ACME FIRST AID CONTAINS 404 PIECES - benzalkonium chloride, lidocaine, water, isopropyl alcohol, aspirin, benzocaine, alcohol, bacitracin zinc, neomycin sulfate, polymyxin b sulfate

Acme United Corp.

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.

ACME FIRST AID Contains 404 PIECES

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

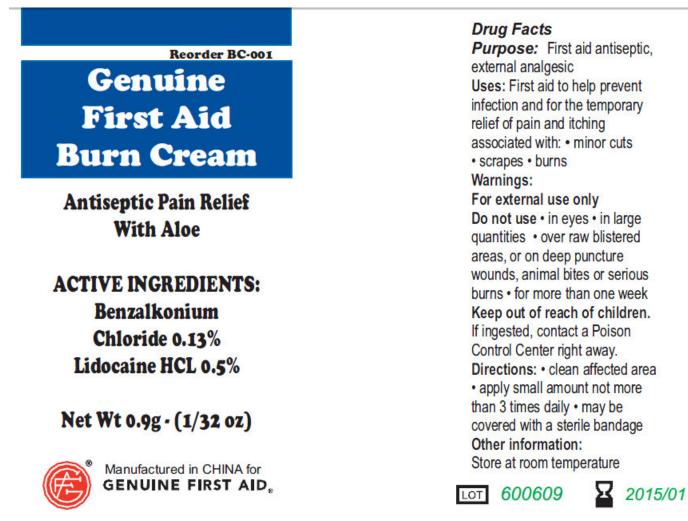
Genuine First Aid Burn Cream

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

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;



DRUG FACTS - Antiseptic Towelette

	VIVIHO OI OPOM	
•		towelette appropriately after sir Inactive ingredient: Purified w
۰.		and use to cleanse desired skin
TE		Directions: Tear open packet,
AR		Do not use: In the eyes, or ove
I		If unusual redness, swelling or
Π	nter right away.	or contact a Poison Control Ce
R	, get medical help	reach of children. If swallowed
Π	ly. Keep out of	Warnings: For external use on
1	nd water.	hands and body without soap a
	,eansing of face,	prevent infection. Antiseptic cle
		Use: For Professional and Hos
		Benzalkonium Chloride 0.40%.
1	Purpose:	Active Ingredient:

GENUINE FIRST AID

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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

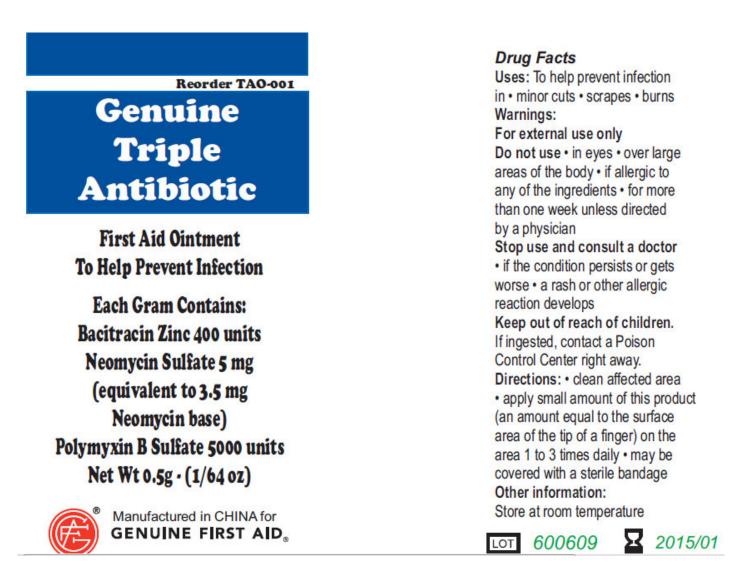
Neomycin base)

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



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



DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient:	Purpose:
Isopropyl Alcohol, 70% v	/v Antiseptic
Use: For preparation of	the skin before injection.
Warnings: For external	use only. I
Flammable - keep away	from fire or flame.
Do not use: with electro	
Stop use if irritation and	redness develop.
If condition persists for n consult your doctor.	hore than 72 hours,
Keep out of reach of cl	ildren If swallowed
get medical help or conta	
Center right away.	Ш
Directions: Wipe injection discard.	on site vigorously and
Other information: Stor	e at room temperature
15°-30° C (59°-86° F)	o at room tomporataro 1
Inactive ingredient: Pu	ified water.
LOT/EXP:	Made in CHINA

20140301

Active Ingredient:

Purpose:

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:

1/pouch

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

GENUINE FIRST AID Made in CHINA . т

physician immediately. initation or other symptoms occur, consult a - TEAR HERE puncture wounds. If unusual redness, swelling, Do not use: In eyes, on broken skin, deep thoroughly with water. contact with eyes. If this happens, rinse Flammable - keep away from fire or flame. Avoid or contact a Poison Control Center right away. reach of children. If swallowed, get medical help Warnings: For external use only. Keep out of Selid associated with minor burns, scrapes and insect Dee: For the temporary relief of pain and itching SD alcohol, 60% w/v Antiseptic Benzocaine, 6% w/v Topical Anesthetic ١. Purpose: Active Ingredient: DRUG FACTS - Insect Sting Relief Pad

REORDER ISRP-001 Insect Sting **Relief Pad Toallitas para** Picaduras de Insectos Genuine First Aid LLC, Clearwater FL 33755 www.GenuineFirstAid.com ISO 9001-2010 GENUINE FIRST AID.

Active ingredient (in each tablet)

*nonsteroidal anti-inflammatory drug

Uses

Purpose

Aspirin 81 mg (NSAID*). Analgesic/Antipyretic

temporary relief of minor aches, pains and headaches

to reduce fever associated with colds, sore throats, teething

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of reye's syndrome, a rare but serious illness.

Stomach bleeding warning: This product contains a non-steroidal anti-inflammatory drug (NSAID), which may cause stomach bleeding. The chance is higher if the user; has had stomach ulcers or bleeding problems takes a blood thinning (anticoagulant) or steroid drug takes other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others) takes more or for a longer time than directed is age 60 or older has 3 or more alcoholic drinks every while using this product.

Do not use

if you are allergic to aspirin or any other pain relievers/fever reducers

for more than 10 days for pain unless directed by a doctor

for more than 3 days for fever unless directed by a doctor

for at least 7 days after a tonsillectomy or oral surgery

Ask a doctor before use if you have;

asthma

stomach problems (such as heartburn, upset stomach, or stomach pain) that persist or recur, ulcers or bleeding problems

a child experiencing arthritis pain

if you are;

allergic to aspirin

taking prescription drug for anticoagulation (thinning of blood), diabetes, gout or arthritis

Stop use and ask a doctor if;

an allergic reaction occurs, Seek medical help right away

pain or fever persists

new or unexpected symptoms occur

redness or swelling is present

ringing in ears or loss of hearing occurs

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

Directions Do not use more than directed

Adults and Children (12 years and older): Chew 4 to 8 tablets with water every 4 hours, Do not exceed 48 tablets in 24 hours unless directed by a doctor.

Children Under 12 years: Consult a doctor.

Store at room temperature

tamper evident sealed packets

do not use any opened or torn packets

Inactive ingredients flavor, FD and C yellow No. 6, saccharin, sodium, silicon dioxide, starch, stearic acid, sucrose

MADE IN USA

Distributed by GENUINE FIRST AID

600 Cleveland Str Suite 400, Clearwater, FL 33755

2 Tablets

GENUINE FIRST AID.

Chewable Aspirin 81mg

2 Tablets



Chewable Aspirin 81mg

Active ingredient (in each tablet) Purpose Aspirin 81 mg (NSAID*)Analgesic/Antipyretic *nonsteroidal anti~inflammatory drug

Uses

temporary relief of minor aches, pains and headaches
 to reduce fever associated with colds, sore throats, teething

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Stomach bleeding warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause stomach bleeding. The chance is higher if the user ■ has had stomach ulcers or bleeding problems ■ takes a blood thinning (anticoagulant) or steroid drug ■ takes other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others) ■ takes more or for a longer time than directed ■ is age 60 or older ■ has 3 or more alcoholic drinks every day while using this product

Do not use ■ if you are allergic to aspirin or any other pain relievers/fever reducers ■ for more than 10 days for pain unless directed by a doctor ■ for more than 3 days for fever unless directed by a doctor ■ for at least 7 days after a tonsillectomy or oral surgery

Warnings (continued)

Ask a doctor before use if you have ■ asthma ■ stomach problems (such as heartburn, upset stomach, or stomach pain) that persist or recur, uclers or bleeding problems ■ a child experiencing arthritis pain

if you are allergic to aspirin taking a prescription drug for anticoagulation (thinning of the blood), diabetes, gout or arthritis

Stop use and ask a doctor if ■ an allergic reaction occurs. Seek medical help right away ■ pain or fever persists ■ new or unexpected symptoms occur ■ redness or swelling is present ■ ringing in the ears or loss of hearing occurs

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

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

Directions Do not use more than directed Adults and Children (12 years and older): Chew 4 to 8 tablets with water every 4 hours. Do not exceed 48 tablets in 24 hours unless directed by a doctor.

Children Under 12 Years: Consult a doctor.

Other information

■ store at room temperature ■ tamper evident sealed packets ■ do not use any opened or torn packets

Inactive ingredients flavor, FD&C yellow #6, saccharin sodium, silicon dioxide, starch, stearic acid, sucrose

MADE IN USA Distributed by GENUINE FIRST AID 600 Cleveland Str Suite 400, Clearwater, FL 33755 FIRST AID all purpose kit with 6 First Aid Pockets

1 Small Cuts and Burns

2 Medium Cuts and Scratches

3 Severe Bleeding and Burns

4 CPR

5 Protection

6 Instruments

New REFILL PACKS AVAILABLE FOR EACH POCKET

HOME PLAY AUTO OFFICE WORK

CONTAINS 1000 PIECES

Soft-Sided Storage Case

With Pockets that are Organized for Quick and Easy Emergency Access

CARRYING CASE 1 Soft-sided Bag 1 Inside Plastic Pockets

SMALL CUTS AND BURNS Pocket One
95 Adhesive Plastic Bandages 3/4"x3"
95 Junior Plastic Bandages 3/8"x1-1/2"
45 Adhesive Plastic Bandages 1"x3"
4 Knuckle Fabric Bandages
56 Adhesive Spot Bandages 7/8"x7/8"
4 Fingertip Fabric Bandages
3 Elbow/Knee Adhesive Bandage
1 Burn Cream
9 Antiseptic Towelettes
2 Triple Antibiotic Ointment
5 Sterile Gauze Pads 2"x2"
20 Cotton Tipped Applicatiors

MEDIUM CUTS AND SCRATCHES Pocket Two

12 Antiseptic Towelettes
5 Sterile Gauze Pad 2"X2"
2 Sterile Gauze Pad 3"x3"
2 Sterile Gauze Pad 4"x4"
1 Roll Gauze Bandages 2"
1 Sterile Eye Pads
1 Sterile Eye Wash
5 Butterfly Wound Closures

SEVERE BLEEDING AND BURNS Pocket Three 1 First Aid Tape Roll 1/2" 1 Combine Pad 5"x9" 1 Combine Pad 8"x10" 1 Roll Gauze Bandage 3"

CPR Pocket Four 1 CPR Breathing Barrier

PROTECTION Pocket Five
2 Medical Grade Gloves
1 Instant Cold Compress
6 Alcohol Cleansing Pads
2 Insect Sting Relief Pads
1 Emergency Blanket 38"X 60"
4 Chewable Aspirin Tablets
1 Triangular Bandage 42"x42"x59"

INSTRUMENTS Pocket Six 1 Emergency First Aid Guide 1 Plastic Tweezers 1 Scissors 6 Assorted Safety Pins 2 Wooden Finger Splints

CAUTION: This kit contains products that have expiration dates. Please check before using.

Acme United Corporation 60 Round Hill Road Fairfield, CT 06824 www.acmeunited.com Designed in USA / Made in India 2009 Acme United Corporation PhysiciansCare and are Registered Trademrks of Acme United Corporation



ACME FIRST AID CONTAINS 404 PIECES

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

Product Information						
Produ	ict T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-1806		
Packa	aging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC	:0924-1806-01	1 in 1 KIT				
Quan	tity of Parts					
-		ackage Quantity	Total Produc	t Quantity		
Part #		ackage Quantity	Total Produc 0.9 g	t Quantity		
Part # Part 1	P	ackage Quantity		t Quantity		
Part # Part 1 Part 2	P 1 PACKAGE 21 PACKAGE	ackage Quantity	0.9 g	t Quantity		
Part # Part 1 Part 2 Part 3	P 1 PACKAGE 21 PACKAGE	ackage Quantity	0.9 g 16.8 mL	t Quantity		
Part # Part 1 Part 2 Part 3 Part 4	1 PACKAGE 21 PACKAGE 1 BOTTLE	ackage Quantity	0.9 g 16.8 mL 10 mL	t Quantity		

Part 1 of 7

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) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE	$0.5\;g$ in 100 g		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0004-1	0.9 g in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part345	04/27/2010	

Part 2 of 7

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Item Code (Source)	NDC:52124-0001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.40 mL			
UNII:7N6 JUD5X6 Y)	CHLORIDE	in 100 mL			

Inactive Ingredients					
			St	rength	
WATER (UNII: 059QF0KO0R	L)				
Packaging					
# Item Code	Package Description	Marketin	g Start Date	Marke	ting End Date
1 NDC:52124-0001-1	0.8 mL in 1 PACKAGE				
Marketing Inform	ation				
Marketing Category A	pplication Number or Monogra	ph Citation	Marketing Start D	ate Ma	rketing End Date
OTC monograph final part	333		04/27/2010		
Part 3 of 7					
STERILE ISOTOR	NIC BUFFERED GEN	UINE EYI	EWASH		
Product Information					
Item Code (Source)	NDC:52124-0005				
Route of Administration	OPHTHALMIC				
Active Ingredient/Acti	ive Moiety				
I	ngredient Name		Basis of Strength	1	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	V	WATER	98.16	5 mL in 100 mL
Inactive Ingredients					
Ingredient Name					Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)					
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)					
SO DIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)					
EDETATE DISO DIUM (UNII: 7FLD91C86K) BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)					
Packaging					
# Item Code	Package Description	Marketin	g Start Date	Marke	ting End Date
1 NDC:52124-0005-1	10 mL in 1 BOTTLE				

Marketing Infor	mation						
Marketing Category	Applicatio	on Number or Monogra	ph Citation	Marketin	ıg Start Date	Mark	eting End Date
OTC monograph final	part349			04/27/2010			
Part 4 of 7							
ALCOHOL CLE	ANSING	F PAD					
isopropyl alcohol liquid	1						
Product Information	n						
Item Code (Source)		NDC:52124-0002					
Route of Administratio	n	TOPICAL					
Active Ingredient/A	ctive Moi	a tw					
Active Ingretient/A		redient Name			Basis of Stre	ngth	Strength
ISOPROPYL ALCOHOL	0		COHOL-		ISOPROPYL		70 mL
UNII:ND2M416302)					ALCOHOL		in 100 mL
Inactive Ingredients	S						
		ngredient Name				Strei	ıgth
WATER (UNII: 059QF0KC	00R)						
Packaging							
# Item Code	Pack	age Description	Marketin	ıg Start Da	ite Ma	arketin	g End Date
1 NDC:52124-0002-1	0.5 mL in	1 PACKAGE					
Marketing Infor	mation						
Marketing Category		n Number or Monogra	ph Citation	Marketin	ıg Start Date	Mark	eting End Date
	part333			04/27/2010	-		0
Part 5 of 7							
	DELIPI						
INSECT STING benzocaine, alcohol liq		PAD					
centro came, arconor inq	ulu						
Product Information	n						

Item Code (Source)		NDC:52124-0008						
Route of Administratio	of Administration TOPICAL							
Active Ingredient/A	Active Mo	iety						
	Ingr	edient Name			Basis of S	Strengt	h	Strength
BENZOCAINE (UNII: U3R	RS Y48 JW5) (I	BENZOCAINE - UNII:U3R	S Y48 J W5)		BENZOCAIN	E	6 r	nL in 100 mL
ALCOHOL (UNII: 3K995	8V90M) (AL	COHOL - UNII:3K9958V9	90M)		ALCOHOL		60	mL in 100 mL
Packaging								
# Item Code	Рас	kage Description	Marketir	ıg Sta	rt Date	Ma	rketir	ng End Date
1 NDC:52124-0008-1		n 1 PACKAGE		18 010	Dutt	Marketing Lift Date		
Marketing Infor Marketing Category		on Number or Monogr	anh Citation	Мал	·keting Start	t Date	Mark	eting End Dat
	part348	on number or monogr			//2010	Date	WIGTK	eting End Dat
CHEWABLE AS								
CHEWABLE AS aspirin tablet, chewabl	le							
CHEWABLE AS aspirin tablet, chewabl Product Informatio	le	NDC 52124 0012						
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source)	le)n	NDC:52124-0012						
CHEWABLE AS aspirin tablet, chewabl Product Informatio	le)n	NDC:52124-0012 ORAL						
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio	n on	ORAL						
aspirin tablet, chewabl Product Informatio Item Code (Source)	le on Active Mo	ORAL			Basis o	fStren	gth	Strength
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio	le on Active Mo Ingro	ORAL ie ty edient Name			Basis o Aspirin	fStren	gth	Strength 81 mg
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y	on Active Mo Ingro 76E) (ASPIR)	ORAL ie ty edient Name				fStren	gth	
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y	on Active Mo Ingro 76E) (ASPIR)	ORAL ie ty edient Name				fStren		
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y Inactive Ingredient	on Active Mo Ingre 776E) (ASPIRI	ORAL ORAL IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII				fStren		81 mg
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y Inactive Ingredient	on Active Mo Ingro 76E) (ASPIRI S UNII: H77VEI	ORAL ORAL IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII				fStren		81 mg
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16C05Y Inactive Ingredient FD&C YELLOW NO.6 (SACCHARIN (UNII: FST46	e on on Active Mo Ingre (76E) (ASPIR) S S UNII: H77VEI 67XS7D)	ORAL ORAL IN - UNII:R16CO5Y76E) Ingredient Name 93A8)				f Stren;		81 mg
CHEWABLE AS aspirin tablet, chewable Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y Inactive Ingredient FD&C YELLOW NO. 6 (SACCHARIN (UNII: FST40 SO DIUM CATION (UNII: SILICON DIO XIDE (UNII	e on on Active Mo Ingre 76E) (ASPIRI S S UNII: H77VEI 67XS7D) LYR4M0 NH3 I: ETJ7Z6 XBU	ORAL ORAL Ingredient Name Ingr				f Stren;		81 mg
CHEWABLE AS aspirin tablet, chewabl Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y Inactive Ingredient FD&C YELLOW NO.6 (I SACCHARIN (UNII: FST40 SODIUM CATION (UNII: SILICON DIO XIDE (UNII STARCH, CORN (UNII: O	le on on Active Mo Ingro 76E) (ASPIRI 75E) (ASPIRI 75E) (ASPIRI 75E) 75E) 75E) 75E) 75E) 75E) 75E) 75E)	ORAL ORAL Ingredient Name Ingr				f Stren;		81 mg
CHEWABLE AS aspirin tablet, chewable Product Informatio Item Code (Source) Route of Administratio Active Ingredient/A ASPIRIN (UNII: R16CO5Y Inactive Ingredient FD&C YELLOW NO. 6 (SACCHARIN (UNII: FST40 SO DIUM CATION (UNII: SILICON DIO XIDE (UNII	e e on on Active Mo Ingre 76E) (ASPIRI 76E) (ASPIRI 58 UNII: H77VEI 67XS7D) LYR4M0 NH3 E ETJ7Z6 XBI 08232NY3SJ) LV7Z65AP)	ORAL ORAL Ingredient Name Ingr				f Stren;		81 mg

	ristics						
Color	orange (ORA	NGE)	Score			no score	
Shape	ROUND		Size			11mm	
Flavor			Imprint Code		ASF	ASPIRIN	
Contains							
Packaging							
# Item Code	Pack	age Description	Marketin	ig Start I	Date	Marketiı	ıg End Date
1 NDC:52124-0012-1	2 in 1 PAC	KAGE					
Marketing Info	rmation						
Marketing Category		on Number or Monogr	aph Citation	Marke	ting Start I	Date Mark	eting End Date
OTC monograph final	part345			04/27/20	10		
Part 7 of 7							
GENUINE TRI bacitracin zinc, neom			intment				
bacitracin zinc, neom	ycin sulfate,ŗ		intment				
bacitracin zinc,neom Product Information	ycin sulfate,ŗ		intment				
	ycin sulfate, _f on	oolymyxin b sulfate oi	intment				
bacitracin zinc,neom Product Information Item Code (Source)	ycin sulfate, _f on	oolymyxin b sulfate oi NDC:52124-0003	intment				
bacitracin zinc,neom Product Informati Item Code (Source) Route of Administrati	ycin sulfate, _F on	oolymyxin b sulfate of NDC:52124-0003 TOPICAL	intment				
bacitracin zinc, neom Product Informational Item Code (Source) Route of Administrational Active Ingredient/	ycin sulfate,p on Active Moi Ingr	oolymyxin b sulfate oi NDC:52124-0003 TOPICAL ety edient Name			Basis of a	Strength	Strength
bacitracin zinc, neom Product Information Item Code (Source) Route of Administration Active Ingredient/A BACITRACIN ZINC (UN	ycin sulfate,p on ion Active Moi Ingr III: 89 Y4M234E	oolymyxin b sulfate of NDC:52124-0003 TOPICAL ety edient Name .S) (BACITRACIN - UNII:	58 H6 RWO 521)		BACITRACIN	N ZINC	400 [iU] in 1 g
bacitracin zinc,neom Product Informational Item Code (Source) Route of Administrational Active Ingredient / BACITRACIN ZINC (UN NEOMYCIN SULFATE (ycin sulfate, on Active Moi Ingr III: 89 Y4M234E (UNII: 057 Y626	oolymyxin b sulfate oi NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII	58 H6 RWO 52I) :116 QD7 X29 7)	Ν	BACITRACIN NEOMYCIN S	N ZINC SULFATE	400 [iU] in 1 g 5 mg in 1 g
bacitracin zinc, neom Product Informational Item Code (Source) Route of Administrational Active Ingredient / BACITRACIN ZINC (UN NEOMYCIN SULFATE (ycin sulfate, on Active Moi Ingr III: 89 Y4M234E (UNII: 057 Y626	oolymyxin b sulfate oi NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII	58 H6 RWO 52I) :116 QD7 X29 7)	Ν	BACITRACIN NEOMYCIN S	N ZINC	400 [iU] in 1 g 5 mg in 1 g
bacitracin zinc, neom Product Information Item Code (Source) Route of Administration Active Ingredient / BACITRACIN ZINC (UNINEOMYCIN SULFATE (POLYMYXIN B SULFATE)	ycin sulfate, on Active Moi Ingr III: 89 Y4M234E (UNII: 057 Y626	oolymyxin b sulfate oi NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII	58 H6 RWO 52I) :116 QD7 X29 7)	Ν	BACITRACIN NEOMYCIN S	N ZINC SULFATE	400 [iU] in 1 g 5 mg in 1 g
bacitracin zinc, neomy Product Information Item Code (Source) Route of Administration Active Ingredient/A BACITRACIN ZINC (UN NEOMYCIN SULFATE (POLYMYXIN B SULFATE)	ycin sulfate, on Active Moi Ingra III: 89 Y4M234E (UNII: 057 Y626 TE (UNII: 1937	NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII 1312D4) (POLYMYXIN B	58 H6 RWO 52I) :116 QD7 X29 7)	06K) F	BACITRACIN NEOMYCIN S OLYMYXIN	I ZINC SULFATE B SULFATE	400 [iU] in 1 g
bacitracin zinc, neomy Product Information Item Code (Source) Route of Administration Active Ingredient/A BACITRACIN ZINC (UN NEOMYCIN SULFATE (POLYMYXIN B SULFATE) POLYMYXIN B SULFATE Packaging # Item Code	ycin sulfate, on on Active Moi Ingr III: 89 Y4M234E (UNII: 057Y626 TE (UNII: 1937	NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII 1312D4) (POLYMYXIN B	58 H6 RWO 521) :116 Q D7 X29 7) - UNII:J2 V Z 0 7 J9	06K) F	BACITRACIN NEOMYCIN S OLYMYXIN	I ZINC SULFATE B SULFATE	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1g
bacitracin zinc, neomy Item Code (Source) Route of Administration Active Ingredient/ BACITRACIN ZINC (UN NEOMYCIN SULFATE (POLYMYXIN B SULFAT Packaging # Item Code 1 NDC:52124-0003-1	ycin sulfate,p on Active Moi Ingra III: 89 Y4M234E (UNII: 89 Y4M234E (UNII: 1937 TE (UNII: 1937 Dack 0.5 g in 1	NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII 1312D4) (POLYMYXIN B	58 H6 RWO 521) :116 Q D7 X29 7) - UNII:J2 V Z 0 7 J9	06K) F	BACITRACIN NEOMYCIN S OLYMYXIN	I ZINC SULFATE B SULFATE	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1g
bacitracin zinc, neomy Product Information Item Code (Source) Route of Administration Active Ingredient/A BACITRACIN ZINC (UN NEOMYCIN SULFATE (POLYMYXIN B SULFATE) POLYMYXIN B SULFATE Packaging # Item Code	ycin sulfate, on Active Moi Ingr III: 89 Y4M234E (UNII: 057Y626 TE (UNII: 1937 Date 0.5 g in 1	NDC:52124-0003 TOPICAL ety edient Name (S) (BACITRACIN - UNII: 693) (NEOMYCIN - UNII 1312D4) (POLYMYXIN B	58 H6 RWO 521) :116 Q D7 X29 7) - UNII:J2 V Z 0 7 J 9 Marketin	96K) F	BACITRACIN NEOMYCIN S OLYMYXIN	I ZINC SULFATE B SULFATE Marketin	400 [iU] in 1g 5 mg in 1g 5000 [iU] in 1g

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333	04/27/2010				

Labeler - Acme United Corp. (001180207)

Establishment

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

Revised: 6/2010

Acme United Corp.