MOUNTAIN SERIES WEEKENDER MEDICAL - benzalkonium chloride, povidone-iodine, acetaminophen, aspirin, diphenhydramine hydrochloride, ibuprofen, bacitracin zinc, neomycin sulfate, polymyxin b sulfate Tender Corporation 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.

Mountain Series Weekender Medical Kit

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 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)

Aspirin (NSAID*) 325 mg *nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

Temporarily relieves minor aches and pains associated with

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

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.

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 thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcohol 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 prescription drugs 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 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 take more than directed
- the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)
- drink a full glass of water with each dose

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

- read all product information before using
- store at room temperature 59°-86°F (15°-30°C)
- avoid excessive heat and humidity
- tamper evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

croscarmellose sodium*, hypromellose, microcrystalline cellulose*, mineral oil*, polyethylene glycol*, propylene glycol*, starch, titanium dioxide*

*may contain

Questions or comments?

1-800-634-7680

116R Medique APAP 325 mg Label

Medique® Aspirin 5 Grain (325 mg) Pain Reliever/Fever Reducer (NSAID) See new warnings information East To Swallow Film Coated Tablets 24 Tablets (12 x 2) Compare active ingredients to Bayer® Registered Trademark Bayer Consumer

Active ingredients

(in each capsule) Diphenhydramine Hydrochloride 25 mg

Purpose

Antihistamine

Uses

Temporarily relieves runny nose and decreases sneezing, itching of the nose and throat, itchy-watery eyes due to hay fever or other respiratory allergies.

Warnings

Do not use

• with any other product containing diphenhydramine, even one that is used on skin.

Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- glaucoma

Ask a doctor or pharmacist before use if you are

- taking any drugs for asthma
- taking sedatives or tranquilizers

When Using This Product

- marked drowsiness may occur
- avoid alcohol beverages
- 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

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

Keep out of the reach of children. In case of overdose, contact a physician or poison control center immediately.

Directions

• do not use more than directed

Adults and children: (12 years and older): Take 1 to 2 capsules every 4 to 6 hours as needed. Do not take more than 12 capsules in 24 hours.

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

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

D&C Red #28, FD&C Blue #1, FD&C Red #40, gelatin, starch

Questions or comments? 1-800-634-7680

182R Medique Diphen Label

Collect Medi-Bucks

See inside flap for more details

Medique®

Diphen

Diphenhydramine HCl 25 mg

- Hay Fever/Allergies
- Fiebre del Heno/Alergias

Pull to Open

TiraParaAbrir

Easy To Swallow Capsules

Capsulas Faciles de Tragar

200 Capsules

(200 x 1)

Tamper Evident Unit Dose Packets

Empaquetado con Sellado Evidente en Dosis Untarias

Active ingredient (in each tablet)

Ibuprofen (NSAID^{*}) 200 mg

*nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

Temporarily relieves minor aches and pains associated with

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

Temporarily reduces fever.

Warnings

Allergy alert: Ibuprofen may cause a severe allergic reaction which may include:

- hives
- asthma (wheezing)
- rash
- blisters
- skin reddening

- facial swelling
- 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 thinning (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

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 aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- taking any other drug

When using this product

• take with food or milk if stomach upset occurs

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
 - feel faint
 - vomit blood
 - have bloody or black stools
 - have stomach pain that does not get better
 - pain gets worse or lasts for more than 10 days
 - fever gets worse or lasts for more than 3 days
 - redness or swelling is present in the painful area
- any new or unexpected symptoms occur

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 specifically directed to 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 (see Warnings)

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

- read all product information before using
- store at 68-77°F (20-25°C)
- avoid excessive heat 40°C (above 104°F)
- tamper evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

carnauba wax^{*}, cellulose^{*}, colloidal silicon dioxide, corn starch^{*}, hypromellose, iron oxide red^{*}, lactose, magnesium stearate, microcrystalline cellulose^{*}, polydextrose, polyethylene glycol, povidone, silica^{*}, sodium lauryl sulfate^{*}, sodium starch glycolate, stearic acid^{*}, titanium dioxide, triacetin^{*}

*may contain

Questions or comments? 1-800-634-7680

Principal Display Panel

Medi-First® Ibuprofen 200 mg Pain Reliever/Fever Reducer (NSAID) Easy To Swallow Film Coated Tablets See new warnings information 100 tablets (50 x 2) Compare active ingredient to: Genuine Advil® Registered trademark Wyeth Consumer Tamper Evident Unit Dose Packets

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/fever reducer

Uses

For the temporary relief of minor aches and pains associated with

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

For the reduction of fever.

Warnings

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

- more than 12 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 nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if

• you are taking the blood thinning drug warfarin

Stop using and ask a doctor if

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

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

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

Directions

• do not use more than directed

Adults and children: (12 years and older)

Take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours.

Children under 12 years:

Do not give to children under 12 years of age.

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

Inactive ingredients

corn starch, hypromellose, maltodextrin*, microcrystalline cellulose*, polyethylene glycol, povidone, pregelatinized starch*, sodium starch glycolate, stearic acid

* may contain

Questions or comments? 1-800-634-7680

Principal Display Panel

145R Medique APAP 325 mg Label

Collect MediBucks See inside flap for more details Medique[®] APAP Acetaminophen 325 mg Pain Reliever/Fever Reducer Alivia el Dolor/Reduce la Fiebre Easy To Swallow Film Coated Tablets Facil de Tragar Tabletas con Cubierta Pelicular Pull to Open Tire Para Abrir See new warnings information 500 Tablets (250 x 2) Tamper Evident Unit Dose Packets Empaquetado con Sellado Evidente en Dosis Unitarias

Weekender Kit Label

Mountain Series Medical Kit Weekender Adventure Medical Kits BE SAFE 1-6 People up to 7 days Povidone-iodine 10% Antiseptic Warnings Do not use • if allergic to iodine

• in the eyes

For external use only

Ask a doctor before use if injuries are

- deep or puncture wounds
- serious burns

Stop use and ask a doctor if

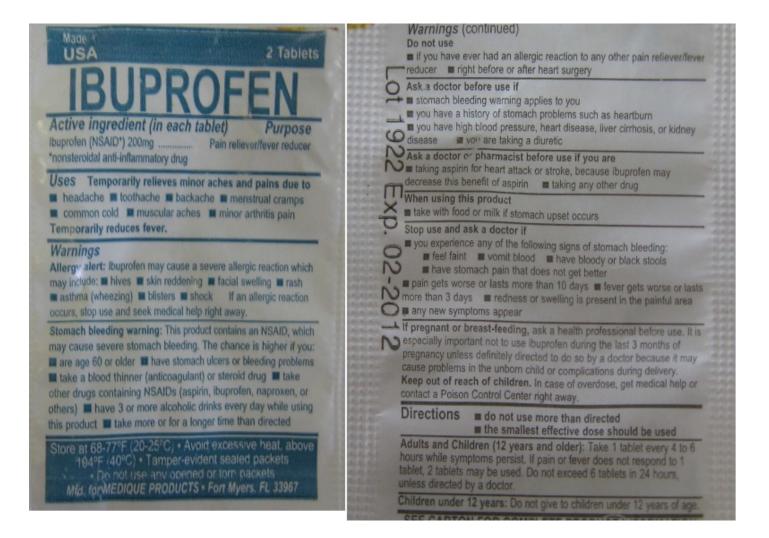
- redness, irritation, swelling or pain persists or increases
- infection occurs

Avoid pooling beneath patient

Keep out of reach of children. In case of accidental ingestion, seek professionalassistance or consult a poison control center immediately.







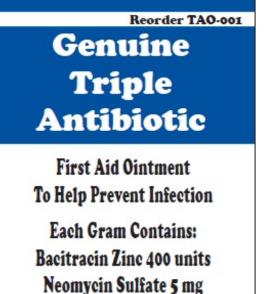


Medique NDC# 47682-182-46 Made in USA Diphen 1 Capsule	Warnings Ask a Doctor Before Use If you have: • a breathing problem such as emphysema or chronic bronchitis • glaucoma • difficulty in urination due to enlargement of the prostate gland If you are: • taking any drugs for asthma • taking sedatives or tranquilizers When Using this product: • drowsiness may occur • avoid alcohol beverages
Active Ingredient Purpose (In each capsule) Diphenhydramine Hydrochloride 25mgAntihistamine	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. o not exceed recommended dosage. Keep this and all drugs
Uses: Temporarily relieves the following symptoms associated with hay fever or other upper respiratory allergies:	It of the reach of children. In case of accidental overdose, ntact a physician or poison control center immediately. As with y drug, if you are pregnant or nursing a baby, seek the advice a health professional before using this product.
 runny nose itching of the nose or throat itchy, watery eyes 	rections: Do not use more than directed ults and children: (12 years and older) Take 1 capsule ry 4 to 5 hours as needed. Do not take more than 12 capsules
Store at room temperature • Tamper-Evident Sealed Packets • Do Not Use any Opened or Torn Packets Mfd. for MEDIQUE PRODUCTS Fort Myers, FL 33967	



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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(equivalent to 3.5 mg Neomycin base) Polymyxin B Sulfate 5000 units Net Wt 0.5g · (1/64 oz)



Manufactured in CHINA for GENUINE FIRST AID_® Drug Facts

Uses: To help prevent infection in • minor cuts • scrapes • burns Warnings:

For external use only Do not use • in eyes • over large areas of the body • if allergic to any of the ingredients • for more than one week unless directed by a physician Stop use and consult a doctor

 if the condition persists or gets worse · a rash or other allergic reaction develops Keep out of reach of children. If ingested, contact a Poison Control Center right away. Directions: • clean affected area apply small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily • may be covered with a sterile bandage Other information: Store at room temperature LOT/EXP: 20130118



SEVERE BLEEDING Instructions, Easy Care**, Bleeding Trauma Pad, 5* x 9* Gloves, Nitrile (Pair), One Hand Wipe SPRAIN / STRAIN Instructions, Easy Care**, Fractures/Sprains Ibuprefen (200 mg), Pkg./2	WOUND CARE / BURN / BLISTE Instructions. Easy Care", Wound Moleskin, Pre-cut and Shaped Dressing, Glacier Gel, Rectangle, 2.5" Tincture of Benzoin Topical Adhesive, 3 Syringe, Irrigation, 20 cc, 18 Gauge Tr After Cuts & Scrapes Antiseptic Wipe Triple Antibiotic Ointment, single use Bandage, Adhesive, Fabric, Knuckle Bandage, Adhesive, Fabric, 1 * x 3" Wound Closure Strips, 1/4 * x 4", Pkg Dressing, Gauze, Sterile, 4* x 4", Pkg Dressing, Gauze, Sterile, 4* x 4", Pkg Bandage, Conforming Gauze, Non-S Dressing, Non-Adherent, Sterile, 3* x	1 Aspirin (325 mg), Pkg./2 2 Antihistamine (Diphenhydramine 25 mg) 1 Ibuprofen (200 mg), Pkg./2 2 Acetaminophen (500 mg), Pkg./2 1 INSTRUMENTS / INSTRUCTION 1 Comp. Guide to Wilderness & Travel Medicine 1 Spinter Picker/Tick Remover Forceps 1 EMT Shears. 4* 1 Outr Tape, 2* x 5 Yards 0./2 3 0./2 3 0./2 3 1 Cotton Tip Applicator. Pkg./2 1 Cotton Tip Applicator. Pkg./2
	Instructions, Easy Care ^w , Bleeding Trauma Pad, 5 ^e x 9 ^e Gloves, Nitrile (Pair), One Hand Wip SPRAIN / STRAIN Instructions, Easy Care ^w , Fractures	

MOUNTAIN SERIES WEEKENDER MEDICAL

benzalkonium chloride, povidone-iodine, acetaminophen, aspirin, diphenhydramine hydrochloride, ibuprofen, bacitracin zinc, neomycin sulfate, polymyxin b sulfate kit

Prod	uct Information	l						
Produ	ict T yp e	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-0118				
Packa	aging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC	:44224-0118-1	1 in 1 KIT						
Quan	tity of Parts							
Part # Package Quantity Total Product Quantity								
	6 PACKAGE		4.8 mL					
Part 1	6 PACKAGE							
Part 1 Part 2	3 TUBE		1.5 g					

Part 4	2 PACKET		2	-			
Part 5	1 PACKET		2	2			
Part 6	2 PACKET		2	ļ.			
Part 7	1 PACKET		2	22 g			
Part	: 1 of 7						
	ISEPTIC T		TTE				
benza	lkonium chloride	swab					
Prod	uct Information	1					
Ite m C	Code (Source)		NDC:52124-0001				
Route	of Administration	1	TOPICAL				
Activ	e Ingredient/A	ctive Moi	etv				
i icuiv			edient Name		Basis of	Strength	Strength
	ALKONIUM CHLOI NG JUD5XG Y)	-	5UM2KM3W7) (BENZAI	LKONIUM -	BENZALKO CHLORIDE	-	0.40 mL in 100 mL
Inacti	ive Ingredients						
		I	ngredient Name			Stı	rength
	ive Ingredients R (UNII: 059QF0KO	I	ngredient Name			Str	rength
		I	ngredient Name			Sti	rength
WATE	R (UNII: 059QF0KO	I	ngredient Name			Stı	rength
	R (UNII: 059QF0KO	In OR)	ngredient Name sage Description	Marketing	Start Date		rength ing End Date
WATE Packa #	R (UNII: 059QF0KO aging	In OR) Pack		Marketing	Start Date		
WATE Packa #	R (UNII: 059QF0KO aging Item Code	In OR) Pack	cage Description	Marketing	Start Date		
WATEI Packa # 1 NDC	R (UNII: 059QF0KO aging Item Code :52124-0001-1	In OR) Pack 0.8 mL in	cage Description	Marketing	Start Date		
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Route of Administration	1	TOPICAL				
Active Ingredient/A	ctive Moi	ety				
	Ingro	edient Name		Basis o	f Strength	Strength
BACITRACIN ZINC (UNII:	89Y4M234E	S) (BACITRACIN - UNII:5	58 H6 RWO 52 I)	BACITRAC	IN ZINC	400 [iU] in 1 g
NEO MYCIN SULFATE (UI	NII: 057Y626	693) (NEOMYCIN - UNII:	:I16QD7X297)	NEOMYCIN	I SULFATE	5 mg in 1 g
POLYMYXIN B SULFATE	E (UNII: 1937)	1312D4) (POLYMYXIN B -	- UNII:J2VZ07J96K)	POLYMYXI	N B SULFATE	5000 [iU] in 1 §
Inactive Ingredients						
	I	ngredient Name			Stre	ngth
WATER (UNII: 059QF0KO	0 R)					
Packaging						
# Item Code	Pack	kage Description	Marketing Sta	rt Date	Marketir	ıg End Date
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Marketing Category		on Number or Monogra	_	-	t Date Mark	eting End Date
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Marketing Category OTC monograph final p Part 3 of 7 MEDIQUE ASPI aspirin tablet, film coate Product Information Item Code (Source) Route of Administration Active Ingredient/Action	Application part333B RIN ed 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	NDC:47682-116 ORAL ety dient Name	02/16	/20 10 Basis 0		Strength
Marketing Category OTC monograph final p Part 3 of 7 MEDIQUE ASPI aspirin tablet, film coate Product Information Item Code (Source) Route of Administration Active Ingredient/Action	Application part333B RIN ed 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	NDC:47682-116 ORAL ety dient Name	02/16	/20 10 Basis 0		Strength
Marketing Category DTC monograph final p Part 3 of 7 MEDIQUE ASPI aspirin tablet, film coate Product Information Item Code (Source) Route of Administration Active Ingredient/Ad ASPIRIN (UNII: R16C05Y7 Inactive Ingredients	Application part333B RIN ed 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	NDC:47682-116 ORAL ety dient Name N - UNII:R16CO5Y76E) Ingredient Name	02/16	/20 10 Basis 0		Strength 325 mg
OTC monograph final p Part 3 of 7 MEDIQUE ASPI	Application art333B RIN ed 1 1 1 1 1 1 1 1 1 1 1 1 1	NDC:47682-116 ORAL ety dient Name N - UNII:R16C05Y76E) Ingredient Name WO) JQ0SDW1A)	02/16	/20 10 Basis 0		Strength 325 mg

STARCH, CORN (UNII:	08232NY3SJ)						
, ,	,						
Product Characte	ristics						
Color	white (white)		Score		no sc	ore	
Shape	ROUND (ROUI	ND)	Size		10 mm	1	
Flavor			Imprint Cod	le	44;15	7;aspirin	
Contains			•			-	
Packaging							
# Item Code	Pac	kage Description	Marketin	ng Start Date	Ma	arketing En	d Date
1 NDC:47682-116-99	2 in 1 PA(ig Start Date			u Dutt
Marketing Info	ormation						
Marketing Category	Applicatio	on Number or Monogra	ph Citation	Marketing Start	Date	Marketing	End Date
OTC monograph final	part343			12/30/2008			
Part 4 of 7							
MEDIQUE DI	PHEN						
diphenhydramine hydra		ماريمه					
		apsuie					
Product Informat	ion						
Item Code (Source)		NDC:47682-182					
Route of Administrat	tion	ORAL					
Active Ingredient							
	•	edient Name				rength	Strength
DIPHENHYDRAMINE H UNII:8GTS82S83M)	IYDRO CHLOR	IDE (UNII: TC2D6JAD40)	(DIPHENHYDRA	AMINE - DIPHENHYD HYDROCHL		E	25 mg
					onabl		
Inactive Ingredie	nts						
		Ingredient Name				Streng	gth
D&C RED NO. 28 (UNI	I: 767IP0Y5NH)	_					
FD&C BLUE NO. 1 (UN	III: H3R47K3TBI	D)					
FD&C RED NO. 40 (UN	NII: WZB9127XC	DA)					
GELATIN (UNII: 2G860	QN327L)						
STARCH, CORN (UNII:	08232NY3SJ)						
Product Characte	ristics						

Color	pink (pink) , w	nite (white)	9	Score	no s	core
	CAPSULE (CA			Size	110 s	
Shape Flavor	GAISULE (CF					;835
			I	mprint Code	CPC	.,035
Contains						
Packaging						
# Item Code	Pac	kage Description	Marketin	ng Start Date	Marketing	End Date
1 NDC:47682-182-46	1 in 1 PA	CKET				
Marketing Inf						
Marketing Category	y Applicat	ion Number or Monogra	aph Citation	Marketing Start	Date Market	ing End Date
OTC monograph final	part341			12/30/2008		
Part 5 of 7						
MEDI-FIRST	IBUPRO	FEN				
ibuprofen tablet, filr	n coated					
r r r r r r r r r r r r r r r r r r r						
Product Informat	tion					
Item Code (Source)		NDC:47682-808				
Route of Administra	tion	ORAL				
Route of Administra	uon	OIML				
Active Ingredient	t/Active Mo	ietv				
		gredient Name		Basis	of Strength	Strength
IBUPROFEN (UNII: Wł		BUPROFEN - UNII:WK2XYI	10 O M)	IBUPROFE		200 mg
	() (
Inactive Ingredie	nts					
Inactive Ingredie	nts	Ingredient Nam	le			Strength
5		0	10			Strength
CELLULOSE, MICRO	CRYSTALLIN	E (UNII: OP1R32D61U)	le			Strength
CELLULOSE, MICRO HYPROMELLOSES (U	CRYSTALLIN JNII: 3NXW29 V	E (UNII: OP1R32D61U)	10			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2	CRYSTALLIN JNII: 3NXW29 V A4N98G)	E (UNII: OP1R32D61U) 3WO)	le			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN	CRYSTALLIN JNII: 3NXW29V A4N98G) XTE (UNII: 7009 MII: VH2XOU12I	E (UNII: OP1R32D61U) 3WO) 97M6I30) E)	10			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN	CRYSTALLIN JNII: 3NXW29V A4N98G) XTE (UNII: 7009 MII: VH2XOU12I	E (UNII: OP1R32D61U) 3WO) 97M6I30) E)	le			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN POLYETHYLENE GLY POVIDONE (UNII: FZ9	CRYSTALLIN JNII: 3NXW29 V A4N98G) ATE (UNII: 7009 NII: VH2XOU12I YCOL (UNII: 3N 89GH94E)	E (UNII: OP1R32D61U) 3WO) 97M6130) E) VJQ0SDW1A)	1e			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN POLYETHYLENE GLY POVIDONE (UNII: FZ9 SILICON DIOXIDE (U	CRYSTALLIN JNII: 3NXW29 V A4N98G) MTE (UNII: 7009 MI: VH2XOU12I YCOL (UNII: 3 89GH94E) NII: ETJ7Z6XB	E (UNII: OP1R32D61U) 3WO) 97M6I30) E) VJQ0SDW1A) U4)	1e			Strength
Inactive Ingredie CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN POLYETHYLENE GL POVIDONE (UNII: FZ9 SILICON DIOXIDE (U STARCH, CORN (UNII	CRYSTALLIN JNII: 3NXW29V A4N98G) ATE (UNII: 7009 III: VH2XOU121 YCOL (UNII: 3V 89GH94E) NII: ETJ7Z6XB : O8232NY3SJ)	E (UNII: OP1R32D61U) 3WO) 97M6I30) E) VJQ0SDW1A) U4)	1e			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN POLYETHYLENE GLY POVIDONE (UNII: FZ9 SILICON DIOXIDE (U STARCH, CORN (UNII STEARIC ACID (UNII:	CRYSTALLIN JNII: 3NXW29 V A4N98G) ME (UNII: 7009 MI: VH2XOU12I YCOL (UNII: 3V 89GH94E) NII: ETJ7Z6XB : O8232NY3SJ) 4ELV7Z65AP)	E (UNII: OP1R32D61U) 3WO) 97M6130) E) WJQ0SDW1A) U4)	1e			Strength
CELLULOSE, MICRO HYPROMELLOSES (U LACTOSE (UNII: J2B2 MAGNESIUM STEARA POLYDEXTROSE (UN POLYETHYLENE GLY POVIDONE (UNII: FZ9 SILICON DIOXIDE (U STARCH, CORN (UNII	CRYSTALLIN JNII: 3NXW29 V A4N98G) ME (UNII: 7009 MI: VH2XOU12I YCOL (UNII: 3V 89GH94E) NII: ETJ7Z6XB : O8232NY3SJ) 4ELV7Z65AP)	E (UNII: OP1R32D61U) 3WO) 97M6130) E) WJQ0SDW1A) U4)				Strength

Product Character	istics						
Color	white (white)		Score			no score	2
Shape	ROUND (RO	UND)	Size			10 mm	
Flavor			Imprin	t Code		44;352	
Contains							
Packaging							
# Item Code	Pac	kage Description	Marketii	ng Start Date	e Ma	rketing H	End Date
1 NDC:47682-808-99	2 in 1 PA			0		U	
Marketing Info	mation						
Marketing Category	Applicatio	on Number or Monogra	ph Citation	Marketing	Start Date	Marketin	ng End Date
ANDA	ANDA075139			12/30/2008			
Part 6 of 7							
MEDIQUE APA	P						
acetaminophen tablet,	, film coated	l					
Product Information	n						
Item Code (Source)		NDC:47682-145					
Route of Administration	n	ORAL					
Active Ingredient/A	Active Moi	etv					
		gredient Name			Basis of S	trength	Strength
ACETAMINOPHEN (UNI) (ACETAMINOPHEN - UI	NII:36209ITL9	D)	ACETAMINO	-	325 mg
Inactive Ingredient	ts						
		Ingredient Nan	ne				Strength
HYPROMELLOSES (UN							
POLYETHYLENE GLYC		JQ0SDW1A)					
POVIDONE (UNII: FZ989			126242)				
STARCH, CORN (UNII: C		E A POTATO (UNII: 5856	J3G2A2)				
STEARIC ACID (UNII: 4E							
Product Character	istics						
Color	white (white)		Score			no score	2
~~~~~	(		Score				

Shape	ROUND	(ROUND)	Size			mm
Flavor			Imprint	Code	AZ	;234
Contains						
Packaging						
# Item Code		Package Description	Marketing	g Start Date	Marketi	ing End Date
1 NDC:47682-145-99	2 in 1	PACKET				
Marketing Inf	ormatio	n				
Marketing Catego	ry Appli	ication Number or Monog	raph Citation	Marketing Star	rt Date Mar	rketing End Date
OTC monograph not fin	nal part343			12/30/2008		
Part 7 of 7						
ADI ICADE D		NE-IODINE SOLU	TION			
			IION			
povidone-iodine so	lution soluti	ion				
Product Informa	tion					
Item Code (Source)		NDC:52380-0001				
Route of Administra	tion	TOPICAL				
Active Ingredien	t/Active M	Ioiety				
	Iı	ngredient Name		Basis o	f Strength	Strength
PO VIDO NE-IO DINE (	UNII: 85H0HZ	- ZU99M) (IODINE - UNII:9679	TC07X4)	POVIDONI	E-IODINE	9.8 g in 100 g
Inactive Ingredie	nts					
		Ingredient Name	e			Strength
CITRIC ACID MONOI						
SO DIUM PHO SPHAT		,				
SO DIUM HYDRO XIDI						
NONOXYNOL-9 (UNI		9T)				
WATER (UNII: 059QF0	KO0R)					
Packaging						
# Item Code	I	Package Description	Marketin	g Start Date	Market	ing End Date
1 NDC:52380-0001-3	22 g i	n 1 PACKET				
Marketing Inf	ormatio	n				
mar Keung In		11				

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	0 3/0 1/19 8 4					
Marketing Information							
maine mon							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
U		Marketing Start Date	Marketing End Date				

# Labeler - Tender Corporation dba Adventure Medical Kits (064437304)

Revised: 8/2011

Tender Corporation dba Adventure Medical Kits