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

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

Genuine First Aid Auto First Aid Kit

Active Ingredient:

Purpose

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

Use: For preparation of skin before injection.

Warnings: For external use only.

Flammable - keep away from fire or flame

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

Do not use: with electrocautery, in the eyes.

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

Keep out of reach of children. If swallowed,

get medical help or contact a Poison Control

Center right away.

Wipe Injection site vigorously and discard.

Inactive Ingredient: Purified water.

LOT/EXP: Made in CHINA

20140301

Alcohol Cleansing Pad Genuine First Aid LLC, Clearwater FL 33755

www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

Active Ingredient:Bacitracin Zinc 400 units

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

Polymyxin B Sulfate 5000 units

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

Warnings:

For external use only.

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

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

Stop use and consult a doctor:

if the condition persists or gets worse; a rash or other allergic reaction develops

Keep out of reach of children.

If ingested, contact a Poison

Control Center right away.

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

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

Active ingredient (in each tablet)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

Warnings

Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include: shock, facial swelling, asthma (wheezing) rash, skin reddening, blisters, hives If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause severe stomach

bleeding. The chance is higher if you: are age 60 or older, have had stomach ulcers or bleeding problems, take a blood thinner (anticoagulant) or steroid drug, take other drugs containing NSAIDs (aspirin, ibuprofen, naproxen, or others), have 3 or more alcoholic drinks every day while using this product, take more or for a longer time than directed

Do not use if you have ever had an allergic reaction to any other pain reliever/fever reducer, right before or after heart surgery.

Ask a doctor before use if stomach bleeding warning applies to you; you have a history of stomach problems such as heartburn; you have a high blood pressure, heart disease, liver cirrhosis, or kidney disease; you are taking a diuretic

Ask a doctor before use if you are:

taking any other drug containing NSAID (prescription or nonprescription); taking aspirin for heart attack or stroke, because Ibuprofen may decrease this benefit of aspirin; taking any other drug

When using this product: take with food or milk if stomach upset occurs

Stop use and ask a doctor If:

you experience any of the following signs of stomach bleeding; feel faint; vomit blood; have bloody or black stools; have stomach pain that does get better; pain gets worse or lasts more than 10 days; fever gets worse or lasts more than 3 days; redness or swelling is present in the painful area; any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions:

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

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

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

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

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

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

IBUPROFEN 2 Tablets

IBUPROFEN 2 Tablets

Active Ingredient (in each tablet) Purpose

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

Purpose: Pain reliever, fever reducer

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

Warnings

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

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

Ask a doctor before use if you have liver disease

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

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

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

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

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

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

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

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

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

Active Ingredient: Purpose

Benzalkonium Chloride 0.40%...... First Aid Antiseptic

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

Warning: For external use only.

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

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

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

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

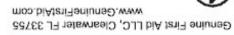
Inactive Ingredient: Purified water

LOT/EXP: Made in CHINA 20130301 Antiseptic Towelette Genuine First Aid LLC, Clearwater FL 33755 www.GenuineFirstAid.com 1/pouch GENUINE FIRST AID



GENUINE FIRST AID.

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REORDER AST-001

DRUG FACTS - Antiseptic Towelette

Active Ingredient: Purpose	E 1
Benzalkonium Chloride 0.40% First Aid Antiseptie	· .
Use: For Professional and Hospital use. Helps	
prevent infection. Antiseptic cleansing of face,	
hands and body without soap and water.	1
Warnings: For external use only. Keep out of	ш
reach of children. If swallowed, get medical help	RE
or contact a Poison Control Center right away.	ш
If unusual redness, swelling or other symptoms	T
occur, consult a physician immediately.	
Do not use: In the eyes, or over large areas of th	TEAR
body.	5
Directions: Tear open packet, unfold towelette	끈
and use to cleanse desired skin area. Discard	· ·
towelette appropriately after single use.	<u>'</u>
Inactive ingredient: Purified water.	
Made in CHINA	ı

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

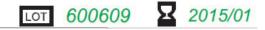
• apply small amount of this product (an amount equal to the surface

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



DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient:	Purpose:
Isopropyl Alcohol, 70% v/v	. Antiseptic
Use: For preparation of the sk	in before injection.
Warnings: For external use	only. I
Flammable - keep away from	n fire or flame.
Do not use: with electrocaute	ery, in the eyes 🔟
Stop use if irritation and redn	ess develop.
If condition persists for more t	han 72 hours, III
consult your doctor.	Т
Keep out of reach of childre	n. If swallowed, Poison Control e vigorously and
get medical help or contact a	Poison Control
Center right away.	1
Directions: Wipe injection sit	e vigorously and
discard.	
Other information: Store at r	
15°-30° C (59°-86° F)	
Inactive ingredient: Purified	water.
LOT/EXP:	Made in CHINA

20140301

Warnings (continued) Do not use
 with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure wheth a drug contains acetaminophen, ank a doctor or pharmaci for more than 10 days for pain unless directed by a doct for more than 3 days for fever unless directed by a doct
Ask a doctor before use if you have in 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 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
Adults and children: (12 years and older) Take 2 lablets every 4 to 6 hours as needed. Do not take more than 8 lablets in 24 hours.
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





GENUINE FIRST AID - AUTO FIRST AID

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

Product T Packagin		HUMAN OTC DRUG	Item Code (Source)	NDC:52124-1000
Packagir	20			
Packagin	ng			
	iig			
# Ite	em Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:5212	24-1000-1	1 in 1 KIT		
Quantity Part #		ckage Quantity	Total Product	Quantity
		ckage Quantity	Total Product	Quantity
Part 1 1 TU			0.5 g	
Part 2 1 PA	ACKET		2	
Part 3 1 PA	ACKET		2	
	ACKAGE		1 C I	
Part 4 2 PA	ACKAGE		1.6 mL	

GENUINE TRIPLE				
bacitracin zinc, neomycin si	ulfate,polymyxin b sulfate oir	ntment		
Product Information				
Item Code (Source)	NDC:52124-0003			
Route of Administration	TOPICAL			
Active Ingredient/Activ	e Moiety			
	Ingredient Name		Basis of Streng	th Strength
BACITRACIN ZINC (UNII: 89 Y	4M234ES) (BACITRACIN - UNII:5	8H6RWO52I)	BACITRACIN ZINC	400 [iU] in 1 g
NEO MYCIN SULFATE (UNII: 0	57Y626693) (NEOMYCIN - UNII:	I16QD7X297)	NEOMYCIN SULFAT	E 5 mg in 1 g
POLYMYXIN B SULFATE (UN	NII: 19371312D4) (POLYMYXIN B -	UNII:J2VZ07J96K)	POLYMYXIN B SULF	ATE 5000 [iU] in 1 g
Inactive Ingredients				
	Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)				
Packaging				
# Item Code	Package Description	Marketing Star	t Date Mar	keting End Date
1 NDC:52124-0003-1 .5	g in 1 TUBE			
Marketing Informa	tion			
	plication Number or Monogra	ph Citation Mark	eting Start Date	Marketing End Date
OTC monograph final part33		0 1/11/2	-	-
Part 2 of 5				
IBUPROFEN				
ibuprofen tablet				
Product Information				
Item Code (Source)	NDC:52124-0009			
Route of Administration	ORAL			
Active Ingredient/Activ	e Moiety			
	Ingredient Name		Basis of Stre	ngth Strength
	<u> </u>			0

IBUPROFEN

POWDERED CELLUIOSE (UNE: SMD1/33X09M) STARCI, CORN (UME: 00222Y35) HYPROMEL.0085 (UNE: SWX29/34W0) LACTOSE (UNE: 228244986) MGENSEUM STEARATE (UME: 70097M610) LACTOSE (UNE: 728244986) MGENSEUM STEARATE (UME: 70097M610) POVIDOXE (UME: 728244986) POVIDOXE (UME: 7282497M610) POVIDOXE (UME: 51926901841) SILICON DIOXIDE (UME: 51926901842) SILICON DIOXIDE (UME: 51926901844) SILIC	PowDERED CELLULOSE (UNIE SMDIX3XO9M) STARCI, CORN (UNIE 09232NYSS) STARCI, CORN (UNIE 09232NYSS) LACTOSE (UNIE 1282A4N98G) AGORSE (UNIE 1282A4N98G) AGORSE (UNIE 12802A4N98G) STARCI (UNIE 12982A4N98G) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1007A62D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A667D) SOLUT	Inactive Ingredien	Ingredient Name	ρ		Strength
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HYPRO MELLOSES (UNIE 3NXW29 V3WO) Image: Status Statu	INTROMELLOSES (UNIE 3NXW29 V3WO) Image: Stream (Stream					
LACTO SE (UNIE J2B2A4N98G) MGGNESUUM STEARATE (UNIE 7097/MEISO) POLYETPKYLENE GLYCOL (UNIE 3W/Q0 SDW1A) POVTOONE (UNIE ETJ726 SDW) SULCON DIO XODE (UNIE ETJ726 SDW) STEARIC ACID (UNIE ETJ726 SDW) STEARIC ACID (UNIE ETJ726 SDW) STRACE TIN (UNIE ETJ726 SDW) TTANUM DIO XIDE (UNIE ETJ726 SDW) TAUEST Part 3 of 5 MECI47682-175 TEJ CATEGRA STRENGTH acetaminophen tablet, film coated NDC:47682-175 TEJ CATEGRA STRENGTH TEM CODE (SOURCE) NDC:47682-175	LACTOSE (UNII: J2B2A4N98G) MAGNESIUM STEARATE (UNII: 70097M6B0) PO LYDEXTROSE (UNII: 70097M6B0) PO LYDEXTROSE (UNII: 84//Q0SDW1A) PO VIDONE (UNII: 847/758AP) SULCON DIO XIDE (UNII: ETY726XBUA) STRATIC ACTO (UNII: 847/758AP) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATION DIO XIDE (UNII: ETY726XBV) TTAATION DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATE TTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATE TTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: E					
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PO VIDO NE (UNIE F2989GH94E) SILEON DIO XIDE (UNIE F1726 XBU4) STEARIC ACID (UNIE 4EL V726 XBU4) STEARIC ACID (UNIE 4EL V726 5AP) - STEARIC ACID (UNIE 15R3 V2JP) - STEARIC ACID	PO VIDO NE (UNIE PT298 9G H9 4E)	POLYDEXTROSE (UNII	: VH2XOU12IE)			
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4) → · · · · · · · · · · · · · · · · · ·	SILICON DIO XIDE (UNII: ETI7Z6 XBU4)	POLYETHYLENE GLY(C OL (UNII: 3WJQ0SDW1A)			
STEARIC ACID (UNII: 4EL V726 5AP) TTANUOM DIO XIDE (UNII: 15FIX9 V21P) TRIACETIN (UNII: XHX3C3X6 73) Product Characteristics Color white (WHITE) Score no score Shape ROUND Size 100mm Flavor 044;352 Contains 144;352 Contains 144;352 Packag ing # Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:52124-0009-1 2 in 1 PACKET 144;352 Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA 75010 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Soure) 10/10/2015	STEARIC ACID (UNII: 4EL V7265AP) Image: Content of the state of	POVIDONE (UNII: FZ98	9 GH9 4E)			
TTANIUM DIOXIDE (UNE: 15FK9 Y2JP) TRIACETIN (UNE: XHX3C3X673) Product Characteristic status and	TTANUM DIO XIDE (UNIE: ISFIX9 V2/P) TRIACETIN (UNIE: XHX3C3X673)	SILICON DIOXIDE (UN	II: ETJ7Z6XBU4)			
TRIACETIN (UNIE XHX3C3X673) Product Characteristic Color white (WHTE) Score no score Shape ROUND Size 10mm Flavor ROUND Size 44;352 Contains Imprint Code 44;352 Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code Package Description Marketing Start Date Marketing End Date Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Part 3 of 5 MEDIQUE APART STRENGTH acetaminophen tablet, film coated Product Information NDC:47682-175	TRIACETIN (UNIE XHX3G3X673) Note: State S	STEARIC ACID (UNII: 4	ELV7Z65AP)			
Product Characteristics Color vhite (WHITE) Score no score no score Shape ROUND ROUND Size Umprint Code V4:352 Omm V4:352 V V Packaging Item Code Package Description Marketing Start Date Marketing End Date I NDC:52124-0009-1 2 in 1 PACKET V V V V V V V V V V V V V V V V V V V	Product Characteristics view of white (WHTE) core of source of shape and the (WHTE) core core of source o	TITANIUM DIO XIDE (U	NII: 15FIX9V2JP)			
Colorwhite (WHITE)Scoreno scoreShapeROUNDSize10mmFlavorImprint Code (Warden)44,352ContainsImprint Code (Source)Variable (Source)PackagingPackage DescriptionMarketing Start DateMarketing End DateI NDC:52124-0009-12 in 1 PACKETVariable (Source)Marketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DatePart 3 of 5Variable (Source)Variable (Source)Variable (Source)Variable (Source)Marketing Code (Source)MDC:47682-175Variable (Source)Variable (Source)	Colorwhite (WHITE)Scoreno scoreShapeROUNDSize10mmFlavorImprint Code44352ContainsImprint CodeVPackagingImprint CodeNarketing Start DateMarketing End DateI NDC:52124-0009-12 in 1 PACKETImprint CodeMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DatePart 3 of 5STRENGTH01/11/2011Imprint CodeImprint CodeProduct InformativeSTRENGTHStrengen VStrengen VImprint CodeProduct InformativeImprint Code (Source)Imprint Code (Source)Imprint CodeImprint Code	TRIACETIN (UNII: XHX3	C3X673)			
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Shape ROUND Size 10mm Flavor Imprint Code Imprint Code 44,352 Contains V V V Packaging Item Code Package Description Marketing Start Date M Marketing Information 1 NDC:52124-0009-1 2 in 1 PACKET V Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Part 3 of 5 MEDIQUE APAP EXTRENGTH Arburder table, filter STRENGTH Start Strength Product Information Imprint Code (source) MDC:47682-175	Shape ROUND Size 10mm Flavor Imprint Code 44,352 Contains Imprint Code 44,352 Packaging * Item Code Package Description * Item Code 2 in 1 PACKET Marketing Informative Marketing Category Application Number or Monograve Marketing Start Date Marketing Category Application Number or Monograve Marketing Start Date Marketing Category AnDA Andor5010 01/11/2011 Product Informative Kerr STRENGTH MC:47682-175	Product Character	istics			
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Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code 2 in 1 PACKET Image: Comparison of the tem Code Comparison of tem Comparison of tem Code Comparison of	Packaging # Item Code Package Description Marketing Start Date Marketing End Date t NDC:52124-0009-1 2 in 1 PACKET Image: Constant Constan	Flavor		Imprint Co	de	44;352
Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code 2 in 1 PACKET Image: Comparison of the tem Code Comparison of tem Comparison of tem Code Comparison of	Packaging # Item Code Package Description Marketing Start Date Marketing End Date t NDC:52124-0009-1 2 in 1 PACKET Image: Constant Constan	Contains				
Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End D ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information MDC:47682-175	Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information MDC:47682-175					
Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA75010 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate FYTHENGTH ANDA MOLESSION OF THE STRENGTH ANDA MARETEX STRENGTH ANDA MARETEX Product Information MC:47682-175	Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coated Product Information MDC:47682-175		Package Description	Marketii	ng Start Date	Marketing End Date
Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA7551 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate Product Information MDC:47682-175	Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate Product Information MDC:47682-175	# Item Code	.	Marketii	ng Start Date	Marketing End Date
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MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	 Item Code NDC:52124-0009-1 	2 in 1 PACKET			
MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	 Item Code NDC:52124-0009-1 Warketing Info Marketing Category 	2 in 1 PACKET rmation Application Number or Monogr		Marketing Start	
acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	 Item Code NDC:52124-0009-1 Warketing Info Marketing Category 	2 in 1 PACKET rmation Application Number or Monogr		Marketing Start	
acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175	 Item Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA 	2 in 1 PACKET rmation Application Number or Monogr		Marketing Start	
Product Information Item Code (Source) NDC:47682-175	Product Information Item Code (Source) NDC:47682-175	<pre># Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5</pre>	2 in 1 PACKET rmation Application Number or Monogr ANDA0 750 10	aph Citation	Marketing Start	
Item Code (Source) NDC:47682-175	Item Code (Source) NDC:47682-175	Item Code I NDC:52124-0009-1 Marketing Info Info Marketing Category Info ANDA Info Part 3 of 5 Info MEDIQUE APA	2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH	aph Citation	Marketing Start	
Item Code (Source) NDC:47682-175	Item Code (Source) NDC:47682-175	Item Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA	2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH	aph Citation	Marketing Start	
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Route of Administration ORAL	Route of Administration ORAL	# Item Code 1 NDC:52124-0009-1 Marketing Info Info Marketing Category ANDA Part 3 of 5 MEDIQUE AP A Acetaminophen tablet Info	2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH , film coated	aph Citation	Marketing Start	
		# Item Code 1 NDC:52124-0009-1 Marketing Info Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA Acetaminophen tablet Information	2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH , film coated	aph Citation	Marketing Start	
		I tem Code I Tem Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA acetaminophen tablet Product Informatie Item Code (Source)	2 in 1 PACKET Timation Application Number or Monogr ANDA075010 ANDA075010 ANDA075010 NDC:47682-175	aph Citation	Marketing Start	

	Ingredient/Act	•			Basic of C	trongth	Strongel
		Ingredient Name			Basis of S CETAMINO	-	Strength
ACEIAM	IINOPHEN (UNII: 3)	5209ITL9D) (ACETAMINOPHEN - U	NII:362O911L9L)) A	CETAMINO	PHEN	500 mg
Inactiv	e Ingredients						
		Ingredient Nam	e			5	Strength
CROSPO	VIDO NE (UNII: 68	40 19 6 0 MK)					
HYDRO X	YPROPYL CELLU	J LOSE (UNII: RFW2ET671P)					
HYPROM	TELLOSES (UNII: 3	3NXW29V3WO)					
MAGNES	IUM STEARATE (UNII: 70097M6I30)					
CELLUL	OSE, MICROCRYS	STALLINE (UNII: OP1R32D61U)					
MINERAI	L OIL (UNII: T5L87	[28 FGP)					
POLYET	HYLENE GLYCOI	L (UNII: 3WJQ0SDW1A)					
POVIDO	NE (UNII: FZ989GH	194E)					
PROPYL	ENE GLYCOL (UN	NII: 6DC9Q167V3)					
	DIO XIDE (UNII: E						
		ULOSE SODIUM (UNII: K6790BS31	1)				
	C ACID (UNII: 4ELV	7Z65AP)					
TALC (U	NII: 7SEV7J4R1U)						
TITANIU	M DIO XIDE (UNII:	15FIX9V2JP)					
		X673)					
Produc	t Characterist						
	t Characterist		Score			no score	
Color	t Characterist	ics	Score Size			no score 12mm	
Color Shape	t Characterist	ics white (WHITE)		e			
Color Shape		ics white (WHITE)	Size	le		12mm	
Color Shape Flavor Contains	5	ics white (WHITE)	Size	le		12mm	
Color Shape Flavor Contains Packag	ing	ics white (WHITE) ROUND	Size Imprint Cod		Ма	12mm AZ;235	and Date
Color Shape Flavor Contains Packag #	5	ics white (WHITE)	Size Imprint Cod	le g Start Date	Ma	12mm	End Date
Color Shape Flavor Contains Packag # 1 NDC:4	s ing Item Code	ics white (WHITE) ROUND Package Description 2 in 1 PACKET	Size Imprint Cod		Ma	12mm AZ;235	End Date
Color Shape Flavor Contains Packag # 1 NDC:4	s ing Item Code 7682-175-46	ics white (WHITE) ROUND Package Description 2 in 1 PACKET	Size Imprint Cod			12mm AZ;235	End Date
Color Shape Flavor Contains Packag # 1 NDC:4 ¹ Marke	s ing Item Code 7682-175-46 eting Inforn ting Category	ics white (WHITE) ROUND Package Description 2 in 1 PACKET	Size Imprint Cod	g Start Date		12mm AZ;235	
Color Shape Flavor Contains Packag # 1 NDC:4 Marke	s ing Item Code 7682-175-46 eting Inforn ting Category	ics white (WHITE) ROUND Package Description 2 in 1 PACKET Application Number or Monogr	Size Imprint Cod	g Start Date Marketing S		12mm AZ;235	
Color Shape Flavor Contains Packag # 1 NDC:4 Marke OTC mon	s ing Item Code 7682-175-46 eting Inform ting Category tograph not final	ics white (WHITE) ROUND Package Description 2 in 1 PACKET Application Number or Monogr	Size Imprint Cod	g Start Date Marketing S		12mm AZ;235	
Color Shape Flavor Contains Packag # 1 NDC:4 Marke OTC mon	s ing Item Code 7682-175-46 eting Inform ting Category tograph not final final	ics white (WHITE) ROUND Package Description 2 in 1 PACKET hation Application Number or Monogr part343	Size Imprint Cod	g Start Date Marketing S		12mm AZ;235	
Color Shape Flavor Contains Packag # 1 NDC:43 Marke OTC mon	s ing Item Code 7682-175-46 eting Inforn ting Category tograph not final a of 5 SEPTIC TO	ics white (WHITE) ROUND Package Description 2 in 1 PACKET Application Number or Monogr part343 WELETTE	Size Imprint Cod	g Start Date Marketing S		12mm AZ;235	
Color Shape Flavor Contains Packag # 1 NDC:43 Marke OTC mon	s ing Item Code 7682-175-46 eting Inform ting Category tograph not final final	ics white (WHITE) ROUND Package Description 2 in 1 PACKET Application Number or Monogr part343 WELETTE	Size Imprint Cod	g Start Date Marketing S		12mm AZ;235	

Product Information							
Item Code (Source)		NDC:52124-0001					
Route of Administration		TOPICAL					
Active Ingradiant/Ac	tivo Moi	a 4=7					
Active Ingredient/Ac		ety dient Name		P	Basis of Stren	ath	Strength
BENZALKONIUM CHLOR	-		KONIUM -		ZALKONIUM	5.11	0.40 mL
UNII:7N6JUD5X6Y)				CHL	ORIDE		in 100 mL
Inactive Ingredients							
	I	ngredient Name				Stre	ngth
WATER (UNII: 059QF0KO0	R)						
Packaging # Item Code	Daal	age Deceription	Markating	Start Da		[aultatis	a End Data
# Item Code 1 NDC:52124-0001-1		Cage Description 1 PACKAGE	Marketing	Start Da		larketii	ng End Date
I NDC.52124-0001-1	0.0 IIIL III	IFACKAGE					
Marketing Inform	nation						
U		on Number or Monogra	aph Citation I	Marketiı	ng Start Date	Mark	keting End Date
	rt333E		_	1/11/2011	-		-
Part 5 of 5							
ALCOHOL CLEA	ANSINC	G PAD					
isopropyl alcohol liquid							
Product Information							
Item Code (Source)		NDC:52124-0002					
Route of Administration		TOPICAL					
Active Ingredient/Ac	tive Moi	ety					
		redient Name			Basis of Str	ength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M	416302) (ISOPROPYL AL	COHOL -		ISOPROPYL		70 mL
UNII:ND2M416302)					ALCOHOL		in 100 mL
Inactive Ingredients							
	I	ngredient Name				Stre	ngth
WATER (UNII: 059QF0KO0	R)						

Packaging							
#	Item Code	Package Description	Marketing	g Start Date	Ma	rketing End Date	
1 ND	C:52124-0002-1	0.5 mL in 1 PACKAGE					
Ма	keting Infor	nation					
	0						
Mar	rketing Category	Application Number or Monog	raph Citation	Marketing Star	t Date	Marketing End Date	
OTC n	nonograph not final	part333A		0 1/11/20 11			
Maı	cketing Inform	mation					
	cketing Infor Keting Category	nation Application Number or Monog	raph Citation	Marketing Star	t Date	Marketing End Date	

Labeler - Genuine First Aid, LLC (619609857)

Revised: 1/2011

Genuine First Aid, LLC