10 PERSON ANSI - benzalkonium chloride, lidocaine, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, water, benzocaine, alcohol, ibuprofen, acetaminophen, aspirin 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.

10Person ANSI

ACTIVE INGREDIENTS:

Benzalkonium Chloride 0.13% Lidocaine HCL 0.5%

Purpose: First aid antiseptic, external analgesic

Uses: First aid to help prevent infection and for the temporary relief of pain and itching associated with:

Minor Cuts

Scrapes

Burns

Warnings:

For external use only

Do not use: In eyes, in large quantities, over raw blistered areas, or on deep puncture wounds, animal bites or serious burns, for more than one week

Keep out of reach of Children.

If ingested, contact a Poison Control Center right away.

Directions: Clean affected area, Apply small amount not more than 3 times daily.

May be covered with a sterile bandage.

Other Information:

Store at room temperature

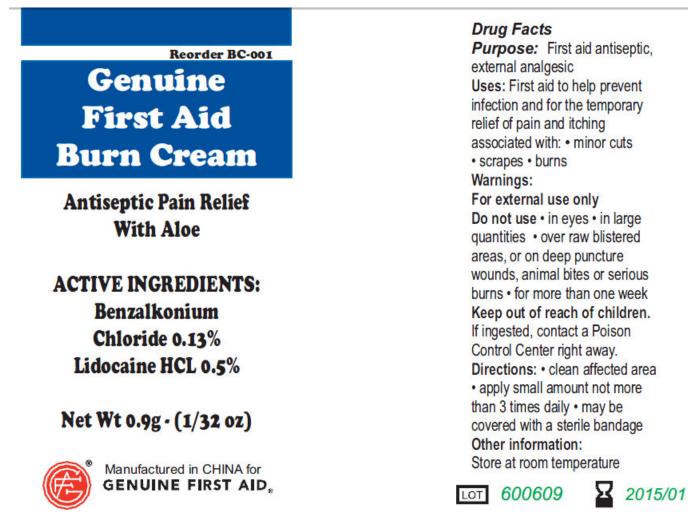
Genuine First Aid Burn Cream

Antiseptic Pain Relief With Aloe

Net Wt 0.9g (1/32 oz)

Manufactured in CHINA for

Genuine First Aid.



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

GENUINE FIRST AID

1/pouch



DRUG FACTS - Antiseptic Towelette

Active Ingredient:	Purpose:	
Benzalkonium Chloride 0.40%First Aid		
Use: For Professional and Hospital use	. Helps	
prevent infection. Antiseptic cleansing of	of face,	1
hands and body without soap and water		1
Warnings: For external use only. Keep	out of	ш
reach of children. If swallowed, get med	dical help	2
or contact a Poison Control Center right	away.	ERE
If unusual redness, swelling or other syr	nptoms	王
occur, consult a physician immediately.		
Do not use: In the eyes, or over large a	reas of the	TEAR
body.		4
Directions: Tear open packet, unfold to	welette	ш
and use to cleanse desired skin area.	Discard	F
towelette appropriately after single use.	Jiscaru	
		1
Inactive ingredient: Purified water.		
Made	in CHINA	

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

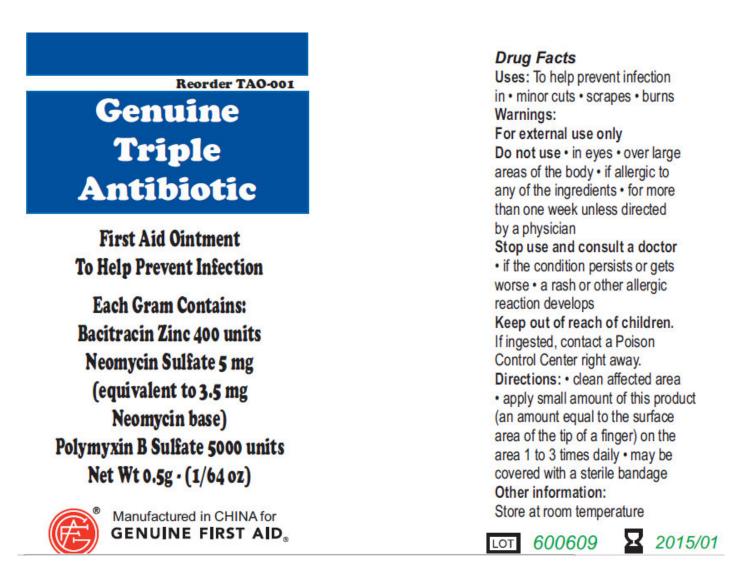
Neomycin base)

Polymyxin B Sulfate 5000 units

Net Wt. 0.5g ; (1/64 oz)

Manufactured in CHINA for

GENUINE FIRST AID.



Active Ingredient:

Purified Water USPq.s. Inactive Ingredients:	
Sodium Chloride USP	44mg
Monobasic Sodium Phosphate USP Sodium Phosphate Dibasic USP	18mg 111mg
Edetate Disodium USP	10 mg
Benzalkonium Chloride NF (as preservative)	0.5mg
Store in a cool place. For irrigation only.	
Discard unused portion of the solution.	
Not for injection.	

Warning:

If you experience eye pain, changes in vision, continued redness or irritation of the eye,

or if the condition worsens or persists for more than 72 hours, discontinue use and consult a doctor.

Do not use if solution changes color or becomes cloudy.

Directions

Remove contacts before using.

Twist top to remove.

Flush the affected area as needed. Control

Rate of flow by pressure on the bottle. Do not touch

tip of the container to any surface. Do not reuse.

If necessary continue flushing with emergency eyewash or shower.

Discard bottle after use.

Uses:

For flushing or irrigating the eyes to

remove loose foreign material, air pollutants,

or chlorinated water.

Code No.: GUJ/DRUG/G/1080

Batch No.:

Mfg Date:

Exp: Date:

10 ml

Sterile Isotonic Buffered Genuine

Eyewash

For single use only



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



Insect Sting

REORDER ISRP-001

DRUG FACTS - Insect Sting Relief Pad

Active Ingredient: Purpose	
Benzocaine, 6% w/v Topical Anesthetic SD alcohol, 60% w/v Antisepti	
Use: For the temporary relief of pain and itching	ĩ
associated with minor burns, scrapes and insect bites.	ш
Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	R
Flammable - keep away from fire or flame. Avoid contact with eyes. If this happens, rinse thoroughly with water.	TEAR
Do not use: In eyes, on broken skin, deep puncture wounds. If unusual redness, swelling,	F
irritation or other symptoms occur, consult a physician immediately.	÷
Made in CHINA	· '

Active ingredient (in each tablet)

Purpose

Ibuprofen 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.

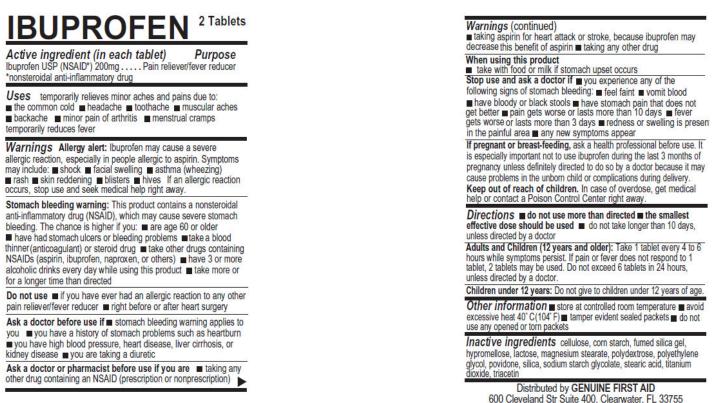
Children under 12 years: Do not give to children under 12 years of age.

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



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 evenu 4 to 6 bours as
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 Farenh	eat (15-30 degree Celcius)
tamper evident sealed packets	; do not use any open or torn packets
Distributed by GENUINE FIF 600 Cleveland Str Suite 400,	
GENUINE FIRST AID	2 Tablets

NON-ASPIRIN

GENUINE FIRST AID.	Warnings (cont	inued)
NON–ASPIRIN Active ingredient (in each tablet) Purpose	pain gets worse or la worse or lasts for mo	doctor if ■ symptoms do not improve ■ ists for more than 10 days ■ fever gets ire than 3 days ■ new symptoms occur ■ s present ■ a rare sensitivity reaction occurs
Acetaminophen 325 mg Analgesic/antipyretic Uses temporary relief of minor aches and pains associated with ■ common cold ■ headache ■ toothache	Keep out of reach o contact a doctor or P medical attention is o	Leeding, ask a health professional before use of children. In case of accidental overdose, toison Control Center immediately. Prompt xritical for adults as well as for children even i y signs or symptoms. Do not exceed age.
 muscular aches ■ backache ■ arthritis menstrual cramps ■ and reduction of fever 	Directions	
Warnings Liver warning: This product contains acetaminophen.	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.
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	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.
Do not use I with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen,	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.
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		ON ■ store at 59°-86°F (15°-30°C) ■ tamp ets ■ do not use any open or torn packets ients corn starch, hydroxypropyl
Ask a doctor before use if the user has liver disease	methylcellulose, poly	ethylene glycol, pregelatinized starch, stearic
Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin		ovidone and sodium starch glycolates.
		uted by GENUINE FIRST AID d Str Suite 400, Clearwater, FL 33755

Active Ingredient (in each tablet)

Purpose

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

*nonsteroidal anti-inflammatory drug

Temporarily relieves minor aches and pains associated with: Uses 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.

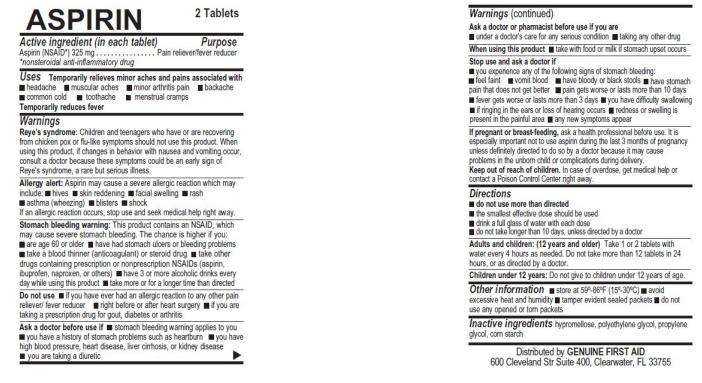
Store at 59 - 86 degree Fahrenheit (15 - 30 degree Celsius); avoid

excessive heat and humidity; tamper evident sealed packets;

Do not use any opened or torn packets Inactive Ingredients: hypromellose, polyethylene glycol, propylene glycol, corn starch Distributed by GENUINE FIRST AID

600 Clevelad Str Suite 400, Clearwater, FL 33755

ASPIRIN 2 Tablets



ANSI/ISEA Z308.1-2009 TYPE III

Caution! This Kit meets ANSI/ISEA Z308.1-2009 only when required minimum fill is maintained with first aid products marked "ANSI/ISEA Z308.1-2009."

CARRYING CASE 1 Easy Access Pocket System 1 Hard Case

POCKET1: SMALL CUTS AND BURNS 16 Adhesive Plastic Bandages 1"x3" 1 Burn Cream 5 Antiseptic Towelettes 6 Triple Antibiotic Ointment 0.5gr 10 Cotton Tipped Applicators

POCKET 2: MEDIUM CUTS AND SCRATCHES 5 Antiseptic Towelettes

4 Sterile Gauze Pad 3"x3" 1 Roller Gauze Bandage 2"X4.1yds 1 Sterile Eye Pads 1 Sterile Eye Wash 10ml Twist Top

POCKET 3: SEVERE BLEEDING AND BURNS 1 First Aid Tape Roll 1/2"x5 yds. 1 Combine Pad 5"X9"

POCKET 4: CPR 1 CPR Breathing Barrier

POCKET 5: PROTECTION 2 Medical Grade Vinyl Gloves

1 Insect Sting Relief Pads 1 Triangular Bandage 42"x42"x59" 2 Ibuprofen 200mg 2Non Aspirin 325mg 2 Aspirin 325mg

POCKET 6: INSTRUMENTS 1 Emergency First Aid Guide **1** Plastic Tweezers 1 Scissors

Manufactured in China for:

Genuine First Aid LLC. 600 Cleveland Street Suite 400 Clearwater FL 33755

www.GenuineFirstAid.com

GENUINE FIRST AID

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10 PERSON ANSI								
benzalkonium chloride, lidocaine, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, water, benzocaine,								
alcohol, ibuprofen, acetaminophen, aspirin kit								
Product Information								
Product Type HUMAN OTC DRUG Item Code (Source) NDC:52124-0111								

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC	2:52124-0111-1	1 in 1 KIT				
Quan	tity of Parts					
Quan Part #		ickage Quantity	Total Produc	t Quantity		
		nckage Quantity	Total Produc 5.4 g	t Quantity		
Part #	6 PACKAGE	ickage Quantity		t Quantity		
Part # Part 1	e Pa 6 PACKAGE 10 PACKAGE	nckage Quantity	5.4 g	t Quantity		

0.5 mL

2 2

2

Part 5 1 PACKAGEPart 6 1 PACKAGE

Part 71 PACKAGEPart 81 PACKAGE

GENUINE FIRST AID BURN ANTISEPTIC PAIN RELIEF WITH ALOE

benzalkonium chloride, lidocaine cream

Product Information

Item Code (Source)	NDC:52124-0004
Route of Administration	TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g			
LIDO CAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.5 in 100 g			

Pack	Packaging						
#	Item Code	Package Description	Marketin	g Start Date	Ma	arketing End Date	
1 ND	C:52124-0004-1	0.9 g in 1 PACKAGE					
Marketing Information							
Mar	keting Category	ategory Application Number or Monograph Citation		Marketing Start	Date	Marketing End Date	
OTC -	nonograph final	part345		04/24/2010			

Part 2 of 8					
ANTISEPTIC TOWELE	TTE				
benzalkonium chloride liquid					
Product Information					
Item Code (Source)	NDC:52124-0001				
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
Ingre	dient Name		Basis o	f Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F UNII:7N6JUD5X6Y)	5UM2KM3W7) (BENZAL	KONIUM -	BENZALKO CHLORIDE		0.40 mL in 100 mL
Inactive Ingredients					
I	ngredient Name			Stre	ngth
WATER (UNII: 059QF0KO0R)					
Packaging					
	age Description	Marketing St	tart Date	Marketi	ng End Date
1 NDC:52124-0001-1 0.8 mL in	1 PACKAGE				
Marketing Information					
Marketing Category Application	on Number or Monogra	aph Citation M	arketing Sta	rt Date Mar	keting End Date
OTC monograph final part333 04/24					
Part 3 of 8					
GENUINE TRIPLE ANT	TIBIOTIC				
bacitracin zinc,neomycin sulfate,polymyxin b sulfate ointment					
Product Information					
Item Code (Source)	NDC:52124-0003				
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
•	edient Name		Basis	of Strength	Strength

Packaging # Item Code Package Description Marketing Start Date 1 NDC:52124-0003-1 0.5 g in 1 TUBE INDC:52124-0003-1 0.5 g in 1 TUBE Marketing Information Marketing Category Application Number or Monograph Citation Marketing OTC monograph final par333 0424/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid NDC:52124-0005 Route of Administration OPHTHALMIC Ingredient Name Basis of S Marketing curve Molety Ingredient Name Basis of S NDC:52124-0005 Route of Administration OPHTHALMIC Ingredient Name Basis of S Marketing Clunk: 059QF0K00R) (WATER - UNIK 059QF0K00R) WATER Ingredient Name Basis of S SOTUM CHLORIDE (UNIK: 451W47/1Q8X) SOTUM CHLORIDE (UNIK: 451W47/1Q8X) SOTUM CHLORIDE (UNIK: 7FLD9 IC86K) BERZALKONIUM CHLORIDE (UNIK: 7FLD9	NEOMYCIN SULFATE					
Item Code Package Description Marketing Start Date 1 NDC:52124-0003-1 0.5 g in 1 TUBE Marketing Start Date Marketing Information Application Number or Monograph Citation Marketing Or Marketing Category Application Number or Monograph Citation Marketing Category Application Number or Monograph Citation Marketing Category Marketing Category Marketing Category Application Number or Monograph Citation Marketing Category 0.4/24/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information NDC:52124-0005 OPHTHALMIC Route of Administration OPHTHALMIC Soperation Server Serve	POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) POLYMYXIN B SULF					
Item Code Package Description Marketing Start Date 1 NDC:52124-0003-1 0.5 g in 1 TUBE Marketing Start Date Marketing Information Application Number or Monograph Citation Marketing Or Marketing Category Application Number or Monograph Citation Marketing Category Application Number or Monograph Citation Marketing Category Marketing Category Marketing Category Application Number or Monograph Citation Marketing Category 0.4/24/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information NDC:52124-0005 OPHTHALMIC Route of Administration OPHTHALMIC Soperation Server Serve						
I NDC:52124-0003-1 0.5 g in I TUBE Marketing Information Marketing Category Application Number or Monograph Citation Marketing Marketing Category Application Number or Monograph Citation Marketing OTC monograph final part333 04/24/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information Item Code (Source) NDC:52124-0005 Route of Administration OPHTHALMIC Application Number or Monograph Citation Ingredient/Active Moiety Ingredient/Active Moiety Ingredient Name Marketing Colspan="2">Basis of S WATER (UNIE 059QF0KOOR) (WATER - UNIE059QF0KOOR) WATER Ingredient Name SODIUM CHLORIDE (UNIE 451W471Q8X) SODIUM PHOSPHATE, DIBASIC (UNIE 3980JIH2SW) SODIUM PHOSPHATE, DIBASIC (UNIE GR6861.BA74) SODIUM CHLORIDE	M	faultatiu	a End Data			
Marketing Information Marketing Category Application Number or Monograph Citation Marketing OTC monograph final part333 04/24/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information Item Code (Source) NDC:52124-0005 Route of Administration OHTHALMIC Active Ingredient/Active Moiety Ingredient Name Basis of S MATER (UNIE 059QF0KOOR) (WATER - UNIE059QF0KOOR) WATER Ingredient Name Basis of S SODIUM CHLORIDE (UNIE 451W471Q8X) SODIUM PHOSPHATE, DIBASIC (UNIE 3980JH25W) SODIUM PHOSPHATE, DIBASIC (UNIE 3980JH2	IVI	larketin	g End Date			
Marketing Category OTC monograph finalApplication Number or Monograph Citation 04/24/2010OTC monograph finalpart33304/24/2010Part 4 of 8STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquidProduct InformationNDC:52124-0005Route of AdministrationNDC:52124-0005Route of AdministrationOPHTHALMICActive Ingredient/Active MoietyBasis of SWATER (UNIE 059QF0KOOR) (WATER - UNIE 059QF0KOOR)WATERIngredientsIngredientsIngredientsIngredient NameBasis of SSOTIUM CHLORIDE (UNIE 451W47108 X)SOTIUM PHO SPHATE, MONOBASIC (UNIE 3980JH2SW)SOTIUM PHO SPHATE, DIBASIC (UNIE 3880JH2SW)SOTIUM PHO SPHATE, DIBASIC (UNIE STUDE USE)BAZALKONUM CHLORIDE (UNIE FFUD21C8K)BEZALKONUM CHLORIDE (UNIE FFUD21C8K)BAZALKONUM CHLORIDE (UNIE FFUD21C8K)BAZALKONUM CHLORIDE (UNIE FFUD21C8K)PackagingItem CodePackaging						
OTC monograph final part333 04/24/2010 Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information Item Code (Source) NDC:52124-0005 Route of Administration OPHTHALMIC Active Ingredient/Active Moiety Active Ingredient/Active Moiety Active Ingredient/Active Moiety Ingredient Name Basis of S WATER (UNIE 059QF0KOOR) (WATER - UNIE059QF0KOOR) WATER Inactive Ingredients Ingredient Name SOIUM CHLORIDE (UNIE 451W471Q8 X) SOIUM PHOSPHATE, MONOBASIC (UNIE 3980JIH2SW) SOIUM PHOSPHATE, DIBASIC (UNIE GR686LBA74) EDETATE DISO DIUM (UNIE 7FLD91CB6K) BENZALKONIUM CHLORIDE (UNIE FSUMZKM3W7) Packaging # Item Code Package Description Marketing Start Date						
Part 4 of 8 STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information Item Code (Source) NDC:52124-0005 Route of Administration OPHTHALMIC Active Ingredient/Active Moiety Ingredient Name Basis of S WATER (UNIE 059QF0K00R) (WATER - UNIE059QF0K00R) WATER Ingredient Name Ingredient Name Solum CHLORIDE (UNIE 451W471Q8X) SODIUM CHLORIDE (UNIE 451W471Q8X) SODIUM PHO SPHATE, MO NOBASIC (UNIE: 3980 JIH2SW) SODIUM PHO SPHATE, DIBASIC (UNIE: GR686LBA74) EDETATE DISODIUM (UNIE: 7FLD91C8 6K) BENZALKONIUM CHLORIDE (UNIE: FSUM2KM3W7) Packaging Marketing Colspan="2">Colspan="2">Marketing Start Date	tart Date	Mark	eting End Date			
Number of AdministrationNDC:52124-0005NDC:52124-0005NDC:52124-0005NDC:52124-0005OPHTHALMICOPHTHALMICOPHTHALMICNameBasis of SWATER (UNII: 059QF0KO0R) (WATER - UNI:059QF0KO0R)Marter (UNII: 059QF0KO0R) (WATER - UNI:059QF0KO0R)WATERIngredient NameBasis of SSOLUM CHLORIDE (UNII: 451W471Q8X)SOLUM PHOSPHATE, DIBASIC (UNII: 3980JH25W)SOLUM PHOSPHATE, DIBASIC (UNII: 5780JH25W)SOLUM PHOSPHATE, SOLUM PHOSPHATE, SOL			-			
STERILE ISOTONIC BUFFERED GENUINE EYEWASH water liquid Product Information Item Code (Source) NDC:52124-0005 Route of Administration OPHTHALMIC Active Ingredient/Active Moiety Active Ingredient/Active Moiety Active Ingredient/Active Moiety Ingredient Name Basis of S WATER (UNII: 059QF0KOOR) (WATER - UNII:059QF0KOOR) WATER MATER (UNII: 059QF0KOOR) (WATER - UNII:059QF0KOOR) Inactive Ingredients Ingredient Name SO DIUM CHLORIDE (UNII: 451W471Q8X) SO DIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW) SO DIUM PHOSPHATE, DIBASIC (UNII: 3980JIH2SW) SO DIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74) EDETATE DISO DIUM (UNII: 7FLD91C86K) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) Packaging M Item Code Package Description Marketing Start Date						
Number of AdministrationNDC:52124-0005NDC:52124-0005NDC:52124-0005NDC:52124-0005OPHTHALMICOPHTHALMICOPHTHALMICNameBasis of SWATER (UNII: 059QF0KO0R) (WATER - UNI:059QF0KO0R)Marter (UNII: 059QF0KO0R) (WATER - UNI:059QF0KO0R)WATERIngredient NameBasis of SSOLUM CHLORIDE (UNII: 451W471Q8X)SOLUM PHOSPHATE, DIBASIC (UNII: 3980JH25W)SOLUM PHOSPHATE, DIBASIC (UNII: 5780JH25W)SOLUM PHOSPHATE, SOLUM PHOSPHATE, SOL						
Product Information NDC:52124-0005 Route of Administration OPHTHALMIC Active Ingredient/Active Moiety Basis of S Active Ingredient/Active Moiety Basis of S WATER (UNIE 059QF0KO0R) (WATER - UNIE059QF0KO0R) WATER Ingredient Name Basis of S WATER (UNIE 059QF0KO0R) (WATER - UNIE059QF0KO0R) WATER Ingredient Name WATER SODIUM CHLORIDE (UNIE 451W471Q8X) WATER SODIUM PHO SPHATE, MONDASIC (UNIE 3980JH2SW) SODIUM PHO SPHATE, DIBASIC (UNIE 3980JH2SW) SODIUM PHO SPHATE, DIBASIC (UNIE GR686LBA74) EDETATE DISODIUM (UNIE 7FLD91C86K) BENZALKONIUM CHLORIDE (UNIE 75UM2KM3W7) FORGARIA (CONIE) Packaging # Item Code Package Description Marketing Start Date						
Iven Code (Source) NDC:52124-0005 Route of Administration OPHTHALMIC Active Ingredient/Active Wolver Karter (UNII: 059QF0KO0R) WATER (UNII: 059QF0KO0R) WATER (UNII: 059QF0KO0R) WATER (UNII: 059QF0KO0R) WATER UNII: 059QF0KO0R) WATER VATER UNII: 059QF0KO0R) WATER UNII: 059QF0KO0R) WATER WATER UNII: 059QF0KO0R) WATER WATER UNII: 059QF0KO0R) WATER WATER WATER WATER UNII: 059QF0KO0R) WATER WATE						
Image: Image						
OPHTHALMIC Route of Administration OPHTHALMIC Ingredient/Active Moiety Ingredient Name Basis of S WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R) WATER WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R) WATER Ingredient Name Basis of S WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R) WATER Ingredient Name Solum ChloRide (UNII: 451W47IQ8X) SO DIUM CHLORIDE (UNII: 451W47IQ8X) SO DIUM PHO SPHATE, DIBASIC (UNII: 3980 JIH2SW) SO DIUM PHO SPHATE, DIBASIC (UNII: 3980 JIH2SW) SO DIUM PHO SPHATE, DIBASIC (UNII: GR6866LBA74) EDETATE DISODIUM (UNII: 7FLD91C86K) BENZALKONIUM CHLORIDE (UNII: FSUM2KM3W7) Y Kaging # Item Code Package Description						
Active Ingredient/Active Moiety Basis of S Ingredient Name Basis of S WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R) WATER Ingredients Ingredient Name Ingredients Ingredient Name						
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)WATERWATERInactive IngredientsIngredient NameSOTUM CHLORIDE (UNII: 451W47IQ8X)SOTUM PHO SPHATE, MONOBASIC (UNII: 3980JIH2SW)SOTUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)EDETATE DISO DIUM (UNII: 7FLD9 1C86K)BENZALKO NIUM CHLORIDE (UNII: F5UM2KM3W7)Packaging#Item CodePackage DescriptionMarketing Start Date						
Ingredient NameBasis of SWATER (UNII: 059QF0KOOR) (WATER - UNII:059QF0KOOR)WATERInactive IngredientsIngredient NameIngredient NameSO JUM CHLORIDE (UNII: 451W47IQ8X)SO JUM PHO SPHATE, MO NOBASIC (UNII: 3980JIH2SW)SO JUM PHO SPHATE, DIBASIC (UNII: 3980JIH2SW)SO JUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)EDETATE DISO DIUM (UNII: 7FLD9 1C86K)BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)Packaging#Item CodePackage DescriptionMarketing Start Date						
WATER (UNII: 059QF0K00R) (WATER - UNII:059QF0K00R)WATERWATERInactive IngredientsIngredient NameSO JIUM CHLORIDE (UNII: 451W47IQ8X)SO JIUM PHO SPHATE, MO NO BASIC (UNII: 3980JIH2SW)SO JIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)EDETATE DISO DIUM (UNII: 7FLD91C86K)BENZALKO NIUM CHLORIDE (UNII: F5UM2KM3W7)Packaging#Item CodePackage DescriptionMarketing Start Date						
Ingredients Ingredient Name SO JIUM CHLO RIDE (UNII: 451W47IQ8 X) SO DIUM CHLO RIDE (UNII: 451W47IQ8 X) SO DIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW) SO DIUM PHO SPHATE, DIBASIC (UNII: 3980 JIH2SW) SO DIUM PHO SPHATE, DIBASIC (UNII: GR686 LBA74) EDETATE DISO DIUM (UNII: 7FLD9 1C86 K) BENZ ALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description	rength		Strength			
Ingredient Name SO JUM CHLO RIDE (UNII: 451W47IQ8 X) SO JUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2S W) SO JUM PHO SPHATE, DIBASIC (UNII: GR686 LB A74) EDETATE DISO DIUM (UNII: 7FLD9 1C86 K) BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date		98.6 m	L in 100 mL			
Ingredient Name SO JUM CHLO RIDE (UNII: 451W47IQ8 X) SO JUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2S W) SO JUM PHO SPHATE, DIBASIC (UNII: GR686 LB A74) EDETATE DISO DIUM (UNII: 7FLD9 1C86 K) BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date						
SO JIUM CHLORIDE (UNII: 451W47IQ8 X) SO JIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2S W) SO DIUM PHO SPHATE, DIBASIC (UNII: GR686 LBA74) EDETATE DISO DIUM (UNII: 7FLD9 1C86 K) BENZALKO NIUM CHLORIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date						
SO DIUM PHO SPHATE, MONOBASIC (UNII: 3980 JIH2SW) SO DIUM PHO SPHATE, DIBASIC (UNII: GR686 LBA74) EDETATE DISO DIUM (UNII: 7FLD91C86 K) BENZALKO NIUM CHLORIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date			Strength			
SO DIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74) EDETATE DISO DIUM (UNII: 7FLD9 1C86K) BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date						
EDETATE DISODIUM (UNII: 7FLD9 1C8 6 K) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date						
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) Packaging # Item Code Package Description Marketing Start Date						
Packaging # Item Code Package Description Marketing Start Date						
# Item Code Package Description Marketing Start Date						
# Item Code Package Description Marketing Start Date						
	N.4.	farlest-	a End Data			
	IVL	iar ke tin	g End Date			

Marketing Info	rmation						
Marketing Category					Marketing End Date		
TC monograph final part349				04/24/2010			
Part 5 of 8							
INSECT STINC		F PAD					
Product Informati	on						
Item Code (Source)		NDC:52124-0008					
Route of Administrati	on	TOPICAL					
Active Ingredient/		•					
DENZO CADIE (UNIL UN	•	dient Name		Basis of Strength	_		
	ENZOCAINE - UNII:U3RSY48JW5) COHOL - UNII:3K9958V90M)		BENZOCAINE	6 mL in 100 mL 60 mL in 100 mL			
Packaging # Item Code	Pacl	kage Description Market	ting Sta	rt Date Ma	rketing End Date		
1 NDC:52124-0008-1	0.5 mL in	1 PACKAGE					
Marketing Info	rmation						
Marketing Category	Applicatio	on Number or Monograph Citation	Mar	keting Start Date	Marketing End Date		
OTC monograph final	part348		04/24	/2010			
Part 6 of 8							
Part 6 of 8 IBUPROFEN							
IBUPROFEN							
IBUPROFEN ibuprofen tablet	on						
IBUPROFEN ibuprofen tablet Product Informati	on	NDC:52124-0009					
		NDC:52124-0009 ORAL					
IBUPROFEN ibuprofen tablet Product Informati Item Code (Source)							
IBUPROFEN ibuprofen tablet Product Informati Item Code (Source)	on	ORAL					

IBUPROFEN

Inactive Ingredien	its					
		Ingredient Name				Strength
PO WDERED CELLULO	DSE (UNII: SMD	1X3XO9M)				
STARCH, CORN (UNII: (08232NY3SJ)					
HYPROMELLOSE (UNI	II: 3NXW29V3V	VO)				
LACTOSE (UNII: J2B2A	4N98G)					
MAGNESIUM STEARAT	TE (UNII: 7009	7M6I30)				
POLYDEXTROSE (UNI	II: VH2XOU12IE)				
POLYETHYLENE GLY	COL (UNII: 3W	'JQ0SDW1A)				
POVIDONE (UNII: FZ98	89 GH9 4E)					
SILICON DIO XIDE (UN	NII: ETJ7Z6XBU	(4)				
STEARIC ACID (UNII: 4)	ELV7Z65AP)					
TITANIUM DIO XIDE (U	JNII: 15FIX9V2J	P)				
TRIACETIN (UNII: XHX)						
Product Character						
Color	white (Wh	ite)	Score]	no score
Shape	ROUND		Size		-	10 mm
Flavor			Imprint Code		4	44;352
Contains						
Packaging						
	Pac	kage Description	Marketii	ng Start Date	Mai	rketing End Date
# Item Code	Pacl 2 in 1 PAC	kage Description CKAGE	Marketin	ng Start Date	Mai	rketing End Date
<pre># Item Code 1 NDC:52124-0009-1</pre>	2 in 1 PAC		Marketin	ıg Start Date	Mar	rketing End Date
<pre># Item Code 1 NDC:52124-0009-1</pre>	2 in 1 PAC			ng Start Date Marketing Start		rketing End Date Marketing End Dat
1 NDC:52124-0009-1 Marketing Info	2 in 1 PAC	CKAGE on Number or Monogra		-		
 # Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA 	2 in 1 PAC ormation Application	CKAGE on Number or Monogra		Marketing Start		
 # Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA 	2 in 1 PAC ormation Application	CKAGE on Number or Monogra		Marketing Start		
# Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA	2 in 1 PAC Prmation Application ANDA075010	CKAGE on Number or Monogra		Marketing Start		
<pre># Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN</pre>	2 in 1 PAC Application ANDA0 750 10	CKAGE on Number or Monogra		Marketing Start		
<pre># Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN</pre>	2 in 1 PAC Application ANDA0 750 10	CKAGE on Number or Monogra		Marketing Start		
 # Item Code MDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN acetaminophen tablet 	2 in 1 PAC Applicatio ANDA0 750 10	CKAGE on Number or Monogra		Marketing Start		
<pre># Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN</pre>	2 in 1 PAC Applicatio ANDA0 750 10	CKAGE on Number or Monogra		Marketing Start		
# Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN acetaminophen tablet Product Informati	2 in 1 PAC Applicatio ANDA075010 t	CKAGE		Marketing Start		
# Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 7 of 8 NON-ASPIRIN acetaminophen tablet Product Informati Item Code (Source)	2 in 1 PAC Applicatio ANDA075010 t	CKAGE		Marketing Start		

Active Ingredient/A		5					
Ingredient Name Basis of S						Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINO					OPHEN	325 mg	
Inactive Ingredient	ts						
Ingredient Name							rength
STARCH, CORN (UNII: C)8232NY3SJ)						
POLYETHYLENE GLYC		JQ0SDW1A)					
STEARIC ACID (UNII: 4E							
POVIDONE (UNII: FZ989	JGH94E)						
Product Character	istics						
Color	white (WHI	TE)	Score			no score	
Shape	ROUND		Size			11mm	
Flavor			Imprint Co	de		AZ;234	
Contains							
Packaging							
	Pack	age Description	Marketin	g Start Date	М	arketing H	End Date
# Item Code	Pack 2 in 1 PAC	xage Description KAGE	Marketin	g Start Date	Μ	arketing I	End Date
 # Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category 	2 in 1 PAC rmation Applicatio			Marketing			
NDC:52124-0010-1 Marketing Information	2 in 1 PAC	KAGE		-			
 # Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category OTC monograph final 	2 in 1 PAC rmation Applicatio	KAGE		Marketing			
 # Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category 	2 in 1 PAC rmation Applicatio	KAGE		Marketing			
# Item Code 1 NDC:52124-0010-1 Marketing Infor Marketing Category OTC monograph final Part 8 of 8	2 in 1 PAC rmation Applicatio	KAGE		Marketing			
<pre># Item Code 1 NDC:52124-0010-1 Marketing Info Marketing Category OTC monograph final Part 8 of 8 ASPIRIN</pre>	2 in 1 PAC rmation Applicatio	KAGE		Marketing			
# Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet	2 in 1 PAC	KAGE		Marketing			
# Item Code I NDC:52124-0010-1 Marketing Information Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet Product Information	2 in 1 PAC	KAGE		Marketing			
# Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet Product Information Item Code (Source)	2 in 1 PAC	on Number or Monogra		Marketing			
# Item Code 1 NDC:52124-0010-1 Marketing Infor Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	2 in 1 PAC	NDC:52124-0011 ORAL		Marketing			
# Item Code 1 NDC:52124-0010-1 Marketing Information Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet Product Information Item Code (Source)	2 in 1 PAC	NDC:52124-0011 ORAL		Marketing \$ 0 4/24/20 10	Start Date	Marketin	ng End Date
# Item Code 1 NDC:52124-0010-1 Marketing Infor Marketing Category OTC monograph final Part 8 of 8 ASPIRIN aspirin tablet Product Information Item Code (Source) Route of Administration	2 in 1 PAC	NDC:52124-0011 ORAL ety dient Name		Marketing \$ 0 4/24/20 10	Start Date	Marketin	End Date

Inactive Ingredien	ts					
Ingredient Name						Strength
HYPROMELLOSE (UNI						
POLYETHYLENE GLY	C OL (UNII: 3WJQ0SDW1A)					
STARCH, CORN (UNII:	O8232NY3SJ)					
Product Character	victics					
Color	white (White)	Scor	0		no score	
Shape	ROUND	Size	C		11mm	
Flavor			int Code		44;157;ASP	IRIN
Contains		Inp I				
Contains						
Packaging						
# Item Code	Package Descriptio	n	Marketin	g Start Date	Ma	arketing End Date
1 NDC:52124-0011-1	2 in 1 PACKAGE					
Marketing Info	rmation					
Marketing Category	Application Number or M	lonogra	ph Citation	Marketing S	tart Date	Marketing End Date
OTC monograph final	part343			04/24/2010		
Marketing Info	rmation					
Marketing Category	Application Number or M	lonogra	ph Citation	Marketing S	tart Date	Marketing End Date
OTC monograph final	part333			04/24/2010		

Labeler - Genuine First Aid LLC (619609857)

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

Revised: 6/2010

Genuine First Aid LLC