EASY CARE FIRST AID - ALL PURPOSE - benzalkonium chloride, benzocaine, sd alcohol, bacitracin zinc, neomycin sulfate, polymyxin b, ibuprofen, acetaminophen
Tender Corp dba Adventure Medical Kits

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Easy Care First Aid Kit Sports and Travel

Active Ingredient

Active Ingredient:

Benzalkonium Chloride 0.40%

Purpose

Antiseptic

Use

For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

Warnings

Warning: For external use only.

Keep out of reach of children: If swallowed, get medical help or contact a Poison Control Center right away.

Stop use if unusual redness, swelling or other symptoms occur. Consult a physician immediately.

Do not use in the eyes or over large areas of the body.

Directions

Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.

Inactive Ingredients

Inactive Ingredient: Purified water

LOT/EXP: Made in CHINA

20130301

Antiseptic Towelette

Genuine First Aid LLC, Clearwater FL 33755

www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

Active Ingredient: Purpose:

Benzocaine, 6% w/v..... Topical Anesthetic

SD alcohol, 60% w/v..... Antiseptic

Use: For the temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

Warnings: For external use only.

Avoid contact with eyes. If this happens, rinse thoroughly with water.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Flammable - keep away from fire or flame.

Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling, irritation or other symptoms occur, consult a physician immediately.

Made in CHINA

LOT/EXP:

Insect Sting Relief Pad

Genuine First Aid LLC, Clearwater FL 33755
www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

Active Ingredients

Active Ingredient: .....Bacitracin Zinc 400 units

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

Polymyxin B Sulfate 5000 units

Purpose

Triple Antibiotic

Uses: To help prevent infection in:
minor cuts; scrapes; burns

Warnings

For external use only.

Do not use: in eyes; over large areas of the body;

If allergic to any of the ingredients; for more than one week unless directed by a physician.

Stop use and consult a doctor:

if the condition persists or gets worse; a rash or other allergic reaction develops

Keep out of reach of children.

If ingested, contact a Poison

Control Center right away.

Directions

Directions: clean affected area; apply small amount of this 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

Other information:

Store at room temperature.

Inactive Ingredient
water

Genuine Triple Antibiotic

First Aid Ointment

To Help Prevent Infection

Each Gram Contains:

Bacitracin Zinc 400 units

Neomycin Sulfate 5 mg

(equivalent to 3.5 mg

Neomycin base)

Polymyxin B Sulfate 5000 units

Net Wt. 0.5g ; (1/64 oz)

Manufactured in CHINA for

GENUINE FIRST AID.

Triple Antibiotic Ointment 10pcs

Net wt. 0.9g (1/32oz)

100
Triple Antibiotic

Active Ingredient (in each tablet) Purpose

Acetaminophen 325 mg ..... Analgesic/antipyretic

Uses

temporary relief of minor aches and pains associated with:

common cold; headache; toothache; muscular aches; backache; arthritis; menstrual cramps; and reduction of fever

Warnings:

Liver warning: This product contains acetaminophen.

Severe liver damage may occur if: adult takes more than 12 tablets in 24 hours, which is the maximum daily amount; child takes more than 5 doses in 24 hours; taken with other drugs containing acetaminophen; adult has 3 or more alcoholic drinks every day while using this product

Do not use: with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist; for more than 10 days for pain unless directed by a doctor; for more than 3 days for fever unless directed by a doctor

Ask a doctor before use if the user has liver disease

Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin

Stop use and ask a doctor if: symptoms do not improve; pain gets worse or lasts for more than 10 days; fever gets worse or lasts for more than 3 days; new symptoms occur; redness or swelling is present; a rare sensitivity reaction occurs

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt
medical attention is critical for adults as well as for children even if
you do not notice any signs or symptoms. Do not exceed recommended dosage

Directions

Adults and Children 12 years of age or older	Take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours.
Children 6-11 years of age	Take 1 tablet every 4 to 6 hours as needed. Do not take more than 5 tablets in 24 hours.
Children under 6 years of age	Do not use this regular strength product. This will provide more than the recommended dose (overdose) and could cause serious health problems.

Store at 59-86 degree F (15-30 degree C)

tamper evident sealed packets; do not use any open or torn packets

Distributed by GENUINE FIRST AID
600 Cleveland Str Suite 400, Clearwater, FL 33755

GENUINE FIRST AID 2 Tablets

NON-ASPIRIN

Active ingredient (in each tablet) Purpose

Ibuprofen USP (NSAID*) 200mg . . . . . .Pain reliever/fever reducer

*nonsteroidal anti-inflammatory drug

Uses temporarily relieves minor aches and pains due to:

the common cold

headache

toothache

muscular aches

backache

minor pain of arthritis

menstrual cramps temporarily reduces fever

Warnings

Allergy alert**:** Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include: shock, facial swelling, asthma (wheezing) rash, skin reddening, blisters, hives If an allergic reaction occurs, stop use and seek medical help right away.
Stomach bleeding warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause severe stomach
bleeding. The chance is higher if you: are age 60 or older, have had stomach ulcers or bleeding problems, take a blood thinner (anticoagulant) or steroid drug, take other drugs containing NSAIDs (aspirin, ibuprofen, naproxen, or others), have 3 or more alcoholic drinks every day while using this product, take more or for a longer time than directed

Do not use if you have ever had an allergic reaction to any other pain reliever/fever reducer, right before or after heart surgery.

Ask a doctor before use if stomach bleeding warning applies to you; you have a history of stomach problems such as heartburn; you have a high blood pressure, heart disease, liver cirrhosis, or kidney disease; you are taking a diuretic

Ask a doctor before use if you are:
taking any other drug containing NSAID (prescription or nonprescription); taking aspirin for heart attack or stroke, because Ibuprofen may decrease this benefit of aspirin; taking any other drug

When using this product:
take with food or milk if stomach upset occurs

Stop use and ask a doctor If:
you experience any of the following signs of stomach bleeding; feel faint; vomit blood; have bloody or

black stools; have stomach pain that does get better; pain gets worse or lasts more than 10 days; fever gets worse or lasts more than 3 days; redness or swelling is present in the painful area; any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions:
do not use more than directed; the smallest effective dose should be used; do not take longer than 10 days, unless directed by a doctor.

Adults and Children (12 years and older): Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used. Do not exceed 6 tablets in 24 hours, unless directed by a doctor.

Children under 12 years: Do not give to children under 12 years of age.

Other information: Store at controlled room temperature; avoid excessive heat 40 degree Celsius (104 degree Fahrenheit); tamper evident sealed packets; do not use any opened or torn packets

Inactive ingredients: cellulose, corn starch, fumed silica gel, hypromellose, lactose, magnesium stearate, polydextrose, polyethylene glycol, povidone, silica, sodium starch glycolate, stearic acid, titanium dioxide, triacetin.

Distributed by GENUINE FIRST AID
600 Cleveland Str Suite 400, Clearwater, FL 33755

IBUPROFEN 2 Tablets

IBUPROFEN 2 Tablets

Active Ingredients

ACTIVE INGREDIENTS:

Benzalkonium Chloride 0.13%
Lidocaine HCL 0.5%

Purpose

Purpose: First aid antiseptic, external analgesic

Uses

First aid to help prevent infection and for the temporary relief of pain and itching associated with:

Minor Cuts

Scrapes

Burns

Warnings

For external use only

Do not use: In eyes, in large quantities, over raw blistered areas, or on deep puncture wounds, animal bites or serious burns, for more than one week

Do not use:
in the eyes or apply over large areas of the body.
longer than 1 week unless directed by a doctor.
in large quantities, particularly over raw surfaces or blistered areas.

Ask a doctor before use if you have deep puncture wounds, animal bites or serious burns.

When using this product, avoid contact with the eyes.

Stop use and ask a doctor if
condition worsens
symptoms persist for more than 7 days
condition clears up and occurs again within a few days

Keep out of reach of Children.

If ingested, contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older

clean affected area.

apply a small amount of this product on the area 1 to 3 times daily.

may be covered with a sterile bandage

children under 2 years of age: consult a doctor

Other Information:

Store at room temperature (do not freeze).

Taper evident sealed packets.

Do not use packet if opened or torn.

Inactive Ingredients

Aloe vera, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol, propylparaben, purified water, stearic acid, irolamine

LOT/EXP: Made in CHINA

20130301

GFA Production Xiamen Co., Ltd
No. 20 Huli Industrial Park, Meixi Road, Tong'an, Xiamen, Fujian, China 361100
Tel: 86-592-7269515 Fax: 86-592-7269528 Http: //www.gfaproduction.com

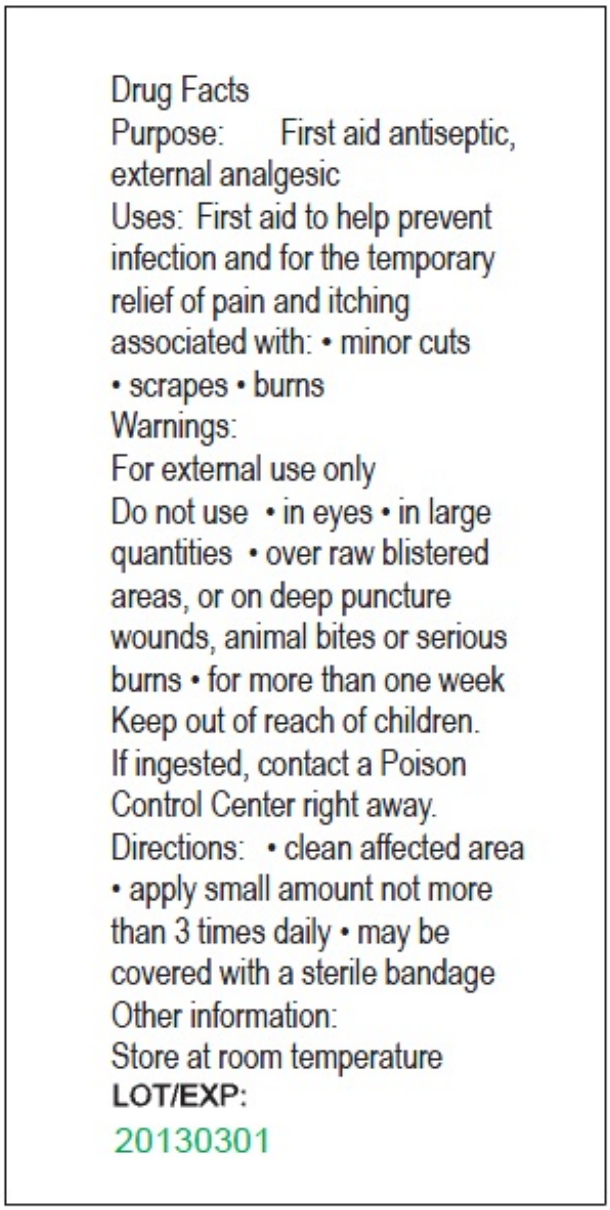
Genuine First Aid Burn Cream

Antiseptic Pain Relief With Aloe

Net Wt 0.9g (1/32 oz)

Manufactured in CHINA for

Genuine First Aid.



Reorder TAO-001

Genuine Triple Antibiotic

First Aid Ointment
To Help Prevent Infection

Each Gram Contains:
Bacitracin Zinc 400 units
Neomycin Sulfate 5 mg
(equivalent to 3.5 mg Neomycin base)
Polymyxin B Sulfate 5000 units
Net Wt 0.5g · (1/64 oz)

Manufactured in CHINA for
GENUINE FIRST AID®

Drug Facts
Uses: To help prevent infection in • minor cuts • scrapes • burns
Warnings:
For external use only
Do not use • in eyes • over large areas of the body • if allergic to any of the ingredients • for more than one week unless directed by a physician
Stop use and consult a doctor • if the condition persists or gets worse • a rash or other allergic reaction develops
Keep out of reach of children. If ingested, contact a Poison Control Center right away.
Directions: • clean affected area • apply small amount of this 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
Other information:
Store at room temperature
LOT/EXP:
20130118

GENUINE FIRST AID. 2 Tablets

NON–ASPIRIN

Active ingredient (in each tablet)	*Purpose*
Acetaminophen 325 mg	Analgesic/antipyretic

Uses

temporary relief of minor aches and pains associated with

■ common cold ■ headache ■ toothache
■ muscular aches ■ backache ■ arthritis
■ menstrual cramps ■ and reduction of fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if ■ adult takes more than 12 tablets in 24 hours, which is the maximum daily amount ■ child takes more than 5 doses in 24 hours ■ taken with other drugs containing acetaminophen ■ adult has 3 or more alcoholic drinks every day while using this product

Do not use ■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist ■ for more than 10 days for pain unless directed by a doctor ■ for more than 3 days for fever unless directed by a doctor

Ask a doctor before use if the user has liver disease

Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin ▶

Warnings (continued)

Stop use and ask a doctor if ■ symptoms do not improve ■ pain gets worse or lasts for more than 10 days ■ fever gets worse or lasts for more than 3 days ■ new symptoms occur ■ redness or swelling is present ■ a rare sensitivity reaction occurs

If pregnant or breast-feeding, ask a health professional before use. **Keep out of reach of children.** In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms. **Do not exceed recommended dosage.**

Directions

Adults and children 12 years of age and older	Take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours.
Children 6-11 years of age	Take 1 tablet every 4 to 6 hours as needed. Do not take more than 5 tablets in 24 hours.
Children under 6 years of age	Do not use this regular strength product. This will provide more than the recommended dose (overdose) and could cause serious health problems.

Other information ■ store at 59°-86°F (15°-30°C) ■ tamper evident sealed packets ■ do not use any open or torn packets

Inactive ingredients corn starch, hydroxypropyl methylcellulose, polyethylene glycol, pregelatinized starch, stearic acid. May contain povidone and sodium starch glycolates.

Distributed by **GENUINE FIRST AID**
600 Cleveland Str Suite 400, Clearwater, FL 33755

REORDER AST-001

Antiseptic Towelette

Toallitas Antisepticas

Genuine First Aid LLC, Clearwater FL 33755
www.GenuineFirstAid.com

CE

ISO 9001:2010

1 /pouch

GENUINE FIRST AID®

2

DRUG FACTS - Antiseptic Towelette

Active Ingredient: Purpose:
Benzalkonium Chloride 0.40%..First Aid Antiseptic
Use: For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.
Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
If unusual redness, swelling or other symptoms occur, consult a physician immediately.
Do not use: In the eyes, or over large areas of the body.
Directions: Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.
Inactive ingredient: Purified water

Made in CHINA

- - - - TEAR HERE - - - -

LOT/EXP:
20130301

GENUINE FIRST AID®

ISO 9001:2010

Genuine First Aid LLC, Clearwater FL 33755
www.GenuineFirstAid.com

Insect Sting Relief Pad

Toallitas para Picaduras de Insectos

REORDER ISRP-001

DRUG FACTS - Insect Sting Relief Pad

Active Ingredient:	**Purpose:**
Benzocaine, 6% w/v	Topical Anesthetic
SD alcohol, 60% w/v	Antiseptic

Use: For the temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Flammable - keep away from fire or flame. Avoid contact with eyes. If this happens, rinse thoroughly with water.

Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling, irritation or other symptoms occur, consult a physician immediately.

Made in CHINA

- - - TEAR HERE - - -

IBUPROFEN 2 Tablets

Active ingredient (in each tablet) **Purpose**
Ibuprofen USP (NSAID*) 200mg . . . . . Pain reliever/fever reducer
*nonsteroidal anti-inflammatory drug

Uses temporarily relieves minor aches and pains due to:
■ the common cold ■ headache ■ toothache ■ muscular aches ■ backache ■ minor pain of arthritis ■ menstrual cramps
temporarily reduces fever

Warnings **Allergy alert:** Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include: ■ shock ■ facial swelling ■ asthma (wheezing) ■ rash ■ skin reddening ■ blisters ■ hives If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause severe stomach bleeding. The chance is higher if you: ■ are age 60 or older ■ have had stomach ulcers or bleeding problems ■ take a blood thinner (anticoagulant) or steroid drug ■ take other drugs containing NSAIDs (aspirin, ibuprofen, naproxen, or others) ■ have 3 or more alcoholic drinks every day while using this product ■ take more or for a longer time than directed

Do not use ■ if you have ever had an allergic reaction to any other pain reliever/fever reducer ■ right before or after heart surgery

Ask a doctor before use if ■ stomach bleeding warning applies to you ■ you have a history of stomach problems such as heartburn ■ you have high blood pressure, heart disease, liver cirrhosis, or kidney disease ■ you are taking a diuretic

Ask a doctor or pharmacist before use if you are ■ taking any other drug containing an NSAID (prescription or nonprescription) ▶

Warnings (continued)
■ taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin ■ taking any other drug

When using this product
■ take with food or milk if stomach upset occurs

Stop use and ask a doctor if ■ you experience any of the following signs of stomach bleeding: ■ feel faint ■ vomit blood ■ have bloody or black stools ■ have stomach pain that does not get better ■ pain gets worse or lasts more than 10 days ■ fever gets worse or lasts more than 3 days ■ redness or swelling is present in the painful area ■ any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions ■ **do not use more than directed** ■ **the smallest effective dose should be used** ■ do not take longer than 10 days, unless directed by a doctor

Adults and Children (12 years and older): Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used. Do not exceed 6 tablets in 24 hours, unless directed by a doctor.

Children under 12 years: Do not give to children under 12 years of age.

Other information ■ store at controlled room temperature ■ avoid excessive heat 40° C (104° F) ■ tamper evident sealed packets ■ do not use any opened or torn packets

Inactive ingredients cellulose, corn starch, fumed silica gel, hypromellose, lactose, magnesium stearate, polydextrose, polyethylene glycol, povidone, silica, sodium starch glycolate, stearic acid, titanium dioxide, triacetin

Distributed by **GENUINE FIRST AID**
600 Cleveland Str Suite 400, Clearwater, FL 33755

EASY CARE FIRST AID - ALL PURPOSE

benzalkonium chloride, benzocaine, sd alcohol, bacitracin zinc, neomycin sulfate, polymyxin b, ibuprofen, acetaminophen kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44224-1999

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:44224-1999-6	1 in 1 KIT		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	20 PACKAGE	16 mL

Part 2	4 PACKAGE	2.0 mL
Part 3	2 TUBE	1.0 g
Part 4	4 PACKAGE	8
Part 5	4 PACKET	8
Part 6	2 PACKET	1.8 g

Part 1 of 6

ANTISEPTIC TOWELETTE

benzalkonium chloride swab

Product Information	
Item Code (Source)	NDC:52124-0001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.4 mL in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52124-0001-1	0.8 mL in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/15/2011	

Part 2 of 6

INSECT STING RELIEF PAD

benzocaine,alcohol swab

Product Information	

Item Code (Source)	NDC:52124-0008
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52124-0008-1	0.5 mL in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	04/15/2011	

Part 3 of 6

GENUINE TRIPLE ANTIBIOTIC

bacitracin zinc,neomycin sulfate,polymyxin b sulfate ointment

Product Information	
Item Code (Source)	NDC:52124-0003
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B SULFATE	5000 [iU] in 1 g

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0003-1	0.5 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	04/15/2011	

Part 4 of 6

NON-ASPIRIN

acetaminophen tablet

Product Information	
Item Code (Source)	NDC:52124-0010
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics			
Color	white (WHITE)	Score	no score
Shape	ROUND	Size	11mm
Flavor		Imprint Code	AZ;234
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0010-1	2 in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	04/15/2011	

Part 5 of 6

IBUPROFEN

ibuprofen tablet

Product Information

Item Code (Source)	NDC:52124-0009
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients

Ingredient Name	Strength
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	white (White)	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	44;352
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0009-1	2 in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075010	02/17/2010	

Part 6 of 6

GENUINE FIRST AID BURN ANTISEPTIC PAIN RELIEF WITH ALOE

benzalkonium chloride, lidocaine cream

Product Information

Item Code (Source)	NDC:52124-0004
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 (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0004-1	0.9 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333A	04/15/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	04/15/2011	

Labeler - Tender Corp dba Adventure Medical Kits (064437304)

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