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

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

Genuine First Aid Auto First Aid Kit

Active Ingredient:

Purpose

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

Use: For preparation of skin before injection.

Warnings: For external use only.

Flammable - keep away from fire or flame

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

Do not use: with electrocautery, in the eyes.

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

Keep out of reach of children. If swallowed,

get medical help or contact a Poison Control

Center right away.

Wipe Injection site vigorously and discard.

Inactive Ingredient: Purified water.

LOT/EXP: Made in CHINA

20140301

Alcohol Cleansing Pad Genuine First Aid LLC, Clearwater FL 33755

www.GenuineFirstAid.com

1/pouch

GENUINE FIRST AID

Active Ingredient:Bacitracin Zinc 400 units

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

Polymyxin B Sulfate 5000 units

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

Warnings:

For external use only.

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

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

Stop use and consult a doctor:

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

Keep out of reach of children.

If ingested, contact a Poison

Control Center right away.

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

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

Active ingredient (in each tablet)PurposeIbuprofen USP (NSAID*) 200mgPain reliever/fever reducer*nonsteroidal anti-inflammatory drug

Uses temporarily relieves minor aches and pains due to:

the common cold

headache

toothache

muscular aches

backache

minor pain of arthritis

Warnings

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

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

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

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

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

Ask a doctor before use if you are:

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

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

Stop use and ask a doctor If:

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

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

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

Directions:

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

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

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

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

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

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

IBUPROFEN 2 Tablets

IBUPROFEN 2 Tablets

Active Ingredient (in each tablet) Purpose

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

Purpose: Pain reliever, fever reducer

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

Warnings

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

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

Ask a doctor before use if you have liver disease

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

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

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

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

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

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

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

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

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

Active Ingredient: Purpose

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

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

Warning: For external use only.

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

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

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

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

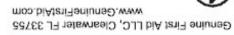
Inactive Ingredient: Purified water

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



GENUINE FIRST AID.

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REORDER AST-001

DRUG FACTS - Antiseptic Towelette

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

Reorder TAO-001

Genuine Triple Antibiotic

First Aid Ointment To Help Prevent Infection

Each Gram Contains: Bacitracin Zinc 400 units Neomycin Sulfate 5 mg (equivalent to 3.5 mg Neomycin base) Polymyxin B Sulfate 5000 units Net Wt 0.5g · (1/64 oz)



Manufactured in CHINA for GENUINE FIRST AID_®

IBUPROFEN^{2 Tablets}

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

Uses temporarily relieves minor aches and pains due to: the common cold headache toothache muscular aches backache minor pain of arthritis menstrual cramps temporarily reduces fever

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

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

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

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

Ask a doctor or pharmacist before use if you are ■ taking any other drug containing an NSAID (prescription or nonprescription)

Drug Facts

Uses: To help prevent infection in • minor cuts • scrapes • burns Warnings:

For external use only Do not use • in eyes • over large areas of the body • if allergic to any of the ingredients • for more than one week unless directed by a physician

Stop use and consult a doctor • if the condition persists or gets

worse • a rash or other allergic reaction develops

Keep out of reach of children. If ingested, contact a Poison

Control Center right away.

Directions: • clean affected area

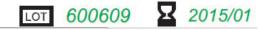
• apply small amount of this product (an amount equal to the surface

area of the tip of a finger) on the area 1 to 3 times daily • may be

covered with a sterile bandage

Other information:

Store at room temperature



Warnings (continued)

■ taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin ■ taking any other drug

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

Stop use and ask a doctor if up you experience any of the following signs of stomach bleeding: feel faint vomit blood

■ have bloody or black stools ■ have stomach pain that does not get better ■ pain gets worse or lasts more than 10 days ■ fever gets worse or lasts more than 3 days ■ redness or swelling is present in the painful area ■ any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

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

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

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

Other information ■ store at controlled room temperature ■ avoid excessive heat 40° C(104° F) ■ tamper evident sealed packets ■ do not use any opened or torn packets

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

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



DRUG FACTS - Alcohol Cleansing Pad

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

20140301

| Warnings (continued) Do not use |
|---|
| with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure wheth a drug contains acetaminophen, ank a doctor or pharmaci for more than 10 days for pain unless directed by a doct for more than 3 days for fever unless directed by a doct |
| Ask a doctor before use if you have in liver disease |
| Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin |
| Stop using and ask a doctor if symptoms do not improve pain or fever persists or gets worse redness or swelling is present |
| Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms. |
| If pregnant or breast-feeding, ask a health professional before use. |
| Directions a do not use more than directed |
| Adults and children: (12 years and older) Take 2 lablets every 4 to 6 hours as needed. Do not take more than 8 lablets in 24 hours. |
| Children under 12 years: Do not give to children under 12 years of age. |
| |

SEE CARTON FOR COMPLETE PRODUCT INFORMATION

2 Tablets

Active Ingredient (in each tablet) Purpose Acetaminophen 500 mg. Pain relieventiever reducer

Médique

Extra Strength

Uses

For the temporary relief of minor aches and paine associated with

headacha muscular aches

minor artuntis pain

menstrual cramps

Made in

IS A

For the reduction of fever.

Il toothacha III common coid

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Store at 59°-86°F (15°-30°C) • Tamper-evident sealed packets • Do not use any opened or tom packets and, for MEDIQUE PRODUCTS • Fort Myers, FL 33967





GENUINE FIRST AID - AUTO FIRST AID

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

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

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

IBUPROFEN

| POWDERED CELLUIOSE (UNE: SMD1/33X09M) STARCI, CORN (UME: 00222Y35) HYPROMEL.0085 (UNE: SWX29/34W0) LACTOSE (UNE: 228244986) MGENSEUM STEARATE (UME: 70097M610) LACTOSE (UNE: 728244986) MGENSEUM STEARATE (UME: 70097M610) POVIDOXE (UME: 728244986) POVIDOXE (UME: 7282497M610) POVIDOXE (UME: 51926901841) SILICON DIOXIDE (UME: 51926901842) SILICON DIOXIDE (UME: 51926901844) SILIC | PowDERED CELLULOSE (UNIE SMDIX3XO9M) STARCI, CORN (UNIE 09232NYSS) STARCI, CORN (UNIE 09232NYSS) LACTOSE (UNIE 1282A4N98G) AGORSE (UNIE 1282A4N98G) AGORSE (UNIE 12802A4N98G) STARCI (UNIE 12982A4N98G) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 10997A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1097A62D) SOLUTION STARCITE (UNIE 1007A62D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A662D) SOLUTION STARCITE (UNIE 1007A667D) SOLUT | Inactive Ingredien | Ingredient Name | ρ | | Strength |
|---|---|---|--|--------------|-----------------|--------------------|
| STARCH, CORN (UNE: 08232NY3SJ) | STARCIL CORN (UNII: 082322Y35) Image: content of the state of | POWDERED CELLULO | | | | ottengti |
| HYPRO MELLOSES (UNIE 3NXW29 V3WO) Image: Status Statu | INTROMELLOSES (UNIE 3NXW29 V3WO) Image: Stream (Stream | | | | | |
| LACTO SE (UNIE J2B2A4N98G) MGGNESUUM STEARATE (UNIE 7097/MEISO) POLYETPKYLENE GLYCOL (UNIE 3W/Q0 SDW1A) POVTOONE (UNIE ETJ726 SDW) SULCON DIO XODE (UNIE ETJ726 SDW) STEARIC ACID (UNIE ETJ726 SDW) STEARIC ACID (UNIE ETJ726 SDW) STRACE TIN (UNIE ETJ726 SDW) TTANUM DIO XIDE (UNIE ETJ726 SDW) TAUEST Part 3 of 5 MECI47682-175 TEJ CATEGRA STRENGTH acetaminophen tablet, film coated NDC:47682-175 TEJ CATEGRA STRENGTH TEM CODE (SOURCE) NDC:47682-175 | LACTOSE (UNII: J2B2A4N98G) MAGNESIUM STEARATE (UNII: 70097M6B0) PO LYDEXTROSE (UNII: 70097M6B0) PO LYDEXTROSE (UNII: 84//Q0SDW1A) PO VIDONE (UNII: 847/758AP) SULCON DIO XIDE (UNII: ETY726XBUA) STRATIC ACTO (UNII: 847/758AP) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATION DIO XIDE (UNII: ETY726XBV) TTAATION DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATE TTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: ETY726XBV) TTAATE TTANIUM DIO XIDE (UNII: ETY726XBV) TTTANIUM DIO XIDE (UNII: E | | | | | |
| PO LYDEXTROSE (UNIE VH2 XOU 12/E) PO LYDEXTROSE (UNIE VEX XOU 12/E) PO VIDONE (UNIE F2989GIB4E) STEARIC ACLD (UNIE VEX ETTYZ & STUAJ STEARIC ACLD (UNIE VEL VT265AP) TTTANUM DIO XDE (UNIE ISFK9 V2.P) TTTANUM DIO XDE (UNIE ISFK9 V2.P) TRIACETIN (UNIE XIIX3C3X6 73) Product Characteristics Color white (WHITE) Score no score Shape ROUND Size Imprint Code Imprint Code Flavor Imprint Code Size Imprint Code Flavor Imprint Code Imprint Code Imprint Code Flavor Imprint Code Imprint Code Imprint Code Packageing # Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:52124-0009-1 2 in 1 PACKET Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date 1 NDC:52124-0009-1 2 in 1 PACKET Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Flavor Imprint Code Imprint Code Product Information Imprint Code Imprint C | PO LYDEXTRO SE (UNIE VH2XOU 12IE) | LACTOSE (UNII: J2B2A | 4N98G) | | | |
| POLYETHYLENE GLYCOL (UNI: 3WJQ05DWIA) POVIDONE (UNI: ET3726389G/E94E) SILICON DIO XIDE (UNI: ET3726389G/E94E) TTANIUM DIO XIDE (UNI: 15FIX9V2JP) TTANIUM DIO XIDE (UNI: 15FIX9V2JP) TRIACETIN (UNI: XHX3C3X673) POUND KINK KIX3C3X673) POUND KINK KIX3C3X673 POUND KINK KIX3C3X673 POUND Size No score Shape ROUND Size No score Shape ROUND Size No score Shape ROUND Size No score Shape ROUND Size No score Packaging # Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:52124-0009-1 2 in 1 PACKET Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA0750 01/11/2011 Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Pound ANDA ANDA07501 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Soure) NDC:47682-175 | PO LYETHYLENE GLYCOL (UNE: 303,003 DW1A) − − − − − − − − − − − − − − − − − − − | MAGNESIUM STEARAT | °E (UNII: 70097M6I30) | | | |
| PO VIDO NE (UNIE F2989GH94E) SILEON DIO XIDE (UNIE F1726 XBU4) STEARIC ACID (UNIE 4EL V726 XBU4) STEARIC ACID (UNIE 4EL V726 5AP) - STEARIC ACID (UNIE 15R3 V2JP) - STEARIC ACID | PO VIDO NE (UNIE PT298 9G H9 4E) | POLYDEXTROSE (UNII | : VH2XOU12IE) | | | |
| SILICON DIO XIDE (UNII: ETJ7Z6 XBU4) → · · · · · · · · · · · · · · · · · · | SILICON DIO XIDE (UNII: ETI7Z6 XBU4) | POLYETHYLENE GLY(| C OL (UNII: 3WJQ0SDW1A) | | | |
| STEARIC ACID (UNII: 4EL V726 5AP) TTANUOM DIO XIDE (UNII: 15FIX9 V21P) TRIACETIN (UNII: XHX3C3X6 73) Product Characteristics Color white (WHITE) Score no score Shape ROUND Size 100mm Flavor 044;352 Contains 144;352 Contains 144;352 Packag ing # Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:52124-0009-1 2 in 1 PACKET 144;352 Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA 75010 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Soure) 10/10/2015 | STEARIC ACID (UNII: 4EL V7265AP) Image: Content of the state of | POVIDONE (UNII: FZ98 | 9 GH9 4E) | | | |
| TTANIUM DIOXIDE (UNE: 15FK9 Y2JP) TRIACETIN (UNE: XHX3C3X673) Product Characteristic status and | TTANUM DIO XIDE (UNIE: ISFIX9 V2/P) TRIACETIN (UNIE: XHX3C3X673) | SILICON DIOXIDE (UN | II: ETJ7Z6XBU4) | | | |
| TRIACETIN (UNIE XHX3C3X673) Product Characteristic Color white (WHTE) Score no score Shape ROUND Size 10mm Flavor ROUND Size 44;352 Contains Imprint Code 44;352 Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code Package Description Marketing Start Date Marketing End Date Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Part 3 of 5 MEDIQUE APART STRENGTH acetaminophen tablet, film coated Product Information NDC:47682-175 | TRIACETIN (UNIE XHX3G3X673) Note: State S | STEARIC ACID (UNII: 4 | ELV7Z65AP) | | | |
| Product Characteristics Color vhite (WHITE) Score no score no score Shape ROUND ROUND Size Umprint Code V4:352 Omm V4:352 V V Packaging Item Code Package Description Marketing Start Date Marketing End Date I NDC:52124-0009-1 2 in 1 PACKET V V V V V V V V V V V V V V V V V V V | Product Characteristics view of white (WHTE) core of source of shape and the (WHTE) core core of source o | TITANIUM DIO XIDE (U | NII: 15FIX9V2JP) | | | |
| Colorwhite (WHITE)Scoreno scoreShapeROUNDSize10mmFlavorImprint Code (Warden)44,352ContainsImprint Code (Source)Variable (Source)PackagingPackage DescriptionMarketing Start DateMarketing End DateI NDC:52124-0009-12 in 1 PACKETVariable (Source)Marketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DatePart 3 of 5Variable (Source)Variable (Source)Variable (Source)Variable (Source)Marketing Code (Source)MDC:47682-175Variable (Source)Variable (Source) | Colorwhite (WHITE)Scoreno scoreShapeROUNDSize10mmFlavorImprint Code44352ContainsImprint CodeVPackagingImprint CodeNarketing Start DateMarketing End DateI NDC:52124-0009-12 in 1 PACKETImprint CodeMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DatePart 3 of 5STRENGTH01/11/2011Imprint CodeImprint CodeProduct InformativeSTRENGTHStrengen VStrengen VImprint CodeProduct InformativeImprint Code (Source)Imprint Code (Source)Imprint CodeImprint Code | TRIACETIN (UNII: XHX3 | C3X673) | | | |
| Color white (WHT) → Score no score Shape RUUN Size 10mm Flavor Imprint Code (Source) 4332 Packaging Imprint Code 4352 Packaging Item Code Package Description Marketing Start Date Marketing End Date I NDC:52124-0009-1 2 in 1 PACKET V Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date NDA ANDA075010 U11/2011 Marketing Start Date Marketing End Date Part 3 of 5 S S S S S WEDIQUE AP + EXTRENGTH S S S S Product Informatible NDC:47682-175 S S S | Color white (WHITE) Score no score Shape ROUND Size 10mm Flavor Imprint Code 4/352 Contains Imprint Code 4/352 Packaging Imprint Code V I Indo: Scite Good Package Description Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Bert 3 of 5 Street Stree | | | | | |
| Shape ROUND Size 10mm Flavor Imprint Code Imprint Code 44,352 Contains V V V Packaging Item Code Package Description Marketing Start Date M Marketing Information 1 NDC:52124-0009-1 2 in 1 PACKET V Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Part 3 of 5 MEDIQUE APAP EXTRENGTH Arburder table, filter STRENGTH Start Strength Product Information Imprint Code (source) MDC:47682-175 | Shape ROUND Size 10mm Flavor Imprint Code 44,352 Contains Imprint Code 44,352 Packaging * Item Code Package Description * Item Code 2 in 1 PACKET Marketing Informative Marketing Category Application Number or Monograve Marketing Start Date Marketing Category Application Number or Monograve Marketing Start Date Marketing Category AnDA Andor5010 01/11/2011 Product Informative Kerr STRENGTH MC:47682-175 | Product Character | istics | | | |
| Flavor imprint Code 44;352 Contains Imprint Code 44;352 Packaging Item Code Package Description Marketing Start Date Marketing End Date # Item Code Package Description Marketing Start Date Marketing End Date I NDC:52124-0009-1 2 in 1 PACKET Marketing Start Date Marketing End Date Marketing Information Application Number or Monograph Citation Marketing Start Date Marketing End Date NDA ANDA750 10 01/11/2011 Marketing End Date Marketing End Date Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH Marketing coated Marketing End Date Product Information NDC:47682-175 NDC:47682-175 Marketing End | Flavor Imprint Code 44;352 Contains Packaging Imprint Code 44;352 Packaging Imprint Code Marketing Start Date Marketing End Date * Item Code Package Description Marketing Start Date Marketing End Date Marketing Information Application Number or Monograph Citation Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 Marketing End Date | Color | white (WHITE) | Score | | no score |
| Contains Item Code Packaging # Item Code Package Description Marketing Start Date Marketing Category Application Number or Monograph Citation Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing Start Date Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date Marketing Start Date Marketing End Date Marketing Start Date Marketing End Date Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date NDA ANDA75010 | Contains Packaging # Item Code Package Description Marketing Start Date Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date | Shape | ROUND | Size | | 10 mm |
| Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code 2 in 1 PACKET Image: Comparison of the tem Code Comparison of tem Comparison of tem Code Comparison of | Packaging # Item Code Package Description Marketing Start Date Marketing End Date t NDC:52124-0009-1 2 in 1 PACKET Image: Constant Constan | Flavor | | Imprint Co | de | 44;352 |
| Packaging # Item Code Package Description Marketing Start Date Marketing End Date # Item Code 2 in 1 PACKET Image: Comparison of the tem Code Comparison of tem Comparison of tem Code Comparison of | Packaging # Item Code Package Description Marketing Start Date Marketing End Date t NDC:52124-0009-1 2 in 1 PACKET Image: Constant Constan | Contains | | | | |
| Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End D ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information MDC:47682-175 | Marketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information MDC:47682-175 | | | | | |
| Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA75010 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate FYTHENGTH ANDA MOLESSION OF THE STRENGTH ANDA MARETEX STRENGTH ANDA MARETEX Product Information MC:47682-175 | Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coated Product Information MDC:47682-175 | | Package Description | Marketii | ng Start Date | Marketing End Date |
| Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA7551 01/11/2011 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate Product Information MDC:47682-175 | Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date ANDA ANDA075010 01/11/2011 01/11/2011 Part 3 of 5 MEDIQUE APARETEXTRENGTH acetaminophen table: Film coate Product Information MDC:47682-175 | # Item Code | . | Marketii | ng Start Date | Marketing End Date |
| ANDA ANDA O75010 01/11/2011 Part 3 of 5 MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | ANDA ANDAO 750 10 01/11/20 11 Part 3 of 5 MEDIQUE APAP EXTRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 | 2 in 1 PACKET | Marketin | ng Start Date | Marketing End Date |
| MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 | 2 in 1 PACKET | | | |
| MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | MEDIQUE APAP EXTRA STRENGTH acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 Warketing Info Marketing Category | 2 in 1 PACKET rmation Application Number or Monogr | | Marketing Start | |
| acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 Warketing Info Marketing Category | 2 in 1 PACKET rmation Application Number or Monogr | | Marketing Start | |
| acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | acetaminophen tablet, film coated Product Information Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA | 2 in 1 PACKET rmation Application Number or Monogr | | Marketing Start | |
| Product Information Item Code (Source) NDC:47682-175 | Product Information Item Code (Source) NDC:47682-175 | <pre># Item Code 1 NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5</pre> | 2 in 1 PACKET rmation Application Number or Monogr ANDA0 750 10 | aph Citation | Marketing Start | |
| Item Code (Source) NDC:47682-175 | Item Code (Source) NDC:47682-175 | Item Code I NDC:52124-0009-1 Marketing Info Info Marketing Category Info ANDA Info Part 3 of 5 Info MEDIQUE APA | 2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH | aph Citation | Marketing Start | |
| Item Code (Source) NDC:47682-175 | Item Code (Source) NDC:47682-175 | Item Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA | 2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH | aph Citation | Marketing Start | |
| | | Item Code I NDC:52124-0009-1 Marketing Info Info Marketing Category Info ANDA Info Part 3 of 5 Info MEDIQUE APA | 2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH | aph Citation | Marketing Start | |
| Route of Administration ORAL | Route of Administration ORAL | # Item Code 1 NDC:52124-0009-1 Marketing Info Info Marketing Category ANDA Part 3 of 5 MEDIQUE AP A Acetaminophen tablet Info | 2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH , film coated | aph Citation | Marketing Start | |
| | | # Item Code 1 NDC:52124-0009-1 Marketing Info Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA Acetaminophen tablet Information | 2 in 1 PACKET rmation Application Number or Monogr ANDA075010 AP EXTRA STRENGTH , film coated | aph Citation | Marketing Start | |
| | | I tem Code I Tem Code NDC:52124-0009-1 Marketing Info Marketing Category ANDA Part 3 of 5 MEDIQUE APA acetaminophen tablet Product Informatie Item Code (Source) | 2 in 1 PACKET Timation Application Number or Monogr ANDA075010 ANDA075010 ANDA075010 NDC:47682-175 | aph Citation | Marketing Start | |

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

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

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

Labeler - Genuine First Aid, LLC (619609857)

Revised: 1/2011

Genuine First Aid, LLC