

CVS ALL PURPOSE FIRST AID KIT- aspirin, diphenhydramine hydrochloride, aspirin, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin b, benzalkonium chloride, ammonia, lidocaine, acetaminophen, ibuprofen, CVS

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.

CVS All Purpose First Aid Kit

Active Ingredients - Genuine Triple Antibiotic

Active Ingredient:Bacitracin Zinc 400 units

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

Polymyxin B Sulfate 5000 units

Purpose - Genuine Triple Antibiotic

Triple Antibiotic

Uses - Genuine Triple Antibiotic

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

Warnings - Genuine Triple Antibiotic

For external use only

DO NOT USE - Genuine Triple Antibiotic

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 - Genuine Triple Antibiotic

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 - Genuine Triple Antibiotic

Keep out of reach of children.

If ingested, contact a Poison

Control Center right away.

Directions - Genuine Triple Antibiotic

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

Storage and Handling - Genuine Triple Antibiotic

Other information:

Store at room temperature.

Inactive Ingredients - Genuine Triple Antibiotic

Vaseline

Mineral Oil

Purified Water

Active Ingredients - Antiseptic

Active Ingredient:

Benzalkonium Chloride 0.13

Purpose - Antiseptic

Antiseptic

Use - Antiseptic

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

Warnings - Antiseptic

Warning: For external use only.

Keep out of reach of children - Antiseptic

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

Stop Use - Antiseptic

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

Do Not Use - Antiseptic

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

Directions - Antiseptic

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

Inactive Ingredients - Antiseptic

Inactive Ingredient: Purified water

Active Ingredient - Ibuprofen

Ibuprofen USP (NSAID*) 200mg

*nonsteroidal anti-inflammatory drug

Purpose - Ibuprofen

Pain reliever/fever reducer

Uses - Ibuprofen

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 - Ibuprofen

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 - Ibuprofen

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 - Ibuprofen

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 - Ibuprofen

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

Stop Use - Ibuprofen

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; redness or swelling is present in the painful area; any new symptoms appear

Pregnancy or Breast Feeding - Ibuprofen

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 - Ibuprofen

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

Directions - Ibuprofen

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

Active Ingredients - Non-Aspirin

Acetaminophen 500 mg

Purpose - Non Aspirin

Analgesic/antipyretic

Uses - Non Aspirin

temporary relief of minor aches and pains associated with:

common cold; headache; toothache; muscular aches; backache; arthritis; menstrual cramps; and reduction of fever

Warnings - Non Aspirin

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 - Non Aspirin

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 - Non Aspirin

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 - Non Aspirin

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

Pregnancy - Non Aspirin

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

Keep Out of Reach of Children - Non Aspirin

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 - Non Aspirin

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 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. 2 years of age This will provide more than the recommended dose (overdose) and could cause serious health problems.

Storage and Handling - Non Aspirin

Store at 59-86 degree F (15-30 degree C)

tamper evident sealed packets; do not use any open or torn packets

Active Ingredients - Burn Cream

ACTIVE INGREDIENTS:

Lidocaine HCL 2.5%

Purpose - Burn Cream

Purpose: First aid antiseptic, external analgesic

Uses - Burn Cream

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

Minor Cuts

Scrapes

Burns

Warnings - Burn Cream

For external use only

Do Not Use - Burn Cream

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

Do not use:

in the eyes or apply over large areas of the body.

longer than 1 week unless directed by a doctor.

in large quantities, particularly over raw surfaces or blistered areas.

Ask a doctor before use if you have deep puncture wounds, animal bites or serious burns.

When using this product, avoid contact with the eyes.

Stop Use - Burn Cream

Stop use and ask a doctor if

condition worsens

symptoms persist for more than 7 days

condition clears up and occurs again within a few days

Keep Out of Reach of Children

Keep out of reach of Children.

If ingested, contact a Poison Control Center right away.

Directions - Burn Cream

Adults and children 2 years of age and older
clean affected area.

apply a small amount of this product on the area 1 to 3 times daily.

may be covered with a sterile bandage

children under 2 years of age: consult a doctor

Storage and Handling - Burn Cream

Other Information:

Store at room temperature (do not freeze).

Taper evident sealed packets.

Do not use packet if opened or torn.

Active Ingredients - After Bite

Active Ingredient:

Ammonia 3.5%

Purpose - After Bite

Counterirritant

Uses - After Bite

Temporarily protects and helps relieve minor skin irritation and itching due to

- insect bites and stings
- poison ivy, oak or sumac

Warnings - After Bite

Warning: For external use only.

Keep Out of Reach of Children - After Bite

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

Stop Use - After Bite

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

When Using - After Bite

Do not get into eyes

Directions - After Bite

Adults and children under 2 years and older dab directly on bite or sting, rub gently and re-apply as needed

Children under 2 years ask a doctor

Active Ingredient - Aspirin

Aspirin (NSAID*) 325mg

*nonsteroidal anti-inflammatory drug

Purpose

Pain Reliever / Fever Reducer

Uses - Aspirin

Temporarily relieves minor aches and pains associated with: 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 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.

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 change 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 everyday while using this product, take more for a longer time than directed

Do Not Use - Aspirin

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 - Aspirin

Stop Use and ask a Doctor if:

You experience any of the following signs of stomach bleeding, you 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 painful areas, or any new symptoms appear.

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use aspirin during the last 3 months of pregnancy unless 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 - Aspirin

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.

Avoid excessive heat and humidity, do not use any open or torn packets.

First Aid Kit Sleeve

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.

Reorder TAO-001

Genuine Triple Antibiotic

**First Aid Ointment
To Help Prevent Infection**

**Each Gram Contains:
Bacitracin Zinc 400 units
Neomycin Sulfate 5 mg
(equivalent to 3.5 mg
Neomycin base)
Polymyxin B Sulfate 5000 units
Net Wt 0.9g - (1/32 oz)**

CE

 **GFA Production Xiamen Co., Ltd**
No. 20 Huli Industrial Park, Meixi Road,
Tong'an, Xiamen, Fujian, China 361100

 **Wellkang Ltd t/a Wellkang Tech Consulting**
Suite B, 29 Harley Street
LONDON W1G 9QR, England, United Kingdom

Drug Facts - Triple Antibiotic

Each Gram Contains:	Purpose:
Bacitracin Zinc.....	400 units
Neomycin Sulfate.....	5 mg (equivalent to 3.5mg Neomycin base)
Polymyxin B Sulfate	5000 units

Uses: To help prevent infections in minor cuts, scrapes or burns.

Warnings: For external use only

Do not use: in eyes, over large areas of the body, if allergic to any of the ingredients, or 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.

Inactive ingredient: Vaseline...96.41%
Mineral oil.....2%
Purified water.

 **LOT** XXXXXXXXXX
 XXXXXXXXXX

Package Label - Antiseptic

Antiseptic Towelette

Genuine First Aid LLC, Clearwater FL 33755

www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

DRUG FACTS - Antiseptic Towelette

Active Ingredient: Benzalkonium Chloride 0.13% **Purpose:** First Aid Antiseptic

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

Warnings: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, consult a physician immediately.

Do not use: In the eyes, or over large areas of the body.

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

Inactive ingredient: Purified water.

LOT/EXP: XXXXXXXX

TEAR HERE

REORDER AST-001

**Antiseptic
Towelette**
**Toallitas
Antisepicas**



GFA Production Xiamen Co., Ltd
www.gfaproduction.com



GFA Production Xiamen Co., Ltd
No. 20 Hull Industrial Park, Meid Road,
Tong'an, Xiamen, Fujian, China 361100



Wellkang Ltd t/a Wellkang Tech Consulting
Suite 8, 29 Hanley Street
LONDON W1G 9QR, England, United Kingdom



Package Label - Ibuprofen

Overall Dimensions: 1.5" x 2.4"
 Cavity Area: 1.125" x 1.9"
 Max Print Area: 1.075" x 1.75"

Overall Dimensions: 1.5" x 2.4"
 Cavity Area: 1.125" x 1.9"
 Max Print Area: 1.075" x 1.75"

EASY CARE
first aid

Ibuprofen

2 Tablets

Active ingredient (in each tablet) **Purpose**
 Ibuprofen (NSAID) 200mg Pain reliever / fever reducer
 *nonsteroidal anti-inflammatory drug

Uses
 Temporarily relieve minor aches and pains associated with
 headache toothache backache menstrual cramps
 common cold muscular aches minor arthritis pain
 Temporarily reduce fever.

Warnings
 Allergy alert: Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:
 hives rash facial swelling skin reddening
 asthma (wheezing) shock blisters
 If allergic reaction occurs stop use and seek medical help right away.
 Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:
 are age 60 or older have had stomach ulcers or bleeding problems
 take a blood 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.

Store at 68-77°F (20-25°C) • Avoid excessive heat, above 104°F (40°C) • Tamper-evident sealed packets • Do not use any opened or torn packets
 Mfg. for Tenda Corp • Littleton, NH 03561

Warnings (continued)
 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 a stomach bleeding warning applies to you
 you have a history of stomach problems such as heartburn you have high blood pressure, heart disease, liver disease, or kidney disease
 you are taking a diuretic you have problems or serious side effects from taking pain relievers or fever reducers you have asthma

Ask a doctor or pharmacist before use if you are taking aspirin for heart attack or stroke, because ibuprofen may decrease the benefit of aspirin
 taking any other drug under a doctor's care for any serious condition

When using this product: the risk of heart attack or stroke may increase if you use more than directed or longer than directed 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 redness or swelling is present in the painful area
 any new or unexpected symptoms appear

If pregnant or breast-feeding, ask a health professional before use. Use especially important to use ibuprofen during the last 6 months of pregnancy unless specifically 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 Poison Control Center right away.

Directions do not use more than directed the smallest effective dose should be used do not take longer than 10 days unless directed by a doctor (see warnings)

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

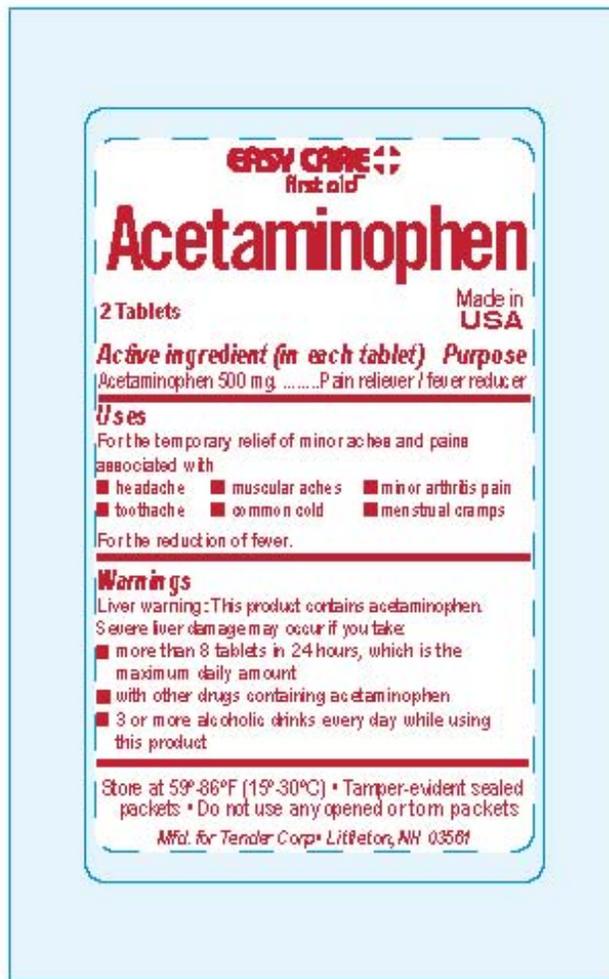
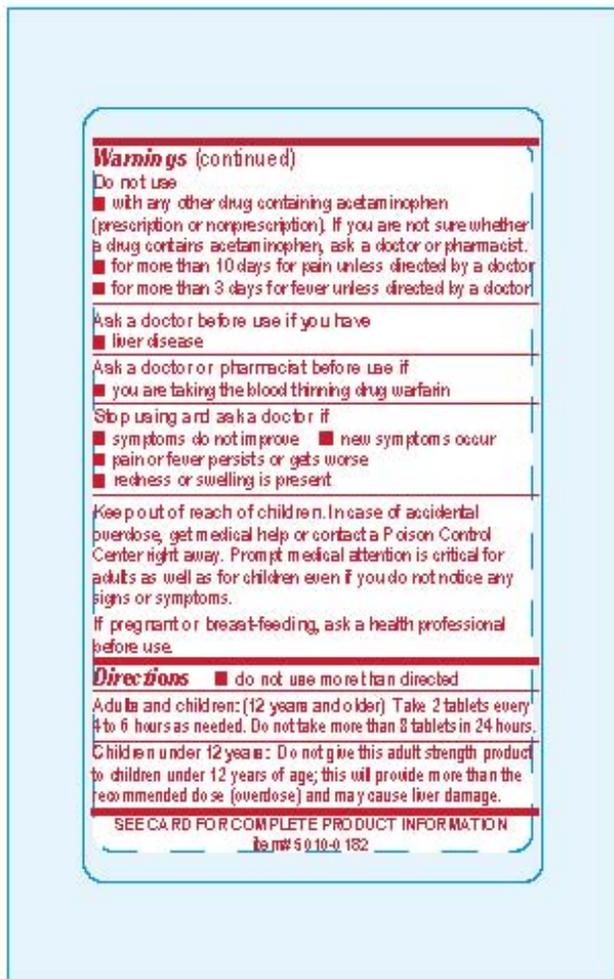
Children under 12 years: Do not give to children under 12 years of age.

SEE CARD FOR COMPLETE PRODUCT INFORMATION
 Item # 50-01-0381

Package Label Non Aspirin

Overall Dimensions: 1.5" x 2.4"
Cavity Area: 1.125" x 1.9"
Max Print Area: 1.075" x 1.75"

Overall Dimensions: 1.5" x 2.4"
Cavity Area: 1.125" x 1.9"
Max Print Area: 1.075" x 1.75"



Package Label - Burn Cream

Genuine First Aid Burn Cream

Antiseptic Pain Relief With Aloe

Net Wt 0.9g (1/32 oz)

Manufactured in CHINA for

Genuine First Aid

GFA Production Xiamen Co., Ltd

No. 20 Huli Industrial Park, Meixi Road, Tong'an, Xiamen, Fujian, China 361100

Tel: 86-592-7269515 Fax: 86-592-7269528 Http: //www.gfaproduction.com

2.125"

2.500"

NDC 044224-0006-2

Child Resistant Package
To Open: Fold across
perforation and tear

AfterBurn[®] gel

Serious Treatment for Burns

- **Cools Burns Fast!**
- **Soothing Clear Aloe Gel**

BE READY for



*kitchen burns, sunburns, windburns,
or any other burn*

Net Wt 1.8 g



Tender Corporation
106 Burndy Rd
Littleton, NH 03561
AfterBurnRelief.com
0005-4452-1

Directions: Spread a thick even layer of AfterBurn over affected area. **Adults and children 2 years of age and older:** apply to affected area not more than 3 to 4 times daily. **Children under 2 years of age:** consult a doctor. **Warnings:** For external use only. When using this product do not get in eyes, do not use in large quantities, particularly over raw surfaces or blistered areas. **Stop use and ask a doctor if** rash, redness, swelling occurs or pain increases, if symptoms persist for more than 7 days or condition clears up and occurs again within a few days. In case of deep or puncture wounds, animal bites or serious burns, consult a doctor. **Active Ingredients:** Lidocaine Hydrochloride: 2.5%, Topical Analgesic.

100% Recycled Paper with 10% Recycled Content, including 5% Post Consumer Waste. Printed in the USA.

Package Label - After Bite

After Bite

The Itch Eraser

Fast Relief from Insect Bites.

Net Contents: 0.037fl. oz.

Contains: One (1) Wipe

Contains Ammonia

Tender Corporation

Littleton, NH 03561

PMS 348

AfterBite®
 The Itch Eraser.com
**Fast Relief from
 Insect Bites.**
**Soulagement rapide
 des piqûres d'insectes.**
 Antipruritic / Antiprurigineux



NET CONTENTS: 0.037 fl. oz.
 CONTAINS/CONTENU: one (1) wipe/serviette
 NPN 02229667 HDC 044224-0001-2
 Contains Ammonia/Contenu Ammoniaque



**Tender
 CORPORATION**
 Littleton, NH 03561 USA

DIRECTIONS: Wipe moist wipelette on bitten area immediately upon opening. Apply with a wiping motion, do not hold on bitten area. Do not bandage or cover tightly until dry. **KEEP OUT OF REACH OF CHILDREN. CAUTION:** for external use only. Avoid mouth, eyes, or mucous membranes. If swallowed, do not induce vomiting. Drink milk and citrus juices and consult a physician. If rash, redness, irritation, swelling or pain increases, discontinue use and consult a physician. Do not apply to wounds or damaged skin. **INGREDIENTS:** Ammonia 3.5% w/v medicinal; Mineral Oil (prevents drying), Alcohol Ethoxylate, Dimethicone, non-medicinal.

MODE D'EMPLOI: retirer la serviette humide de l'emballage et l'utiliser immédiatement pour nettoyer la plaie d'un mouvement fluide, sans presser sur la piqûre. Ne pas panser ou couvrir la plaie encore humide. **HORS DE LA PORTEE DES ENFANTS. ATTENTION:** Pour emploi externe seulement. Évitez le contact avec la bouche, les yeux ou membranes muqueuses. Si le produit est avalé, ne provoquez pas le vomissement. Buvez du lait et des jus d'agrumes et consultez un médecin. Si l'éruption, la rougeur, l'irritation, l'enflure ou la douleur augmente, cessez l'emploi du produit et consultez un médecin. N'appliquez pas sur une blessure ouverte ou peau endommagée. **INGREDIENTS:** Ammoniaque 3.5% w/v (Actif), Médicinal; Huile Minérale (prévient le dessèchement), l'Alcool Ethoxylate, Diméthicone, non-médicinal.

www.tendercorp.com 8005-362-0-1

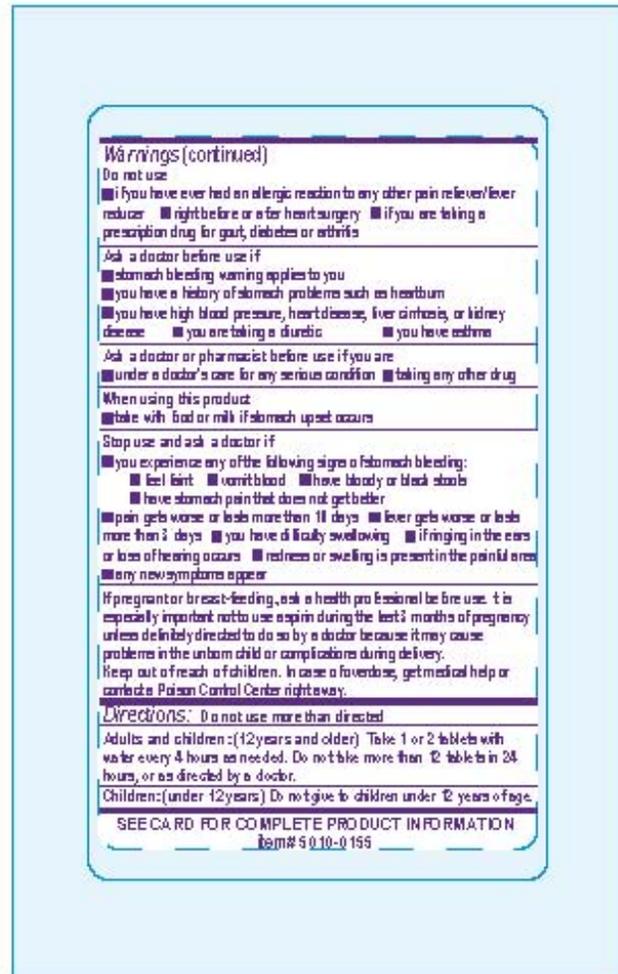
TEAR HERE DÉCHIRURE ICI



Package Label - Aspirin

Overall Dimensions: 1.5" x 2.4"
Cavity Area: 1.125" x 1.9"
Max Print Area: 1.075" x 1.75"

Overall Dimensions: 1.5" x 2.4"
Cavity Area: 1.125" x 1.9"
Max Print Area: 1.075" x 1.75"



REORDER ACP-001



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



GENUINE FIRST AID®



DRUG FACTS - Alcohol Cleansing Pad

Active Ingredient: Isopropyl Alcohol, 70% w/v
Purpose: Antiseptic
Use: For preparation of the skin before injection.
Warnings: For external use only.
Flammable - keep away from fire or flame.
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.
Directions: Wipe injection site vigorously and discard.
Other information: Store at room temperature 15°-30° C (59°-86° F)
Inactive ingredient: Purified water.

--- TEAR HERE ---

CVS ALL PURPOSE FIRST AID KIT

aspirin, diphenhydramine hydrochloride, aspirin, isopropyl alcohol, bacitracin zinc, neomycin sulfate, polymyxin b, benzalkonium chloride, ammonia, lidocaine, acetaminophen, ibuprofen, kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-300
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-300-00	1 in 1 BAG; Type 0: Not a Combination Product	12/28/2016	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 PACKAGE	3 mL in .7
Part 2	1 PACKET	2 mL in 1.8
Part 3	1 PACKET	2

Part 4	1 PACKET	2
Part 5	15 PACKAGE	12 mL in .8
Part 6	6 TUBE	3 g in .5
Part 7	2 PACKET	4
Part 8	17 PACKAGE	12 mL in .7

Part 1 of 8

AFTER BITE WIPE

ammonia swab

Product Information

Item Code (Source)	NDC:44224-0001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
C12-13 ALCOHOLS (UNII: T7ZJT3I9X2)	
WATER (UNII: 059QF0KO0R)	
DIMETHICONE 1000 (UNII: MCU2324216)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-0001-2	0.7 mL in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/28/2016	

Part 2 of 8

AFTER BURN

lidocaine hydrochloride gel

Product Information

Item Code (Source)	NDC:44224-5120
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMO POLYMER TYPE C (UNII: 4Q93RCW27E)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0K00R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-5120-1	1.8 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/12/2011	

Part 3 of 8**MEDIQUE APAP EXTRA STRENGTH**

acetaminophen tablet, film coated

Product Information

Item Code (Source)	NDC:47682-125
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	FR;33
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-125-99	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	12/30/2008	

Part 4 of 8

MEDIQUE IPRIN

ibuprofen tablet, coated

Product Information

Item Code (Source)	NDC:47682-700
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE K30 (UNII: U725QWY32X)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	red (Reddish Brown)	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	G;2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-700-99	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079174	01/26/2016	

Part 5 of 8

ANTISEPTIC

benzalkonium chloride swab

Product Information

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

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0001-1	0.8 mL in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/28/2016	

Part 6 of 8

GENUINE TRIPLE ANTIBIOTIC

bacitracin zinc,neomycin sulfate,polymyxin b sulfate ointment

Product Information

Item Code (Source)	NDC:52124-0003
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1 NDC:52124-0003-1 0.5 g in 1 TUBE; Type 0: Not a Combination Product

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	12/28/2016	

Part 7 of 8

MEDIQUE ASPIRIN

aspirin tablet, film coated

Product Information

Item Code (Source)	NDC:47682-097
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (Round)	Size	10mm
Flavor		Imprint Code	TCL;011
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-097-99	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part343	12/30/2008	

Part 8 of 8

ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information

Item Code (Source)	NDC:52124-0017
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	700 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52124-0017-1	0.7 mL in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/28/2016	

Labeler - CVS (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(69842-300)

Revised: 2/2017

CVS