# 50 PERSON ANSI - benzalkonium chloride, lidocaine, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, water, benzocaine, alcohol, ibuprofen, acetaminophen, aspirin, isopropyl alcohol

## 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.

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#### 50 Person 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

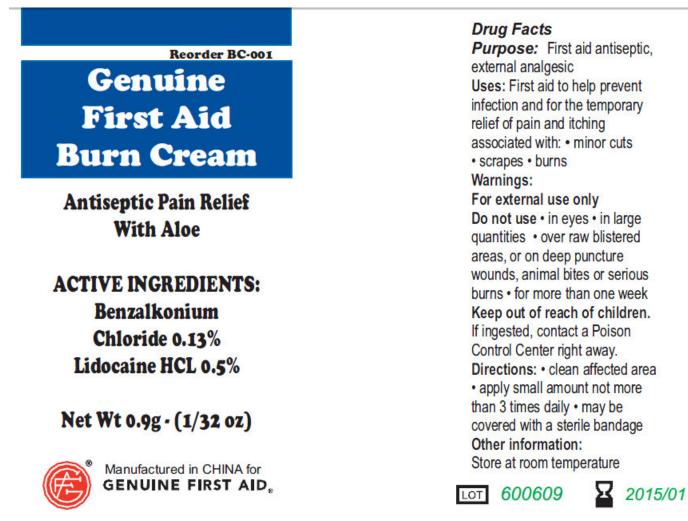
Genuine First Aid Burn Cream

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

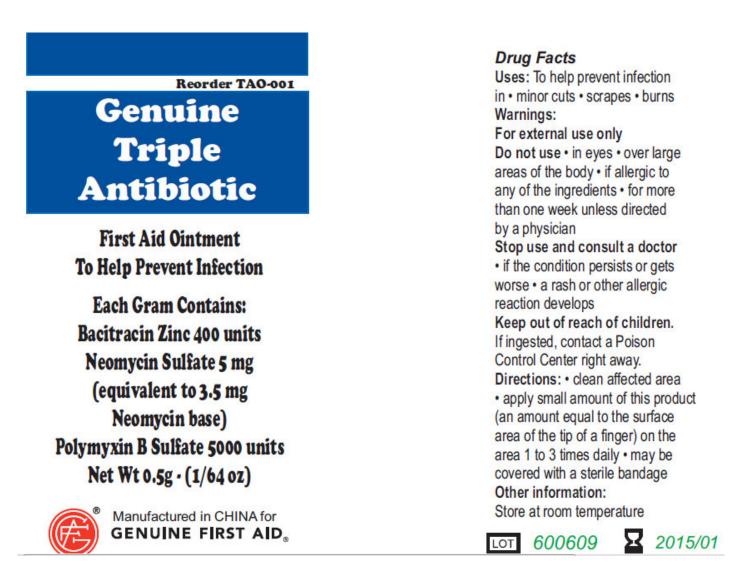
Neomycin base)

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	· '

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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 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.

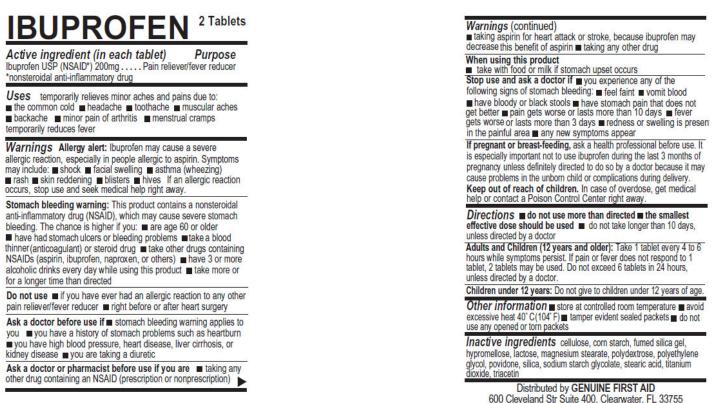
Children under 12 years: Do not give to children under 12 years of age.

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)
<b>NON–ASPIRIN</b> 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.
<ul> <li>muscular aches ■ backache ■ arthritis</li> <li>menstrual cramps ■ and reduction of fever</li> </ul>	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.
<b>Do not use I</b> 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.

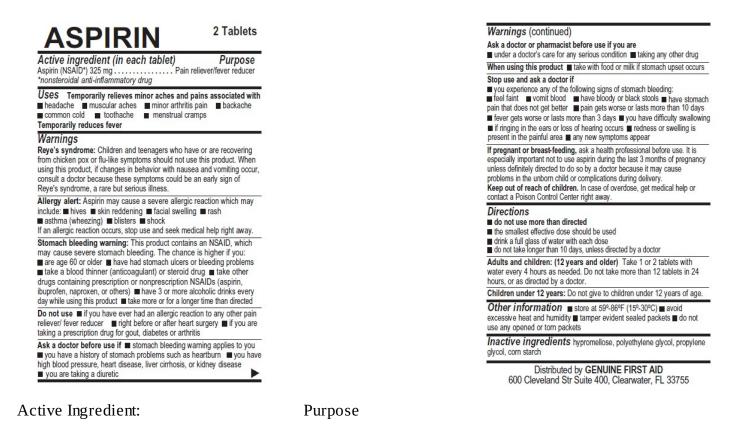
Store at 59 - 86 degree Fahrenheit (15 - 30 degree Celsius); avoid

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



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



#### DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient:	Purpose:
Isopropyl Alcohol, 70% v/v	Antiseptic
Use: For preparation of the skin be	efore injection.
Warnings: For external use only	. 1
Flammable - keep away from fire	or flame.
Do not use: with electrocautery, in	the eyes Ш
Stop use if irritation and redness of	levelop. 🗠
If condition persists for more than a	n the eyes III levelop. L 72 hours, III III
consult your doctor.	T
Keep out of reach of children. If get medical help or contact a Poise Center right away.	swallowed, r
Directions: Wipe injection site vige discard.	orously and
Other information: Store at room 15°-30° C (59°-86° F)	temperature
Inactive ingredient: Purified wate	r. i

LOT/EXP: Made in CHINA 20140301

Active Ingredient:

Ethyl Alcohol 62 percent

Enter section text here

Warnings:

For external use only

Flammable, keep away from fire or flame

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and consult a doctor if irritation and redness develop

and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Directions: Empty contents into palm. Rub hands until gel dissipates. Recommended for repeated use. Other Information: Store at 15 to 25 degree Celsius (59 to 77 degree Fahrenheit) Genuine Hand Sanitizer

Instant Gel

Antiseptic Handwash

with Vitamin E and Aloe

Kills 99.9 percent of germs

Without Water

Active Ingredient: Ethyl Alcohol 62 percent

Net Wt 0.9g (1/32 oz)

Manufactured in China For

Genuine First Aid.



Instant Gel Antiseptic Handwash with Vitamin E & Aloe

Kills 99.9% of Germs
Without Water
<b>Active Ingredient:</b>
Ethyl alcohol 62%
Net Wt 0.9g · (1/32 oz)

Manufactured in CHINA for GENUINE FIRST AID<sub>®</sub>

ANSI/ISEA Z308.1-2009 TYPE III

**Drug Facts** Uses: For hand washing to decrease bacteria on the skin Warnings: For external use only Flammable, keep away from fire or flame Do not use in the eyes. In case of contact, rinse eyes thoroughly with water Stop use and consult a doctor if irritation and redness develop and persist for more than 72 hours Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Directions: Empty contents into palm. Rub hands until gel dissipates. Recommended for repeated use Other information: Store at 15° to 25° C (59° to 77° F) LOT 600600 0 2015/01 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

POCKET 1: SMALL CUTS AND BURNS 100 Adhesive Plastic Bandages 1"x3" 2 Knuckle Fabric Bandage 25 Adhesive Spot Bandages 7/8"X7/8" 2 Fingertip Fabric Bandage 2 Elbow Knee Adhesive Bandage 6 Burn Cream 15 Antiseptic Towelettes 6 Triple Antibiotic Ointment 0.5gr 40 Cotton Tipped Applicators POCKET 2: MEDIUM CUTS AND SCRATCHES 15 Antiseptic Towelettes 4 Sterile Gauze Pad 3"x3" 4 Sterile Gauze Pads 4"X4" 2 Roller Gauze Bandage 2"X4.1yds 1 Roller Gauze Bandage 3"X4.1yds 2 Sterile Eve Pads 2 Sterile Eye Wash 10ml Twist Top 1 Butterfly Wound Closure, Medium POCKET 3: SEVERE BLEEDING AND BURNS 2 First Aid Tape Rolls 1/2"x5 yds. 2 Combine Pad 5"X9" POCKET 4: CPR 1 CPR Breathing Barrier POCKET 5: PROTECTION 4 Medical Grade Vinyl Gloves 1 Instant Cold Compress 30 Alcohol Cleansing Pads 1 Hand Sanitizer 0.9g 2 Insect Sting Relief Pads

2 Triangular

Bandage 42"x42"x59"

2 Ibuprofen 200mg 2 Non Aspirin 325mg 4 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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## **50 PERSON ANSI**

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

	uct T yp e	HUMAN C	OTC DRUG	Item Code (Source)		NDC:52124-	0113
Pack	aging						
#	Item Code	Pack	age Description	Marketing Start I	Date	Marketing	End Date
1 NDC	2:52124-0113-1	1 in 1 KIT					
Quar	ntity of Parts						
Part #	ŧ Pa	ackage Qua	antity	Tota	l Produc	t Quantity	
Part 1	6 PACKAGE			5.4 g			
Part 2	30 PACKAGE			24 mL			
Part 3	6 TUBE			3 g			
Part 4	2 BOTTLE			20 mL			
Part 5	2 PACKAGE			1 mL			
Part 6				2			
	1 PACKAGE			2			
	2 PACKAGE			4			
Part 9 Part	30 PACKAGE			15 mL			
10	1 PACKAGE			0.9 g			
GEI				PTIC PAIN REL	IEF WI	ITH ALOE	2
GEI				PTIC PAIN REL	IEF WI	ITH ALOF	2
<b>GEI</b> benza	NUINE FIRS	, lidocaine		PTIC PAIN REL	IEF WI	ITH ALOE	2
GEN benza Prod	NUINE FIRS alkonium chloride luct Information	, lidocaine		PTIC PAIN REL	IEF WI	TH ALOE	
GEN benza Prod Item (	NUINE FIRS	n lidocaine n	cream	PTIC PAIN REL	IEF WI	ITH ALOE	
GEN benza Prod Item (	NUINE FIRS alkonium chloride luct Information Code (Source)	n lidocaine n	cream NDC:52124-0004	PTIC PAIN REL	IEF WI	TH ALOE	
GEN benza Prod Item ( Route	NUINE FIRS alkonium chloride luct Information Code (Source)	n n active Moi	Cream NDC:52124-0004 TOPICAL	PTIC PAIN REL			
GEN benza Prod Item ( Route	NUINE FIRS alkonium chloride luct Information Code (Source) of Administratio	n n active Moi	Cream NDC:52124-0004 TOPICAL	PTIC PAIN REL		<b>Of Strength</b>	
GEN benza Prod Item ( Route Activ	NUINE FIRS alkonium chloride luct Information Code (Source) of Administration re Ingredient/A	n n Active Moi Ing	Cream NDC:52124-0004 TOPICAL			of Strength KONIUM	
GEN benza Prod Item ( Route Activ BENZ	NUINE FIRS alkonium chloride luct Information Code (Source) of Administration re Ingredient/A ALKONIUM CHLO N6JUD5X6Y)	n n Active Moi Ing RIDE (UNII: 1	NDC:52124-0004 TOPICAL	ALKONIUM -	Basis	of Strength KONIUM E	Strength 0.13 g in 100 g
GEN benza Prod Item ( Route Activ BENZ, UNII:71 LIDO (	NUINE FIRS alkonium chloride luct Information Code (Source) of Administration re Ingredient/A ALKONIUM CHLO N6 JUD5X6 Y) CAINE (UNII: 98 PI20	n n Active Moi Ing RIDE (UNII: 1	Cream NDC:52124-0004 TOPICAL ety redient Name F5UM2KM3W7) (BENZA	ALKONIUM -	Basis BENZALK CHLORID	of Strength KONIUM E	Strength 0.13 g
GEN benza Prod Item ( Route Activ BENZ UNII:71 LIDO ( Pack	NUINE FIRS alkonium chloride luct Information Code (Source) of Administration re Ingredient/A ALKONIUM CHLO N6 JUD5X6 Y) CAINE (UNII: 98 PI2C aging	n n Active Moi Ingı RIDE (UNII: 1 00987) (LIDC	Cream NDC:52124-0004 TOPICAL ety redient Name F5UM2KM3W7) (BENZA OCAINE - UNII:98PI2003	ALKONIUM - 987)	Basis BENZALK CHLORID LIDOCAIN	of Strength KONIUM E NE	Strength           0.13 g           in 100 g           0.5 g in 100
GEN benza Prod Item ( Route Activ BENZ UNII:71 LIDO ( Pack #	NUINE FIRS alkonium chloride luct Information Code (Source) of Administration re Ingredient/A ALKONIUM CHLO N6 JUD5X6 Y) CAINE (UNII: 98 PI20	n Active Moi Ingi RIDE (UNII: I 00987) (LIDC Pacl	Cream NDC:52124-0004 TOPICAL ety redient Name F5UM2KM3W7) (BENZA	ALKONIUM -	Basis BENZALK CHLORID LIDOCAIN	of Strength KONIUM E	Strengt 0.13 g in 100 g 0.5 g in 100

	rmation					
Marketing Category	Applicatio	on Number or Monogra	aph Citation	Marketing	Start Date	Marketing End Date
OTC monograph final	part345			06/02/2010		
Part 2 of 10						
ANTISEPTIC 7	<b>FOWELE</b>	TTE				
benzalkonium chlorid	e liquid					
Product Information	on					
Item Code (Source)		NDC:52124-0001				
Route of Administration	on	TOPICAL				
Active Ingredient//	Active Moi	ety				
		dient Name		Bas	is of Streng	gth Strength
BENZALKONIUM CHLO UNII:7N6 JUD5X6 Y)	ORIDE (UNII: F	F5UM2KM3W7) (BENZAL	KONIUM -	BENZA CHLOR	LKONIUM IDE	0.40 mL in 100 mL
Institut Inguadian	<b>t</b> o					
Inactive Ingredien		ngredient Name				Strength
WATER (UNII: 059QF0K						Strength
	,					
Packaging						
Packaging # Item Code	Pack	age Description	Marketin	g Start Date	Ma	arketing End Date
0 0		<b>xage Description</b> 1 PACKAGE	Marketin	g Start Date	Mi	arketing End Date
# Item Code			Marketin	g Start Date	Ma	arketing End Date
#         Item Code           1         NDC:52124-0001-1	0.8 mL in		Marketin	ig Start Date	Ma	arketing End Date
# Item Code 1 NDC:52124-0001-1 Marketing Info	0.8 mL in rmation	1 PACKAGE		-		
# Item Code 1 NDC:52124-0001-1 Marketing Info	0.8 mL in rmation			g Start Date Marketing S		arketing End Date Marketing End Date
# Item Code 1 NDC:52124-0001-1 Marketing Info	0.8 mL in rmation Applicatio	1 PACKAGE		Marketing		
# Item Code 1 NDC:52124-0001-1 Marketing Info	0.8 mL in rmation Applicatio	1 PACKAGE		Marketing		
# Item Code 1 NDC:52124-0001-1 Marketing Info	0.8 mL in rmation Applicatio	1 PACKAGE		Marketing		
#       Item Code         1       NDC:52124-0001-1         Marketing Info       Marketing Category         OTC monograph final	0.8 mL in rmation Applicatio part333	1 PACKAGE		Marketing		
#       Item Code         1       NDC:52124-0001-1         Marketing Info       Marketing Category         OTC monograph final         Part 3 of 10	0.8 mL in rmation Applicatio part333	1 PACKAGE	aph Citation	Marketing		
#       Item Code         1       NDC:52124-0001-1         Marketing Info       Marketing Category         OTC monograph final         Part 3 of 10         GENUINE TRIE	0.8 mL in rmation Applicatio part333	1 PACKAGE	aph Citation	Marketing		
#       Item Code         1       NDC:52124-0001-1         Marketing Info       Marketing Category         OTC monograph final         Part 3 of 10         GENUINE TRI         bacitracin zinc, neomy	0.8 mL in rmation Applicatio part333 PLE ANT ycin sulfate,p	1 PACKAGE	aph Citation	Marketing		
#       Item Code         1       NDC:52124-0001-1         Marketing Info       Marketing Category         OTC monograph final         Part 3 of 10         GENUINE TRIE	0.8 mL in rmation Applicatio part333 PLE ANT ycin sulfate,p	1 PACKAGE	aph Citation	Marketing		

	Active Mo	iety					
	Ing	redient Name			Basis of S	trength	Strength
BACITRACIN ZINC (UN	CITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO52I) BACITRACIN ZINC					ZINC	400 [iU] in 1 g
NEOMYCIN SULFATE (	UNII: 057Y62	6693) (NEOMYCIN - UNII:	I16QD7X297)	]	NEOMYCIN SU	5 mg in 1 g	
POLYMYXIN B SULFA	<b>TE</b> (UNII: 193)	71312D4) (POLYMYXIN B -	UNII:J2VZ07J9	96K)	POLYMYXIN B	5000 [iU] in 1	
Packaging							
# Item Code	Pac	kage Description	Marketin	ıg Start	Date	Marketi	ng End Date
1 NDC:52124-0003-1	0.5 g in 1	I TUBE					
Marketing Info	rmation						
Marketing Category	Applicati	ion Number or Monogra	aph Citation	Marke	ting Start Da	ate Marl	keting End Date
OTC monograph final	part333			06/02/20	0 10		
Part 4 of 10							
Product Information	on						
	on	NDC:52124-0005					
		NDC:52124-0005 OPHTHALMIC					
Item Code (Source)							
Item Code (Source) Route of Administrati	on	OPHTHALMIC					
Item Code (Source) Route of Administrati	on Active Mo	OPHTHALMIC		Basis	of Strength		Strength
Item Code (Source) Route of Administrati Active Ingredient/A	on Active Mo Ingredi	OPHTHALMIC iety ent Name		<b>Basis</b> WATER	of Strength		<b>Strength</b> nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K	on Active Mo Ingredi (OOR) (WATE	OPHTHALMIC iety ent Name			of Strength		_
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K	on Active Mo Ingredi (OOR) (WATE	OPHTHALMIC ie ty ent Name R - UNII:059QF0KO0R)			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien	on Active Mo Ingredi COOR) (WATE ts	OPHTHALMIC  ie ty  r VNII:059QF0KO0R)  Ingredient Name			of Strength		_
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien	on Active Mo Ingredi COOR) (WATE ts	OPHTHALMIC ie ty ent Name R - UNII:059QF0KO0R) Ingredient Nam X8 X)			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien SO DIUM CHLORIDE (U	on Active Mo Ingredi COOR) (WATE ts INII: 451W47IC MONOBASI	OPHTHALMIC  ie ty ent Name R - UNII:059QF0K00R)  Ingredient Nam (8 X) C (UNII: 3980JIH2SW)			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien SODIUM CHLORIDE (U SODIUM PHO SPHATE, SODIUM PHO SPHATE,	on Active Mo Ingredi COOR) (WATE ts INII: 451W47IC MONOBASIC DIBASIC (UN	OPHTHALMIC  ie ty ent Name R - UNII:059QF0KO0R)  Ingredient Nam (8 X) C (UNII: 3980JIH2SW) III: GR686LBA74)			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien SODIUM CHLORIDE (U SODIUM PHO SPHATE, SODIUM PHO SPHATE, EDETATE DISODIUM (U	on Active Mo Ingredi COOR) (WATE ts NIII: 451W47IC MONOBASIC DIBASIC (UN JNII: 7FLD91C	OPHTHALMIC  ie ↓  ie ↓  import Name  R - UNII:059QF0K00R)  Ingredient Nam  (8 ×)  ( UNII: 3980JIH2SW)  II: GR686LBA74)  S8 5()			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien SODIUM CHLORIDE (U SODIUM PHO SPHATE, SODIUM PHO SPHATE, EDETATE DISODIUM (U	on Active Mo Ingredi COOR) (WATE ts NIII: 451W47IC MONOBASIC DIBASIC (UN JNII: 7FLD91C	OPHTHALMIC  ie ↓  ie ↓  import Name  R - UNII:059QF0K00R)  Ingredient Nam  (8 ×)  ( UNII: 3980JIH2SW)  II: GR686LBA74)  S8 5()			of Strength		nL in 100 mL
Item Code (Source) Route of Administrati Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredien SODIUM CHLORIDE (U SODIUM PHO SPHATE, SODIUM PHO SPHATE, EDETATE DISODIUM (U	on Active Mo Ingredi COOR) (WATE ts NIII: 451W47IC MONOBASIC DIBASIC (UN JNII: 7FLD91C	OPHTHALMIC  ie ↓  ie ↓  import Name  R - UNII:059QF0K00R)  Ingredient Nam  (8 ×)  ( UNII: 3980JIH2SW)  II: GR686LBA74)  S8 5()			of Strength		nL in 100 mL
Product Information Item Code (Source) Route of Administration Active Ingredient/A WATER (UNII: 059QF0K Inactive Ingredient SODIUM CHLORIDE (U SODIUM CHLORIDE (U SODIUM PHO SPHATE, EDETATE DISODIUM (U BENZALKONIUM CHLO	on Active Mo Ingredi COOR) (WATE ts NIII: 451W47IC MONOBASIC DIBASIC (UN JNII: 7FLD91C	OPHTHALMIC  ie ↓  ie ↓  import Name  R - UNII:059QF0K00R)  Ingredient Nam  (8 ×)  ( UNII: 3980JIH2SW)  II: GR686LBA74)  S8 5()			of Strength		nL in 100 mL

				6 P		
# Item Code		kage Description	Marketin	ıg Start Date	Marl	keting End Date
1 NDC:52124-0005-1	10 mL in 1	1 BOTTLE				
Marketing Info						
Marketing Category		on Number or Monogra	aph Citation	Marketing Start	t Date N	Aarketing End Date
OTC monograph final	part349			06/02/2010		
Part 5 of 10						
INSECT STINC	G RELIEI	F PAD				
benzocaine,alcohol li						
Product Information	on					
Item Code (Source)		NDC:52124-0008				
Route of Administrati	on	TOPICAL				
Active Ingredient/	Activo Moi	otv				
Active ingredient/		dient Name		Dasis of S	twongth	Strongth
BENZOCAINE (UNII) US	0	ENZOCAINE - UNII:U3RS	V/8 IW5)	Basis of S BENZOCAIN	-	Strength 6 mL in 100 mL
ALCOHOL (UNII: 3K995				ALCOHOL	L	60 mL in 100 mL
X			,			
Packaging						
# Item Code	Pac	kage Description	Marketin	ıg Start Date	Marl	keting End Date
1 NDC:52124-0008-1		1 PACKAGE		0		5
Marketing Info	rmation					
Marketing Category		on Number or Monogra	aph Citation	Marketing Start	t Date N	Aarketing End Date
OTC monograph final	part348	5		06/02/2010		5
0 1	1					
Part 6 of 10						
IBUPROFEN						
ibuprofen tablet						
Product Information	on					
Item Code (Source)		NDC:52124-0009				
Item Code (Source) Route of Administrati		NDC:52124-0009 ORAL				

Active Ingredient/	Active Moiety				
	Ingredient Name		Basis	of Strength	Strength
IBUPROFEN (UNII: WKZ	2XYI10QM) (IBUPROFEN - UNII:WK2XY	IBUPROFEN		200 mg	
Inactive Ingredier	its				
	Ingredient Nam	ie		S	trength
PO WDERED CELLULO	SE (UNII: SMD1X3XO9M)				
STARCH, CORN (UNII:	O8232NY3SJ)				
HYPROMELLOSE (UN	I: 3NXW29V3WO)				
LACTOSE (UNII: J2B2A	4N98G)				
MAGNESIUM STEARA	<b>FE</b> (UNII: 70097M6I30)				
POLYDEXTROSE (UNI	I: VH2XOU12IE)				
POLYETHYLENE GLY	COL (UNII: 3WJQ0SDW1A)				
PO VIDO NE (UNII: FZ98	9 GH9 4E)				
SILICON DIOXIDE (UN	II: ETJ7Z6XBU4)				
STEARIC ACID (UNII: 4	ELV7Z65AP)				
TITANIUM DIO XIDE (U					
TRIACETIN (UNII: XHX)	3C3X673)				
Product Character					
Color	white (White)	Score		no score	
Shape	ROUND	Size		10 mm	
Flavor		Imprint Cod	e	44;352	
Contains					
Packaging					
# Item Code	Package Description	Marketin	ng Start Date	Marketing	End Date
<b>1</b> NDC:52124-0009-1	2 in 1 PACKAGE				
Marketing Info	rmation				
Marketing Category	Application Number or Monog	raph Citation	Marketing Start	Date Market	ting End Dat
	ANDA075010		06/02/2010		
ANDA					
ANDA					
ANDA					
Part 7 of 10 NON-ASPIRIN					
Part 7 of 10					
Part 7 of 10 NON-ASPIRIN					

Item Code (Source)		NDC:52124-0010					
Route of Administration	l	ORAL					
Active Ingredient/Ad	ctive Moie	ety					
	In	gredient Name			Basis of S	trength	Strength
ACETAMINO PHEN (UNII:	362O9ITL9E	) (ACETAMINOPHEN - U	NII:362O9ITL9	D)	ACETAMINO	NOPHEN 325 mg	
Inactive Ingradiants							
Inactive Ingredients		Ingredient Name				St	rength
STARCH, CORN (UNII: O8	232NY3SJ)	Ingreutent Name				ottengen	
POLYETHYLENE GLYCO		JQ0SDW1A)					
STEARIC ACID (UNII: 4EL)		- ,					
POVIDONE (UNII: FZ989G	H94E)						
Product Characteris							
Color	white (WHI	ГЕ)	Score			no score	
Shape Flavor	ROUND		Size	da		11mm AZ;234	
Contains			Imprint Co	ue		AL,234	
Contains							
Packaging							
# Item Code	Pack	age Description	Marketin	g Start Date	Ma	rketing <b>E</b>	End Date
<b>1</b> NDC:52124-0010-1							
Marketing Inform	liation						
Marketing Inform Marketing Category		on Number or Monogra	ph Citation	Marketing S	Start Date	Marketir	ng End Date
Marketing Category		n Number or Monogra	ph Citation	<b>Marketing S</b> 06/02/2010	Start Date	Marketir	ng End Date
Marketing Category	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketir	ng End Date
Marketing Category	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketir	ng End Date
Marketing Category	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketir	ng End Date
Marketing CategoryOTC monograph finalpPart 8 of 10	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketin	ng End Date
Marketing Category OTC monograph final p Part 8 of 10 ASPIRIN	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketin	ng End Date
Marketing CategoryOTC monograph finalpPart 8 of 10	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketin	ng End Date
Marketing Category OTC monograph final p Part 8 of 10 ASPIRIN	Applicatio	n Number or Monogra	ph Citation	_	Start Date	Marketin	ng End Date
Marketing Category       P         OTC monograph final       P         Part 8 of 10       ASPIRIN         aspirin tablet       P	<b>Applicatio</b> art343	n Number or Monogra	ph Citation	_	Start Date	Marketin	ng End Date
Marketing CategoryOTC monograph finalPart 8 of 10ASPIRIN aspirin tabletProduct Information	<b>Applicatio</b> art343		ph Citation	_	Start Date	Marketin	ng End Date
Marketing CategoryOTC monograph finalPart 8 of 10ASPIRIN aspirin tabletProduct Information Item Code (Source)	Applicatio art343	NDC:52124-0011	ph Citation	_	Start Date	Marketin	ng End Date
Marketing Category OTC monograph final p Part 8 of 10 ASPIRIN aspirin tablet Product Information	Applicatio art343		ph Citation	_	Start Date	Marketin	ng End Date
Marketing CategoryOTC monograph finalPart 8 of 10ASPIRIN aspirin tabletProduct Information Item Code (Source)	Applicatio art343	NDC:52124-0011	ph Citation	_	Start Date	Marketin	ng End Date
Marketing CategoryOTC monograph finalPart 8 of 10ASPIRIN aspirin tabletProduct Information Item Code (Source)	Applicatio art343	NDC:52124-0011 ORAL	ph Citation	_	Start Date	Marketin	ng End Date

Ingredient Name				Basis o	f Strength	Strength	
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)			ASPIRIN		325 mg		
Inactive Ingredient	ts						
		Ingredient Na	ime			Strength	
HYPROMELLOSE (UNII							
POLYETHYLENE GLYC		JQ0SDW1A)					
STARCH, CORN (UNII: C	)8232NY3SJ)						
Product Character	istics						
Color	white (White)	S	core	no	score		
Shape	ROUND		ize	11m	ım		
Flavor		In	nprint Code	44;	157;ASPIRIN		
Contains			•				
Packaging							
# Item Code	Pack	age Description	Marketin	g Start Date	Marketin	ıg End Date	
<b>1</b> NDC:52124-0011-1	2 in 1 PAC	KAGE					
Marketing Info	mation						
Marketing Category		on Number or Mono	ograph Citation	Marketing Start	Date Mark	eting End Date	
OTC monograph final				06/02/2010			
0 1	L						
Part 9 of 10							
ALCOHOL CLE	EANSING	G PAD					
isopropyl alcohol liqui	id						
<b>Product Information</b>	n						
Item Code (Source)		NDC:52124-0002					
Route of Administration	on	TOPICAL					
Active Ingredient/A	Active Moi	ety					
0		redient Name		Basis	of Strength	Strength	
ISOPROPYL ALCOHOI	0		ALCOHOL -	ISOPRO	_	70 mL	
UNII:ND2M416302)				ALCOH	OL	in 100 mL	
<b>Inactive Ingredient</b>	ts						

Ingredient Name						Strength		
WATER (UNII: 059QF0KO0R)								
Packaging								
# Item Code	-			ate	Marketing End Date			
<b>1</b> NDC:52124-0002-1	$0.5 \ mL$ in	1 PACKAGE						
<b>Marketing Inform</b>	nation							
Marketing Category				Marketii	ng Start	Date	Mark	eting End Date
OTC monograph final pa	art333			06/02/2010	)			
Part 10 of 10								
GENUINE HAND SANITIZER INSTANT ANTISEPTIC HANDWASH WITH VITAMIN E AND ALOE alcohol gel								
<b>Product Information</b>	I							
Item Code (Source)		NDC:52124-0006						
Route of Administration TOPICAL								
Active Ingredient/Ac	ctive Moie	ety						
					Basis of	Streng	th	Strength
ALCOHOL (UNII: 3K9958	V90M) (ALC	OHOL - UNII:3K9958V9	0 M)	ALCOHOL 62 g in			62 g in 100 g	
Packaging	-							
	Item Code Package Description Marketing Start Da			ate	Marketing End Date			
1 NDC:52124-0006-1	0.9 g in 1	PACKAGE						
<b>Marketing Inform</b>	nation							
Marketing Category				Marketii	farketing Start Date Marketing End			eting End Date
OTC monograph final pa	art333							
Marketing Inform	nation							
Marketing Category	Applicatio	n Number or Monogra	aph Citation	Marketing Start Date M			Marketing End Date	
OTC monograph final pa	art333			06/02/2010	)			

Labeler -	Genuine	First A	Aid LLC	(619609857)
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## Establishment

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

Revised: 6/2010

Genuine First Aid LLC