ADVENTURE MEDICAL KITS 1-2 PERSON FIRST AID - benzalkonium chloride, aspirin, ibuprofen, isopropyl alcohol, benzocaine, sd alcohol, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, acetaminophen, diphenhydramine chloride 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.

ADVENTURE MEDICAL KITS 1-2 PERSON [1.0] FIRST AID KIT

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: Pu	irpose:
-----------------------	---------

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 ingredient (in each tablet)PurposeIbuprofen USP (NSAID*) 200mgPain 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

Aspirin (NSAID^{*}) 325 mg..... Pain Reliever / fever reducer

*nonsteroidal anti-inflammatory drug

Uses Temporarily relieves minor aches and pains associated with: headache ; muscular aches ; minor arthritis pain ; backache ; common cold ; toothache ; menstrual cramps ; Temporarily reduces fever

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox of flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include: hives, skin reddening, facial swelling, rash, asthma (wheezing), blisters, shock, If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: This contains an 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 prescription or nonprescription 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; if you are taking a prescription drug for gout, diabetes or arthritis

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:

under a doctor's care for any serious condition; 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; you have difficulty swallowing; if ringing in the ears or loss of hearing occurs; redness or swelling is present in the painful areas; any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important to use aspirin 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 drink a full glass of water with each dose do not take longer than 10 days, unless directed by a doctor

Adults and children: (12 years and older) Take 1 or 2 tablets with water every 4 hours as needed. Do not take more than 12 tablets in 24 hours, or as directed by a doctor. Children under 12 years: Do not give to children under 12 years of age. Store at 59 - 86 degree Fahrenheit (15 - 30 degree Celsius); avoid excessive heat and humidity; tamper evident sealed packets; Do not use any opened or torn packets Inactive Ingredients: hypromellose, polyethere glycol, propylene glycol, corn starch Oistributed by GENUINE FIRST AID 600 Clevelad Str Suite 400, Clearwater, FL 33755 ASPIRIN 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 (in each tablet) Purpose

Diphenhydramine Hydrochloride 25mg...... Antihistimine

Purpose: Antihistimine

Uses Temporarily relieves the following symptoms associated with hay fever or other upper respiratory allergies:

runny nose, sneezing, itching of the nose or throat, itchy, watery eyes

Store at room temperature. Tamper-evident sealed packets. do not use any opened or torn packets Mfd. for MEDIQUE PRODUCTS, FORT MYERS, FL 33967

Warnings

Ask a doctor before use if you have: a breathing problem such as emphysema or chronic bronchitis glaucoma difficulty in urination due to enlargement of the prostate gland

if you are: taking any drugs for asthma taking sedatives or tranquilizers

When using this product

Drowsiness may occur

avoid alcoholic beverages

alcohol, sedatives, and tranquilizers may increase the drowsiness effect

use caution when driving a motor vehicle or operating machinery

excitaility may occur, especially in children

Do not exceed recommended dosage. Keep this and all drugs out of the reach of children. In case of accidental overdose, contact a physician or posion control center immediately. As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using this product.

Adults and children: (12 years and older)

take 1 capsule every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours, or as directed by a doctor.

Children under 12 years: do not give to children under 12 years of age unless directed by a doctor.

Inactive Ingredients: DandC Red 28, FDandC Blue 1, FDandC Red 40, gelatin, starch

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.

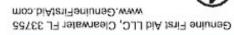
Inactive Ingredient: Purified water

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.

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REORDER AST-001

DRUG FACTS - Antiseptic Towelette

Active Ingredient:	Purpose:	1
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 wate	r.	1
Warnings: For external use only. Keep	o out of	ш
reach of children. If swallowed, get me	dical help	R
or contact a Poison Control Center righ	t away.	ш
If unusual redness, swelling or other sy		I
occur, consult a physician immediately.		
Do not use: In the eyes, or over large a	areas of the	TEAR
body.		4
Directions: Tear open packet, unfold to	owelette	ш
and use to cleanse desired skin area.	Discard	-
towelette appropriately after single use.		
Inactive ingredient: Purified water.		1
•	2010/2010/201	T.
Made	in CHINA	



33755 First bid LLC, Clearwater FL 33755 moo.biAtariBeniunaD.www



DRUG FACTS - Insect Sting Relief Pad

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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_®

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)

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

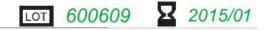
• apply small amount of this product (an amount equal to the surface

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



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 up 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

Active ingredient (in each tablet) Purpose

Uses Temporarily relieves minor aches and pains associated with ■ headache ■ muscular aches ■ minor arthritis pain ■ backache ■ common cold ■ toothache ■ menstrual cramps Temporarily reduces fever

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include: ■ hives ■ skin reddening ■ facial swelling ■ rash asthma (wheezing) blisters shock

If an allergic reaction occurs, stop use and seek medical help right away. Stomach bleeding warning: This product contains an 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 prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others) have 3 or more alcoholic drinks every day while using this product I 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 ■ if you are taking a prescription drug for gout, diabetes or arthritis

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

Warnings (continued)

Ask a doctor or pharmacist before use if you are

■ under a doctor's care for any serious condition ■ taking any other drug When using this product I take with food or milk if stomach upset occurs Stop use and ask a doctor if

y ou 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 ■ you have difficulty swallowing ■ if ringing in the ears or loss of hearing occurs ■ 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 aspirin 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

drink a full glass of water with each dose do not take longer than 10 days, unless directed by a doctor

Adults and children: (12 years and older) Take 1 or 2 tablets with water every 4 hours as needed. Do not take more than 12 tablets in 24 hours, or as directed by a doctor.

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

Other information ■ store at 59°-86°F (15°-30°C) ■ avoid excessive heat and humidity a tamper evident sealed packets do not use any opened or torn packets

Inactive ingredients hypromellose, polyethylene glycol, propylene glycol, corn starch

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

	18
Warnings (continued) Do not use	
 with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure wheth a drug contains acetaminophen, ask a doctor or pha.mac for more than 10 days for pain unless directed by a doc for more than 3 days for fever unless directed by a doc 	ast.
Ask a doctor before use If you have m 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 a do not use more than directed	1 1-10
Adults and children: (12 years and older) Take 2 lablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.	9
Children under 12 years: Do not give to children under 12 years of age.	

SEE CARTON FOR COMPLETE PRODUCT INFORMATION

2 Tablets

Active Ingredient (in each tablet) Purpose Acetaminophen 500 mg. Pain relieventiever reducer

Médique

Extra Strength

Uses

For the temporary relief of minor aches and paine associated with

headacha muscular aches

minor artuntis pain

menstrual cramps

Made in

IS A

For the reduction of fever.

Il toothacha III common coid

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 tom packets and, for MEDIQUE PRODUCTS • Fort Myers, FL 33967



Warnings Ask a Doctor Before Use If you have: · a breathing problem such as emphysema or chronic bronchies diaucoma · difficulty in urination due to enlargement of the prostate pland H vou are: · taking any drugs for asthma · taking sedatives or tranguilizers When Using this product: · avoid alcohol beverages · drowsiness may occur · alcohol, sedatives and tranquilizers may increase the drowsiness effect use caution when driving a motor vehicle or operating machinery · excitability may occur, especially in children. Do not exceed recommended dosage. Keep this and all drugs out of the reach of children. In case of accidental overdose, contact a physician or poison control center immediately. As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using this product. Directions: Do not use more than directed Adults and children: (12 years and older) Take 1 capsule every 4 to 6 hours as needed. Do not take more than 12 capsules in 24 hours, or as directed by a doctor. Children under 12 years: Do not give to children under 12 years of age unless directed by a doctor.

Lot 3951 Exp. 11-2011





ADVENTURE MEDICAL KITS 1-2 PERSON FIRST AID

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

Product	Information			
Product T	Гуре	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-2000
Packagi	ng			
# It	em Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC ·442	224-2000-1	1 in 1 KIT		
1 NDC.442				
	of Parts			
Quantity		ckage Quantity	Total Product	t Quantity
Quantity Part #	Pa	ckage Quantity	Total Product	t Quantity
Quantity Part # Part 1 2 T	Pa	ckage Quantity		t Quantity
Quantity Part # Part 1 2 T Part 2 2 P	Pa CUBE	ckage Quantity	1 g	t Quantity
Quantity Part # Part 1 2 T Part 2 2 P Part 3 4 P	Pae TUBE PACKAGE	ckage Quantity	1 g 1 mL	t Quantity
Quantity Part # Part 1 2 T Part 2 2 P Part 3 4 P Part 4 2 P	Pac TUBE PACKAGE PACKET	ckage Quantity	1 g 1 mL 8	t Quantity

Part 1 of 7

GENUINE TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

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

Ingredient Name	Basi	s of Strength	Strength
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	BACITR	ACIN ZINC	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMY	CIN SULFATE	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYM	YXIN B SULFATE	5000 [iU] in 1 g
Inactive Ingredients			
Ingredient Name		Stre	ngth
			0
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 Info	rmation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	08/26/2010	

Part 2 of 7

INSECT STING RELIEF PAD

benzocaine, alcohol swab

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

Active Ingredient/A	Active Moi	ety					
	Ingre	dient Name		В	asis of Strengt	h	Strength
BENZOCAINE (UNII: U31	RS Y48 JW5) (B	ENZOCAINE - UNII:U3RS	SY48JW5)	BEI	NZOCAINE		in 100 mL
ALCOHOL (UNII: 3K995	8V90M) (ALC	OHOL - UNII:3K9958V9	0 M)	AL	COHOL	60 m	L in 100 mL
Inactive Ingredient	ts						
	I	ngredient Name				Streng	th
WATER (UNII: 059QF0K	O0R)						
Packaging							
# Item Code	Pacl	kage Description	Marketin	ng Start I)ate Ma	rketing	End Date
1 NDC:52124-0008-1		1 PACKAGE		ig sturt i			
Marketing Info	rmation						
Marketing Category	Applicatio	on Number or Monogra	aph Citation	Market	ing Start Date	Market	ing End Date
OTC monograph final	part348			08/26/20	10		
Part 3 of 7							
IBUPROFEN							
ibuprofen tablet							
D 1 1 1							
Product Information	n						
Item Code (Source)		NDC:52124-0009					
Route of Administratio	on	ORAL					
Active Ingredient/A	Active Moi	ety					
	Ing	redient Name			Basis of Str	ength	Strength
IBUPROFEN (UNII: WK2)	XYI10QM) (IBU	UPROFEN - UNII:WK2XYI	10 Q M)		IBUPROFEN		200 mg
Inactive Ingredient	ts						
		Ingredient Name				S	trength
POWDERED CELLULO	SE (UNII: SMD	-					
STARCH, CORN (UNII: C)8232NY3SJ)						
HYPROMELLOSES (UN	II: 3NXW29V3	WO)					
LACTOSE (UNII: J2B2A4	4N98G)						
MAGNESIUM STEARAT	E (UNII: 70097	7M6I30)					

POLYETHYLENE GLYCO						
DO LUDO NE (LIDIU EROCO)		JQ0SDW1A)				
POVIDONE (UNII: FZ9890	GH94E)					
SILICON DIOXIDE (UNII:	ETJ7Z6XBU	4)				
STEARIC ACID (UNII: 4EL	V7Z65AP)					
TITANIUM DIO XIDE (UN	II: 15FIX9V2J	P)				
TRIACETIN (UNII: XHX3C	3X673)					
Product Characteris	stics					
Color	white (WHI	TE)	Score		no score	
Shape	ROUND		Size		10 mm	
Flavor			Imprint Code		44;352	
Contains						
Packaging						
# Item Code	Pack	kage Description	Marketing Star	t Date Ma	arketing En	d Date
1 NDC:52124-0009-1	2 in 1 PAC	CKET				
Marketing Infor	mation					
Marketing Category	Applicatio	n Number or Monogra	ph Citation Mark	eting Start Date	Marketing	End Date
ANDA A	ANDA075010		08/26/	2010		
Part 4 of 7						
	IEN					
MEDIQUE DIPH		nsule				
		apsule				
MEDIQUE DIPH		ıpsule				
MEDIQUE DIPH	ochloride ca	ıpsule				
MEDIQUE DIPH diphenhydramine hydro Product Information	ochloride ca	npsule NDC:47682-182				
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source)	ochloride ca	-				
MEDIQUE DIPH diphenhydramine hydro Product Information	ochloride ca	NDC:47682-182				
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source)	ochloride ca	NDC:47682-182				
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration	n n	- NDC:47682-182 ORAL				
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source)	n n ctive Moie	NDC:47682-182 ORAL		Basis of Stu	rength	Strength
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration	n n ctive Moie Ingre	NDC:47682-182 ORAL ety edient Name	(DIPHENHYDRAMINE -	Basis of Stu DIPHENHYDRAMIN	-	Strength
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A	n n ctive Moie Ingre	NDC:47682-182 ORAL ety edient Name	(DIPHENHYDRAMINE -		-	Strength 25 mg
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A DIPHENHYDRAMINE HYD	n n ctive Moie Ingre	NDC:47682-182 ORAL ety edient Name	(DIPHENHYDRAMINE -	DIPHENHYDRAMIN	-	
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A DIPHENHYDRAMINE HYT UNII:8 GTS82S83M)	ochloride ca n ctive Moio Ingro DRO CHLO R	NDC:47682-182 ORAL ety edient Name	(DIPHENHYDRAMINE -	DIPHENHYDRAMIN	-	
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A DIPHENHYDRAMINE HYD	ochloride ca n ctive Moio Ingro DRO CHLO R	NDC:47682-182 ORAL ety edient Name IDE (UNII: TC2D6JAD40)	(DIPHENHYDRAMINE -	DIPHENHYDRAMIN	Ε	25 mg
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A DIPHENHYDRAMINE HYT UNII:8 GTS82S83M)	ochloride ca 1 n ctive Moie Ingre DROCHLOR	NDC:47682-182 ORAL ety edient Name	(DIPHENHYDRAMINE -	DIPHENHYDRAMIN	-	25 mg
MEDIQUE DIPH diphenhydramine hydro Product Information Item Code (Source) Route of Administration Active Ingredient/A DIPHENHYDRAMINE HYT UNII:8 GTS82S83M)	ochloride ca n ctive Moie Ingre DROCHLOR 3 327L)	NDC:47682-182 ORAL ety edient Name IDE (UNII: TC2D6JAD40)	(DIPHENHYDRAMINE -	DIPHENHYDRAMIN	Ε	25 mg

FD&C BLUE NO.1 (U	JNII: H3R47K3TB	D)					
FD&C RED NO. 40 (UNII: WZB9127XOA)							
STARCH, CORN (UNII: 08232NY3SJ)							
Product Charact	eristics						
Color	pink (PINK) , wh	ite (WHITE)		Score		no so	ore
Shape	CAPSULE			Size		14mm	1
Flavor				Imprint Code	!	CPC;	835
Contains							
Packaging							
# Item Code	Pac	kage Description	Marketin	ng Start Date	Ma	arketing E	End Date
1 NDC:47682-182-46	1 in 1 PAC			0		0	
Mauliating Ind							
Marketing Inf							
Marketing Categor		on Number or Monogra	ph Citation	Marketing S	tart Date	Marketiı	ng End Date
OTC monograph final	part341			08/25/2010			
Part 5 of 7							
MEDIOUE AI	PAP EXTR	A STRENGTH					
acetaminophen tab							
		•					
Product Informa							
Item Code (Source))	NDC:47682-175					
Route of Administra	ation	ORAL					
Active Ingredier	nt/Active Moi	ety					
	In	gredient Name			Basis of S	Strength	Strength
ACETAMINOPHEN (UNII: 36209 ITL9D) (ACETAMINOPHEN - UNII:36209 ITL			NII:362O9ITL9	D) .	ACETAMINO	_	500 mg
							_
Inactive Ingredie	ents						
		Ingredient Nam	1e			5	Strength
CROSPOVIDONE (U	NII: 68401960MK	-					0
HYDROXYPROPYL							
HYPROMELLOSES (
MAGNESIUM STEAR							
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)							
MINERAL OIL (UNII: T5L8T28FGP)							
POLYETHYLENE GL	YCOL (UNII: 3W	/JQ0SDW1A)					

POVIDONE (UNII: FZ989GH94E)							
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)							
	SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)						
CARBOXYMETHYLCELLU		DIUM (UNII: K679OBS311)				
STEARIC ACID (UNII: 4ELV	/7Z65AP)						
TALC (UNII: 7SEV7J4R1U)							
TITANIUM DIO XIDE (UNII:		P)					
TRIACETIN (UNII: XHX3C3	X673)						
Product Characterist	tics						
Color	white (WHI	TE)	Score		n	o score	
Shape	ROUND		Size		1	mm	
Flavor			Imprint Code		A	Z;235	
Contains							
Packaging							
# Item Code	Pacl	kage Description	Marketing S	tart Date	Marl	eting End Da	te
1 NDC:47682-175-46	2 in 1 PAC		Walketing 5	tart Date			ii.
I NDC.47082-175-40	2 III 1 FAC	JKE I					
Marketing Information							
88							
Marketing Category		ion Number or Monogr	aph Citation M	larketing Star	t Date I	Aarketing End	Date
Marketing Category		ion Number or Monogr	-	Marketing Star 3/26/2010	t Date I	Marketing End	Date
Marketing Category	Applicat	ion Number or Monogr	-	-	t Date I	Marketing End	Date
Marketing Category	Applicat	ion Number or Monogr	-	-	t Date 🛛	Marketing End	Date
Marketing Category	Applicat	ion Number or Monogr	-	-	t Date I	Marketing End	Date
Marketing Category OTC monograph not final Part 6 of 7	Applicat	ion Number or Monogr	-	-	t Date I	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN	Applicat	ion Number or Monogr	-	-	t Date I	Marketing End	Date
Marketing Category OTC monograph not final Part 6 of 7	Applicat	ion Number or Monogr	-	-	t Date I	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN	Applicat	ion Number or Monogr	-	-	t Date I	Marketing End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet	Applicat	ion Number or Monogr	-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN	Applicat	ion Number or Monogr	-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet	Applicat	ion Number or Monogr NDC:52124-0011	-	-	t Date	Marketing End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information	Applicat part343		-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source)	Applicat part343	NDC:52124-0011	-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source)	Applicat part343	NDC:52124-0011	-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source)	Applicat part343	NDC:52124-0011 ORAL	-	-	t Date	Marke ting End	Date
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	Applicat part343	NDC:52124-0011 ORAL	-	3/26/2010			
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	Applicat part343	NDC:52124-0011 ORAL ety dient Name	-	3/26/2010	f Strengt		
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	Applicat part343	NDC:52124-0011 ORAL ety dient Name	-	B/26/2010 Basis of		h Stren	
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	Applicat part343	NDC:52124-0011 ORAL ety dient Name	-	B/26/2010 Basis of		h Stren	
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration Active Ingredient/Act	Applicat part343	NDC:52124-0011 ORAL ety dient Name	-	B/26/2010 Basis of		h Stren	
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	Applicat part343	NDC:52124-0011 ORAL ety dient Name N - UNII:R16CO5Y76E)	-	B/26/2010 Basis of		h Stren 325 mg	ngth
Marketing Category OTC monograph not final Part 6 of 7 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration Active Ingredient/Act	Applicat part343	NDC:52124-0011 ORAL ORAL ety dient Name N - UNII:R16CO5Y76E) Ingredient Name	-	B/26/2010 Basis of		h Stren	ngth

POLYETHYLENE GLYC	OL (UNII: 3W	JQ0SDW1A)				
PROPYLENE GLYCOL (UNII: 6 DC9Q167V3)						
STARCH, CORN (UNII: O	8232NY3SJ)					
Product Characteri						
Color	white (white)				score	
Shape	ROUND	Siz				
Flavor		Imp	orint Code	44	;157;ASPIRIN	
Contains						
Packaging						
# Item Code	Pack	age Description	Marketin	g Start Date	Marke	ting End Date
1 NDC:52124-0011-1	2 in 1 PAC			g otart Date	IVINI KC	Life Dute
Marketing Infor	mation					
Marketing Infor						
Marketing Category		on Number or Monog	graph Citation	Marketing Sta	rt Date Ma	rketing End Date
OTC monograph final	part343			08/26/2010		
Part 7 of 7						
ANTISEPTIC T	OWELE	TTE				
benzalkonium chloride	e swab					
Product Informatio	n					
Item Code (Source)		NDC:52124-0001				
. ,						
Route of Administration	n	TOPICAL				
Active Ingradiant/	etivo Moi	o.t				
Active Ingredient/A		-		Dasia	f Stuon oth	Strongth
BENZALKONIUM CHLO	•	edient Name		BENZALK	of Strength	Strength 0.40 mL
UNII:7N6JUD5X6Y)			ALKONIOM -	CHLORIDE		in 100 mL
Inactive Ingredient	S					
	I	ngredient Name			St	rength
WATER (UNII: 059QF0K00R)						
Packaging						
# Item Code	Pack	age Description	Marketin	g Start Date	Market	ting End Date

1 NDC:52124-0001-1	0.8 mL in 1 PACKAGE		
Marketing Info	rmation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333E	09/09/2010	
Marketing Info	rmation		
Marketing Category	Application Number or Monograph Citation	n Marketing Start Date	Marketing End Date
OTC monograph not final	part333B	08/26/2010	

Labeler - Tender Corp dba Adventure Medical Kits (064437304)

Registrant - GFA Production (Xiamen) Co., Ltd. (421256261)

Establishment					
Name	Address	ID/FEI	Business Operations		
GFA Production (Xiamen) Co., Ltd.		421256261	manufacture		

Revised: 1/2011

Tender Corp dba Adventure Medical Kits