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

ADVENTURE MEDICAL KITS 1-2 PERSON NATIONAL SKI PATROL 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:
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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: 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.

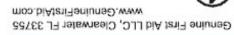
Inactive Ingredient: Purified water

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
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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

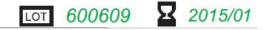
• apply small amount of this product (an amount equal to the surface

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

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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 1-10H
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

ISA

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





ADVENTURE MEDICAL KITS 1-2 PERSON NATIONAL SKI PATROL FIRST AID

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

Pro	duct Information			
Prod	luct T yp e	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-2500
Dad				
Pacl	kaging			
Pacl #	kaging Item Code	Package Description	Marketing Start Date	Marketing End Date
#		Package Description 1 in 1 KIT	Marketing Start Date	Marketing End Date

Part # Pa	ckage Quantity			Tota	al Prod	uct Quant	tity	
Part 1 2 TUBE		1	l g					
Part 2 2 PACKAGE		1	l mL					
Part 3 4 PACKET		8	3					
Part 4 2 PACKET		4	1					
Part 5 2 PACKAGE		4	1					
Part 6 10 PACKAGE		8	3 mL					
Part 1 of 6								
GENUINE TRIP bacitracin zinc, neomyc			intment					
Product Information	I							
Item Code (Source)	NDC:52	2124-0003						
Route of Administration	TOPICA	AT						
Active Ingredient/Ac	ctive Moiety							
Active Ingredient/Ac	Ingredient					of Streng	gth	Strength
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BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO)	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI : (UNII: 19371312D4) (Ingredie 0R)	ITRACIN - UNII:: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9	6K)	BACITRA NEO MYO PO L YMY	ACIN ZINC CIN SULFAT XIN B SULF	TE FATE Stre	400 [iU] in 5 mg in 1 g 5000 [iU] i
BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO)	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI : (UNII: 19371312D4) (Ingredie 0R)	ITRACIN - UNII:: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9	6K)	BACITRA NEO MYO PO L YMY	ACIN ZINC CIN SULFAT XIN B SULF	TE FATE Stre	400 [iU] in 1 5 mg in 1g 5000 [iU] in 1
BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO) Packaging # Item Code 1 NDC:52124-0003-1	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI : (UNII: 19371312D4) (Ingredie 0R) Package De .5 g in 1 TUBE	ITRACIN - UNII:: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9	6K)	BACITRA NEO MYO PO L YMY	ACIN ZINC CIN SULFAT XIN B SULF	TE FATE Stre	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1
BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO) Packaging # Item Code 1 NDC:52124-0003-1	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI : (UNII: 19371312D4) (Ingredie 0R) Package De .5 g in 1 TUBE	ITRACIN - UNII: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9 Marketin	6K)	BACITRA NEO MYO PO L YMY	ACIN ZINC CIN SULFAT XIN B SULF Mai	TE FATE Stre	400 [iU] in 1 g 5 mg in 1 g 5000 [iU] in 1
BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO) Packaging # Item Code 1 NDC:52124-0003-1 Marketing Infori Marketing Category	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI : (UNII: 19371312D4) (Ingredie 0R) Package De .5 g in 1 TUBE	ITRACIN - UNII: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9 Marketin	6K)	BACITRA NEO MYO POLYMY Date	ACIN ZINC CIN SULFAT XIN B SULF Mai	TE FATE Stre	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1 ngth
BACITRACIN ZINC (UNII: NEOMYCIN SULFATE (UN POLYMYXIN B SULFATE Inactive Ingredients WATER (UNII: 059QF0KO) Packaging # Item Code 1 NDC:52124-0003-1 Marketing Infori Marketing Category	Ingredient I 89 Y4M234ES) (BAC NII: 057Y626693) (NI (UNII: 19371312D4) (Ingredie 0R) Package De .5 g in 1 TUBE	ITRACIN - UNII: EOMYCIN - UNII (POLYMYXIN B ent Name	:116 QD7X297) - UNII:J2VZ07J9 Marketin	6K) g Start Marke	BACITRA NEO MYO POLYMY Date	ACIN ZINC CIN SULFAT XIN B SULF Mai	TE FATE Stre	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1 ngth

benzocaine,alcohol swab					
Product Information					
Item Code (Source)	NDC:52124-0008				
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	etv				
-	dient Name		Basis of	Strength	Strength
BENZOCAINE (UNII: U3RS Y48 JW5) (B		Y48JW5)	BENZOCAL	-	6 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M) (ALC			ALCOHOL		60 mL in 100 mL
Inactive Ingredients					
I	ngredient Name			St	trength
WATER (UNII: 059QF0KO0R)					
Packaging					
	kage Description	Marketing	Start Date	Marke	ting End Date
	1 PACKAGE	i mini ne ting	Start Date	WAIIN	Ling Ling Dutt
Marketing Information					
Marketing Category Application	on Number or Monogra	ph Citation	Marketing Sta	rt Date Ma	arketing End Date
OTC monograph final part348		0	8/26/2010		
Part 3 of 6					
IBUPROFEN					
ibuprofen tablet					
1					
Product Information					
Item Code (Source)	NDC:52124-0009				
Route of Administration	ORAL				
Active Ingredient/Active Moi	ety				
	redient Name		Bas	is of Streng	th Strength
IBUPROFEN (UNII: WK2XYI10QM) (IB	UPROFEN - UNII:WK2XYI	10 Q M)	IBUPRO	DFEN	200 mg

		Ingredient Name				Sti	rength
POWDERED CELLULOSE	(UNII: SMD	1X3XO9M)					
STARCH, CORN (UNII: 082	232NY3SJ)						
HYPROMELLOSES (UNII:	3NXW29V3	WO)					
LACTOSE (UNII: J2B2A4NS	98G)						
MAGNESIUM STEARATE ((UNII: 7009	7M6I30)					
POLYDEXTROSE (UNII: V	H2XOU12IE)					
POLYETHYLENE GLYCO	L (UNII: 3W	JQ0SDW1A)					
POVIDONE (UNII: FZ989G	,						
SILICON DIO XIDE (UNII: E		(4)					
STEARIC ACID (UNII: 4EL)							
TITANIUM DIO XIDE (UNII		P)					
TRIACETIN (UNII: XHX3C3	X673)						
Product Characterist							
Color	white (WH	TE)	Score			no score	
Shape	ROUND		Size			10 mm	
Flavor			Imprint Co	de		44;352	
Contains							
Packaging							
# Item Code	Pac	kage Description	Marketin	ng Start Date	Mar	rketing E	and Date
1 NDC:52124-0009-1	2 in 1 PAC	CKET					
Marketing Inform	nation						
Marketing Category	Applicatio	on Number or Monogra	ph Citation	Marketing Start	t Date	Marketin	ıg End Date
	Applicatio NDA075010		ph Citation	Marketing Start	t Date	Marketin	ıg End Date
			ph Citation	_	t Date	Marketin	ng End Date
			ph Citation	_	t Date	Marketir	ng End Date
			ph Citation	_	t Date	Marketin	ng End Date
ANDA A	NDA075010		ph Citation	_	t Date	Marketir	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f Product Information	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f Product Information Item Code (Source)	NDA075010 • EXTR	A STRENGTH	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f Product Information Item Code (Source) Route of Administration	NDA075010 EXTR ilm coated	A STRENGTH NDC:47682-175 ORAL	ph Citation	_	t Date	Marketin	ng End Date
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f Product Information Item Code (Source)	NDA075010 EXTR ilm coated	A STRENGTH NDC:47682-175 ORAL	ph Citation	08/26/2010			
ANDA A Part 4 of 6 MEDIQUE APAP acetaminophen tablet, f Product Information Item Code (Source) Route of Administration	DA075010 EXTR ilm coated tive Moi In	A STRENGTH NDC:47682-175 ORAL ety gredient Name		08/26/2010 Ba	t Date sis of St	trength	ng End Date

Inactive Ingredients	5				
0	Ingredient Nan	ne			Strength
CROSPOVIDONE (UNII: 6					
,	LULOSE (UNII: RFW2ET671P)				
HYPROMELLOSES (UNII					
MAGNESIUM STEARATE					
	YSTALLINE (UNII: OP1R32D61U)				
MINERAL OIL (UNII: T5L8					
POLYETHYLENE GLYCO					
POVIDONE (UNII: FZ9890	GH94E)				
PROPYLENE GLYCOL (U	JNII: 6DC9Q167V3)				
SILICON DIO XIDE (UNII:					
	LULOSE SODIUM (UNII: K679OBS31	11)			
STEARIC ACID (UNII: 4EL					
TALC (UNII: 7SEV7J4R1U))				
TITANIUM DIO XIDE (UNI					
TRIACET IN (UNII: XHX3C					
Product Characteris	stics				
Color	white (WHITE)	Score			no score
Shape	ROUND	Size			12mm
Flavor		Imprint Cod	le		AZ;235
Contains					
Packaging	Package Description	Maximuting	a Start Data	Max	diating End Data
Packaging # Item Code	Package Description	Marketing	g Start Date	Mai	keting End Date
 Packaging # Item Code 1 NDC:47682-175-46 	2 in 1 PACKET	Marketinş	g Start Date	Maı	keting End Date
Packaging # Item Code 1 NDC:47682-175-46	2 in 1 PACKET mation				
 Packaging # Item Code 1 NDC:47682-175-46 	2 in 1 PACKET		g Start Date Marketing Star		keting End Date Marketing End Date
Packaging # Item Code 1 NDC:47682-175-46	2 in 1 PACKET mation				
Packaging # Item Code 1 NDC:47682-175-46 Marketing Infor	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 NDC:47682-175-46 Narketing Infor Narketing Category OTC monograph not final	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 Marketing Information of the second of the	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 Parketing Information of the second	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 Marketing Information of the second of the	2 in 1 PACKET mation Application Number or Monog		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 Marketing Information of the second of the	2 in 1 PACKET mation Application Number or Monog part343		Marketing Star		
Packaging I Item Code I NDC:47682-175-46 INDC:47682-175-46 INDC:47	2 in 1 PACKET Mation Application Number or Monog part343		Marketing Star		
Packaging # Item Code 1 NDC:47682-175-46 Marketing Information of the second of the	2 in 1 PACKET mation Application Number or Monog part343		Marketing Star		

Active Ingredient/A	Active Moie	ety					
		dient Name		Ba	sis of Stren	gth	Strength
ASPIRIN (UNII: R16CO5Y		N - UNII:R16CO5Y7	76E)	ASPIRI	N	_	325 mg
Inactive Ingredient	ts						
		Ingredient	Name			5	Strength
HYPROMELLOSES (UN							
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A) PROPYLENE GLYCOL (UNII: 6DC9Q167V3)							
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) STARCH, CORN (UNII: 08232NY3SJ)							
Product Characteri	istics						
Color	white (white)		Score		no score		
Shape	ROUND		Size		11mm		
Flavor			Imprint Code		44;157;ASPI	RIN	
Contains							
Packaging							
# Item Code		age Description	n Marketin	ig Start Date	e Ma	rketing	g End Date
1 NDC:52124-0011-1	2 in 1 PACI	KAGE					
Marketing Infor	rmation						
Marketing Category	Applicatio	on Number or Mo	onograph Citation	Marketing	Start Date	Marke	ting End Date
OTC monograph final	part343			08/26/2010			
Part 6 of 6							
ANTISEPTIC T	OWFLF	TTF					
benzalkonium chloride		112					
	e swab						
Product Informatio							
	n						
Item Code (Source)		NDC:52124-0001					
Route of Administration	on	TOPICAL					
A T							
Active Ingredient/A		-					
	_	dient Name			sis of Streng		Strength
BENZALKONIUM CHLC	ORIDE (UNII: F	5UM2KM3W7) (BI	ENZALKONIUM -	BENZA	ALKONIUM	0).40 mL

UNII:7N6JUD5X6Y)			CHLORIDE		in 100 mL
The second transmission					
Inactive Ingredients					
	Ingredient Name				Strength
WATER (UNII: 059QF0KO	0 R)				
Packaging					
# Item Code	Package Description	Marketin	g Start Date	Ma	rketing End Date
1 NDC:52124-0001-1	0.8 mL in 1 PACKAGE				
Marketing Inform	mation				
Marketing Category	Application Number or Monogra	aph Citation	Marketing Start	t Date	Marketing End Date
OTC monograph final p	part333E		09/09/2010		
Marketing Inform	mation				
Marketing Category	Application Number or Monog	raph Citation	Marketing Star	t 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: 11/2010

Tender Corp dba Adventure Medical Kits