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

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

Easy Care First Aid Kit Outdoor and Travel

Active Ingredient

Active Ingredient: Benzalkonium Chloride 0.40%

Purpose

Antiseptic

Use

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

Warnings

Warning: For external use only.

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

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

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

Directions

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

Inactive Ingredients

Inactive Ingredient: Purified water

LOT/EXP: Made in CHINA 20130301 Antiseptic Towelette Genuine First Aid LLC, Clearwater FL 33755 www.GenuineFirstAid.com 1/pouch GENUINE FIRST AID Active Ingredient: **Purpose: Topical** Anesthetic Benzocaine, 6% w/v.... SD alcohol, 60% w/v..... Antiseptic Use: For the temporary relief of pain and itching associated with minor burns, scrapes and insect bites. Warnings: For external use only. Avoid contact with eyes. If this happens, rinse thoroughly with water. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Flammable - keep away from fire or flame. Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling, irritation or other symptoms occur, consult a physician immediately. Made in CHINA LOT/EXP: Insect Sting Relief Pad

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

1/pouch

GENUINE FIRST AID

Active Ingredients

Active Ingredient:Bacitracin Zinc 400 units Neomycin Sulfate 5mg (equivalent to 3.5 mg Neomycin base) Polymyxin B Sulfate 5000 units

Purpose

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

Warnings

For external use only.

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

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

Stop use and consult a doctor: if the condition persists or gets worse; a rash or other allergic reaction develops Keep out of reach of children. If ingested, contact a Poison Control Center right away.

Directions

Directions: clean affected area; apply small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily; may be covered with a sterile bandage

Other information:

Store at room temperature.

Inactive Ingredient

water Genuine Triple Antibiotic First Aid Ointment To Help Prevent Infection Each Gram Contains: Bacitracin Zinc 400 units Neomycin Sulfate 5 mg (equivalent to 3.5 mg Neomycin base) Polymyxin B Sulfate 5000 units Net Wt. 0.5g ; (1/64 oz) Manufactured in CHINA for GENUINE FIRST AID.

Triple Antibiotic Ointment 10pcs Net wt. 0.9g (1/32oz)

100 Triple Antibiotic Active Ingredient (in each tablet) Purpose Acetaminophen 325 mg Analgesic/antipyretic Uses temporary relief of minor aches and pains associated with:

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

reduction of fever

Warnings:

Liver warning: This product contains acetaminophen.

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

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

Ask a doctor before use if the user has liver disease

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

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

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

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt

medical attention is critical for adults as well as for children even if

you do not notice any signs or symptoms. Do not exceed recommended dosage

Directions

Adults and Children	Take 2 tablets every 4 to 6 hours as
12 years of age	needed. Do not take more than 12 tablets
or older	in 24 hours.
Children 6-11 years	Take 1 tablet every 4 to 6 hours as
of age	needed. Do not take more than 5
	tablets in 24 hours.
Children under 6	Do not use this regular strength product.
years of age	This will provide more than the
	recommended dose (overdose) and could
	cause serious health problems.
Store at 59-86 degree F (15-30 d	legree C)

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

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

GENUINE FIRST AID 2 Tablets

NON-ASPIRIN

Active ingredient (in each tablet)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.

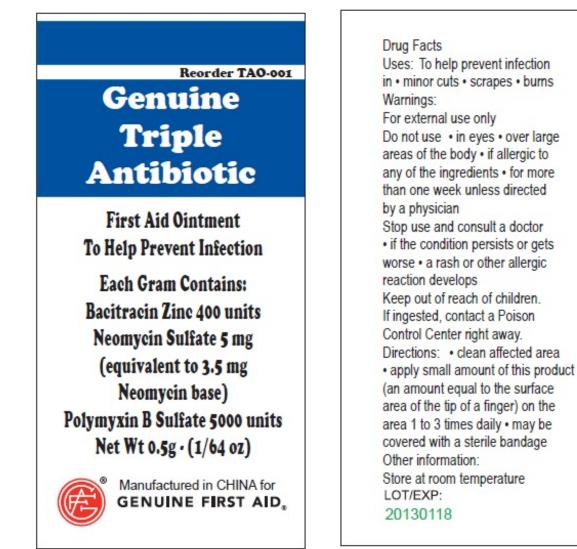
Other information: Store at controlled room temperature; avoid excessive heat 40 degree Celsius (104 degree Fahrenheit); tamper evident sealed packets; do not use any opened or torn packets

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



2 Tablets

toothache

arthritis

Warnings (continued)

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

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

Directions

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

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

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

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

Warnings

common cold

muscular aches backache

Uses

with

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

GENUINE FIRST AID.

Active ingredient (in each tablet) Purpose

Acetaminophen 325 mg Analgesic/antipyretic

temporary relief of minor aches and pains associated

menstrual cramps and reduction of fever

headache

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

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



DRUG FACTS - Antiseptic Towelette

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 LOT/EXP: 20130301
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DRUG FACTS - Insect Sting Relief Pad

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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 wou experience any of the following signs of stomach bleeding: I 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 redeness or swelling is present in the painful area any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

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

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

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

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

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

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



EASY CARE FIRST AID KIT - OUTDOOR AND TRAVEL

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

Prod	uct Informatio	n			
Produ	ct T yp e	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-0699	
Packa	iging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC	:44224-0699-5	1 in 1 KIT			
Quan	tity of Parts				
Part #	Pa	ackage Quantity	Total Product Quantity		
Part 1	10 PACKAGE		8 mL		
	4 PACKAGE		2.0 mL		
Part 2					

Part 42 PACKAGEPart 52 PACKET	4				
IIII ZIMGALI	4				
Part 1 of 5					
ANTISEPTIC TOWELI	TTF				
benzalkonium chloride swab					
Product Information					
Item Code (Source)	NDC:52124-0001				
Route of Administration	TOPICAL				
Active Ingredient/Active Mo	-		D	f Sturmently	Store with
Ingr BENZALKONIUM CHLORIDE (UNII:	<mark>edient Name</mark> F5UM2KM3W7) (BENZAI	KONIUM -	BENZALK	of Strength	Strength
UNII:7N6JUD5X6Y)			CHLORIDE		in 100 mL
Inactive Ingredients					
	ngredient Name			Stre	ngth
WATER (UNII: 059QF0KO0R)	ngreutenertune			5410	
Packaging					
# Item Code Pac	kage Description	Marketing S	Start Date	Marketir	ıg End Date
1 NDC:52124-0001-1 0.8 mL ir	1 PACKAGE				
Marketing Information		-	-	-	
	tion Number or Monog	_	Marketing Sta	rt Date Marl	keting End Date
OTC monograph not final part333E		0	14/13/2011		
Part 2 of 5					
INSECT STING RELIE	F PAD				
	F PAD				
INSECT STING RELIE	F PAD				
INSECT STING RELIE	F PAD				
INSECT STING RELIE benzocaine,alcohol swab Product Information					
INSECT STING RELIE	F PAD NDC:52124-0008 TOPICAL				

Active Ingredient/Active Mo	oiety						
Ing	redient Name		Basis o	f Strength	Strength		
BENZOCAINE (UNII: U3RS Y48 JW5) (BENZOCAINE - UNII: U3RS Y48 JW5) BENZOC					6 mL in 100 mL		
ALCOHOL (UNII: 3К9958 V90М) (AI	LCOHOL - UNII:3K9958V9	0 M)	ALCOHO	L	60 mL in 100 mL		
Inactive Ingredients							
Ingredient Name					Strength		
WATER (UNII: 059QF0KO0R)							
Packaging							
# Item Code Pa	ckage Description	Marketin	g Start Date	Mark	eting End Date		
1 NDC:52124-0008-1 0.5 mL	in 1 PACKAGE						
Marketing Information							
Marketing Category Applicat	tion Number or Monogra	aph Citation	Marketing St	art Date M	arketing End Date		
OTC monograph final part348			04/13/2011				
Part 3 of 5							
GENUINE TRIPLE AN	TIBIOTIC						
bacitracin zinc,neomycin sulfate,polymyxin b sulfate ointment							
Product Information							
Item Code (Source)	NDC:52124-0003						
Route of Administration	TOPICAL						
Active Ingredient/Active Mo	biety						
Ing	redient Name		Basi	s of Strengtl	n Strength		
BACITRACIN ZINC (UNII: 89 Y4M234	IES) (BACITRACIN - UNII:5	68 H6 RWO 52 I)	BACITR	RACIN ZINC 400 [iU] in 1			
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII: 116QD7X297) NEOMY				YCIN SULFATE 5 mg in 1 g			
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ07J96K) POLYMYXIN B SULFATE					TE 5000 [iU] in 1 g		
Inactive Ingredients							
	Ingredient Name			S	trength		
WATER (UNII: 059QF0KO0R)							

Packaging							
# Item Code						rketing E	and Date
1 NDC:52124-0003-1	0003-1 0.5 g in 1 TUBE						
Marketing Inform	nation						
Marketing Category		n Number or Monogra	ph Citation	Marketing	Start Date	Marketin	g End Date
OTC monograph final p	art333B			04/13/2011			
Part 4 of 5							
NON-ASPIRIN							
acetaminophen tablet							
Product Information	1						
Item Code (Source)		NDC:52124-0010					
Route of Administration	l	ORAL					
Active Ingredient/Active	ctive Moie	ety					
		gredient Name			Basis of S	trength	Strength
ACETAMINOPHEN (UNII:	362O9ITL9E) (ACETAMINOPHEN - U	NII:362O9ITL9	D)	ACETAMINO	PHEN	325 mg
T							
Inactive Ingredients		T.,				6.6	
STARCH, CORN (UNII: 08	222MV2S I)	Ingredient Name				50	rength
POLYETHYLENE GLYCO		IO0SDW1A)					
STEARIC ACID (UNII: 4EL)							
POVIDONE (UNII: FZ989G	H94E)						
	.•						
Product Characteris			0				
Color	white (WHI ROUND	IE)	Score Size			no score	
Shape Flavor	ROUND		Imprint Co	da		11mm AZ;234	
Contains			Imprint Co	ue		AL,254	
Contains							
Packaging							
# Item Code	Pack	age Description	Marketin	g Start Date	Ma	rketing E	and Date
1 NDC:52124-0010-1	2 in 1 PAC	KAGE					

Marketing Inform	ation						
Marketing Category	Application Number or Monog	raph Citation Mar	keting Start Date	Marketing End Date			
	art343	-	3/2011				
Part 5 of 5							
IBUPROFEN							
ibuprofen tablet							
Product Information							
Item Code (Source)	NDC:52124-0009						
Route of Administration	ORAL						
Active Ingredient/Act	•						
	Ingredient Name		Basis of Str				
IBUPROFEN (UNII: WK2XYI1	l0QM) (IBUPROFEN - UNII:WK2XY	I10 Q M)	IBUPRO FEN	200 mg			
Inactive Ingredients							
	Ingredient Name	2		Strength			
POWDERED CELLULOSE (UNII: SMD1X3XO9M)						
STARCH, CORN (UNII: 08232NY3SJ)							
HYPROMELLOSES (UNII: 3)	HYPROMELLOSES (UNII: 3NXW29V3WO)						
LACTOSE (UNII: J2B2A4N98	3G)						
MAGNESIUM STEARATE (U	JNII: 70097M6I30)						
POLYDEXTROSE (UNII: VH	2XOU12IE)						
POLYETHYLENE GLYCOL	(UNII: 3WJQ0SDW1A)						
POVIDONE (UNII: FZ989GH	94E)						
SILICON DIO XIDE (UNII: ET	TJ7Z6XBU4)						
STEARIC ACID (UNII: 4ELV7	7Z65AP)						
TITANIUM DIO XIDE (UNII: 1							
TRIACETIN (UNII: XHX3C3X	(673)						
Product Characteristi	cs						
Color	white (White)	Score		no score			
Shape	ROUND	Size		10 mm			
Flavor		Imprint Code		44;352			
Contains							
Packaging							
# Item Code	Package Description	Marketing Sta	rt Date Ma	rketing End Date			
1 NDC:52124-0009-1	2 in 1 PACKET	Marine ting Sta		The ting Life Dute			
1120.52124-0005-1							

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
ANDA	ANDA075010	02/17/2010				
Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part348	04/13/2011				

Labeler - Tender Corp dba Adventure Medical Kits (064437304)

Revised: 4/2011

Tender Corp dba Adventure Medical Kits