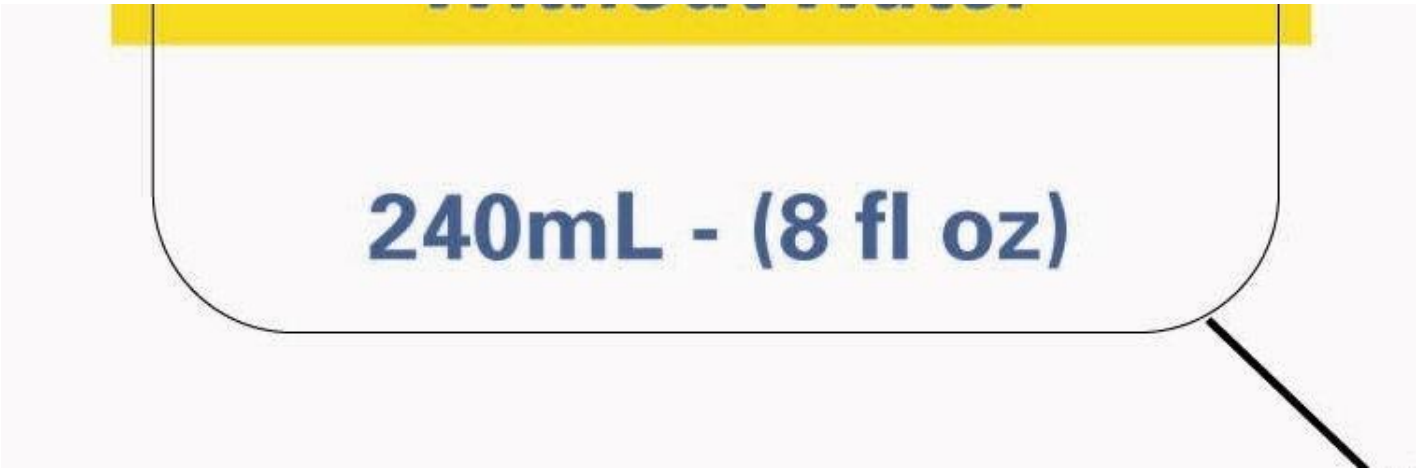
WATER·JEL®

INSTANT

Hand Sanitizer

Antiseptic Gel
With Vitamin E & Aloe

Kills 99.9% of Germs
Without Water



4295 Kit Label
SF00002877

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

4295 FIRST AID KIT

4295 first aid kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-4295

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4295-01	1 in 1 KIT	09/13/2018	

Quantity of Parts		
Part #	Package Quantity	Total Product Quantity
Part 1	6 PACKET	21 g
Part 2	10 PACKET	9 g
Part 3	1 PACKET	1.4 mL
Part 4	1 BOTTLE, PLASTIC	118 mL

Part 1 of 4

BURN JEL

gel for burns gel

Product Information	
Item Code (Source)	NDC:0498-0203
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
TEA TREE OIL (UNII: VIF565UC2G)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
OCTOXYNOL-9 (UNII: 7JPC6Y25QS)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0203-00	3.5 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		09/19/2018	

Part 2 of 4

TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information	
Item Code (Source)	NDC:0498-0750
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0750-35	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		09/19/2018	

Part 3 of 4

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		12/22/2017	

Part 4 of 4

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 (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59898-420-12	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/15/2010	

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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