GENUINE FIRST AID - AUTO FIRST AID - benzalkonium chloride, ibuprofen, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, acetaminophen
Genuine First Aid, LLC

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.

Genuine First Aid Auto First Aid Kit

Active Ingredient: Purpose

Isopropyl Alcohol, 70% v/v..... Antiseptic

Use: For preparation of skin before injection.

Warnings: For external use only.

Flammable - keep away from fire or flame

Store at room temperature 15-30 degree Celsius (59-86 degree Fahrenheit)

Do not use: with electrocautery, in the eyes.

Stop use if irritation and redness develop. If condition persists for more than 72 hours, consult your doctor.

Keep out of reach of children. If swallowed,

get medical help or contact a Poison Control

Center right away.

Wipe Injection site vigorously and discard.

Inactive Ingredient: Purified water.

LOT/EXP: Made in CHINA

20140301

Alcohol Cleansing Pad
Genuine First Aid LLC, Clearwater FL 33755

www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

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

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

Polymyxin B Sulfate 5000 units

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.

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.

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 Ingredient (in each tablet) Purpose

Acetaminophen 500 mg..... Pain Reliever / fever reducer

Purpose: Pain reliever, fever reducer

Uses for the temporary relief of minor aches and pains associated with
headache ; muscular aches ; minor arthritis pain ; toothache ; common cold ; menstrual cramps ; for the reduction of fever

Warnings

Liver Warning: This product contains acetaminophen. Sever liver damage may occur if you take: more than 8 tablets in 24 hours, which is the maximum daily amount; with other drugs containing acetaminophen; 3 or more alcoholic drinks every day while using this product.

Do not use: with any other drug containing acetaminophen (prescription or non prescription). If you are not sure whether a drug contains acetaminiophen, ask a doctor or phramacist.
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 you have liver disease

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin

Stop using and ask a doctor if
symptoms do not improve
new symptoms occur
pain or fever persists or gets worse
redness or swellign is present

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms

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

Adults and children: (12 years and older)
take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

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

Store at 59 - 86 degree Fahrenheit (15 - 30 degree Celsius); Tamper-evident sealed packets. do not use any opened or torn packets Mfd. for MEDIQUE PRODUCTS, FORT MYERS, FL 33967

Inactive Ingredients:Cellulose*, corn starch*, crospovidone*, hydroxypropyl cellulose*,

hypromellose*, magnesium stearate*, microcrystalline cellulose*, mineral oil*, opadry clear*, polyethylene glycol*, polyvinylpyrrolidone*, povidone*, pregelatinized starch*, propylene glycol*, silicon dioxide*, sodium carboxymethylcellulose*, sodium starch glycolate*, starch 1500*, stearic acid, talc*, titanium dioxide*, triacetin*.

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.

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 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

GENUINE FIRST AID®

ISO 9001:2010

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

Antiseptic Towelette
Toallitas Antisepticas

REORDER AST-001

DRUG FACTS - Antiseptic Towelette

Active Ingredient: Benzalkonium Chloride 0.40%..**Purpose:** 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 - - -

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 600609 2015/01

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

REORDER ACP-001

Alcohol Cleansing Pad

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CE

ISO 9001:2010

1 pouch

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

GENUINE FIRST AID®

2

DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient:	Purpose:
Isopropyl Alcohol, 70% v/v	Antiseptic

Use: For preparation of the skin before injection.
Warnings: For external use only.
Flammable - keep away from fire or flame.
Do not use: with electrocautery, in the eyes
Stop use if irritation and redness develop.
If condition persists for more than 72 hours, consult your doctor.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
Directions: Wipe injection site vigorously and discard.
Other information: Store at room temperature 15°-30° C (59°-86° F)
Inactive ingredient: Purified water.

- - - TEAR HERE - - -

LOT/EXP:
20140301

Made in CHINA

Warnings (continued)

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 you have

- liver disease

Ask a doctor or pharmacist before use if

- you are taking the blood thinning drug warfarin

Stop using and ask a doctor if

- symptoms do not improve
- new symptoms occur
- pain or fever persists or gets worse
- redness or swelling is present

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

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

Directions

- **do not use more than directed**

Adults and children: (12 years and older)
Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

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

SEE CARTON FOR COMPLETE PRODUCT INFORMATION

Lot 1857 Exp. 01-2013

Medique®

2 Tablets

Made in USA

Extra Strength APAP

Active Ingredient (in each tablet)	Purpose
Acetaminophen 500 mg.	Pain reliever/fever reducer

Uses

For the temporary relief of minor aches and pains associated with

- headache
- muscular aches
- minor arthritis pain
- toothache
- common cold
- menstrual cramps

For the reduction of fever.

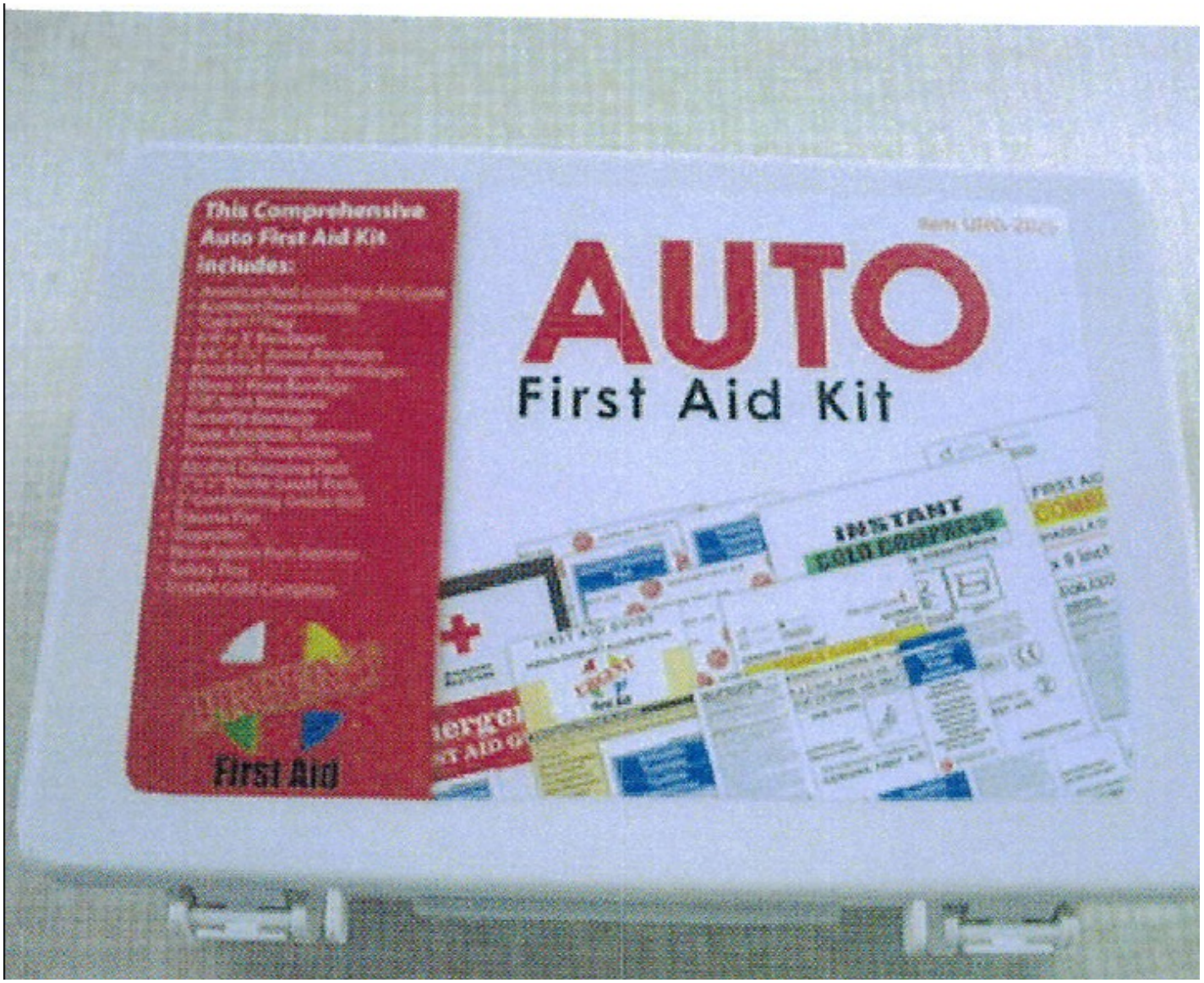
Warnings

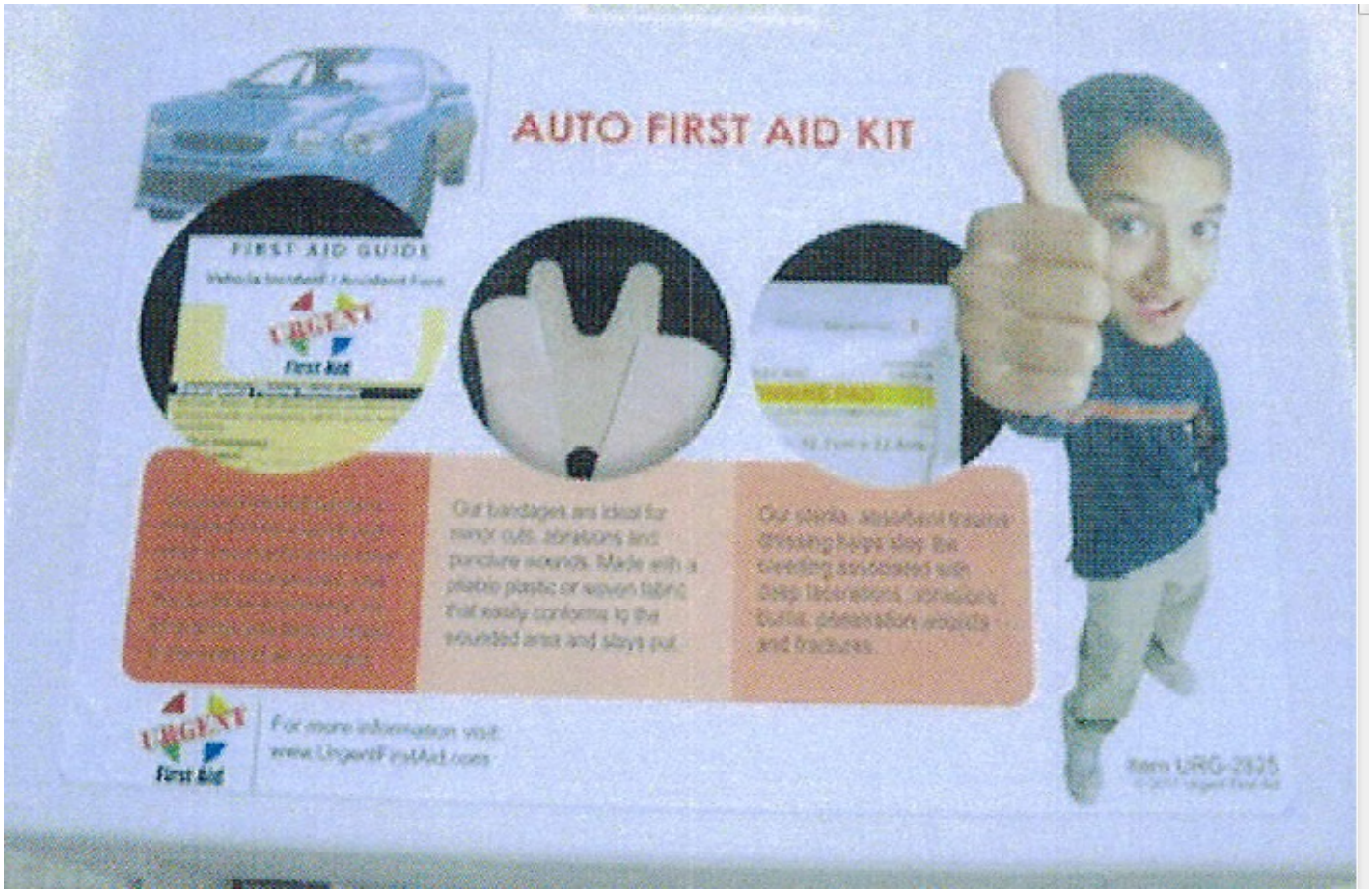
Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Store at 59°-86°F (15°-30°C) • Tamper-evident sealed packets • Do not use any opened or torn packets

Mfd. for MEDIQUE PRODUCTS • Fort Myers, FL 33967


This Comprehensive Auto First Aid Kit Includes:
AUTO
First Aid Kit
First Aid
INSTANT



GENUINE FIRST AID - AUTO FIRST AID

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52124-1000

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52124-1000-1	1 in 1 KIT		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 TUBE	0.5 g
Part 2	1 PACKET	2
Part 3	1 PACKET	2
Part 4	2 PACKAGE	1.6 mL
Part 5	2 PACKAGE	1 mL

Part 1 of 5

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	.5 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	01/11/2011	

Part 2 of 5

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	01/11/2011	

Part 3 of 5

MEDIQUE APAP EXTRA STRENGTH

acetaminophen tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-175
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CROSPOVIDONE (UNII: 68401960MK)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics			
Color	white (WHITE)	**Score**	no score
Shape	ROUND	**Size**	12mm
Flavor		**Imprint Code**	AZ;235
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47682-175-46	2 in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/11/2011	

Part 4 of 5

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	0.40 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 final	part333E	01/11/2011	

Part 5 of 5

ALCOHOL CLEANSING PAD

isopropyl alcohol liquid

Product Information	
Item Code (Source)	NDC:52124-0002
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 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:52124-0002-1	0.5 mL in 1 PACKAGE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/11/2011	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/11/2011	

Labeler - Genuine First Aid, LLC (619609857)

Revised: 1/2011 Genuine First Aid, LLC