EASY CARE FIRST AID KIT - COMPREHENSIVE- ammonia, aspirin, diphenhydramine hydrochloride, ibuprofen, acetaminophen, lidocaine hydrochloride, benzalkonium chloride
Tender Corporation

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.

Easy Care Comprehensive First Aid Kit

Active Ingredients - After Bite Wipe

Ammonia 3.5%

Purpose - After Bite Wipe

Counterirritant

Use - After Bite Wipe

Temporarily protects and helps relieve minor skin irritation and itching due to - insect bites and stings, poison ivy, oak or sumac

Warnings - After Bite Wipes

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 Wipe

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 Wipe

Adults and Children 2 years and older - dab directly on bite or sting, rub gently and reapply as needed

Children under 2 years - ask a doctor

Active Ingredient - Aspirin

Aspirin (NSAID*) 325mg *nonsteroidal anti-inflammatory drug

Purpose - Aspirin

Pain Reliever/Fever Reducer

When Using - Aspirin

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 or shock. If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning: this contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you: are age 60 or older; have had stomach ulcers or bleeding problems, take a blood thinner (anticoagulant) or steroid drug; take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen or others), have 3 or more alcoholic drinks everyday while using this problem, take more for a longer time than directed.

Do not use if you have ever had an allergic reaction to any other pain reliever/fever reducer; right before or after heart surgery, if you are taking a prescription drug for gout, diabetes or arthritis.

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

Ask a Doctor or Pharmacist before use if you are: under a doctor's care for any serious condition, taking any other drug.

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

Stop use and Ask a Doctor if: you experience any of the following signs of stomach bleeding, you feel faint, vomit blood, have bloody or black stools, have stomach pain that does not get better, 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 Breastfeeding, 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 deliver.

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 dosage 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 as 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.

Use - Aspirin

Temporariy relieves minor aches and pains associated with: headache, muscular aches, minor arthritis

pain, backache, common cold, toothache, menstrual cramps, temorarily reduces fever.

Active Ingredient - Ibuprofen

Ibuprofen

Purpose - Ibuprofen

Pain Reliever/Fever Reducer

Use - 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 and Precautions - 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, and 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, fight before or after heart surgery.

Ask a doctor before use if stomach bleeding warning applies to you, you have had 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, feeling 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.

Store at controlled room temperature, avoid excessive heat 40 degrees Celsius (104 degrees Fahrenheit); temper evident sealed packets, do not use any opened or torn packets.

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, 3 tablets may be use. Do not

exceed 6 tablets in 24 hours, unless directed by a doctor. Children under 12 years: do not give children under 12 years of age.

Active Ingredient - APAP

Acetaminophen 500mg

Purpose - APAP

Analgesic/antipyretic

Uses - APAP

Temporary relief of minor aches and pains associated with: common cold, headache, toothache, muscular aches, backache, arthritis, menstrual cramps, and reduction of fever

Warnings and Precautions - APAP

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 ever day while using this product.

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

Ask a doctor before use if the user has liver disease.

Ask a doctor or pharmacist before use if the user is taking the blood thinning drug warfarin.

Stop use and ask a doctor if: symptoms do not improve, pain gets worse or lasts more than 10 days, fever gets worse or lasts more than 3 days, new symptoms occur, redness or swelling is present, a rare sensitivity reaction occurs.

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

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt medical attention is critical for adults as well for children even if you do not notice any signs or symptoms. Do not exceed recommended dosage.

Store at 59-86 degrees F (15-30 degrees C), tamper evident sealed packets, do not use any open or torn packets.

Directions - APAP

Adult and Children (12 years and older) take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours.

Children 6-11 years take 1 tablet every 4 to 6 hours as needed. Do not take more than 5 tablets in 24 hours.

Childen under 6 do not use this product

Active Ingredient - Diphenhydramine

Diphenhydramine Hydrochloride 25mg

Purpose - Diphenhydramine

Antihistamine

Use - Diphenhydramine

Temporarily relieves these symptoms due to hay fever or other respiratory allergies: runny nose, sneezing, itching nose or throat, itchy-watery eyes.

Temporarily relieves these symptoms due to the common cold: runny nose, sneezing

Warnings and Precautions - Diphenhydramine

Do not use to make a child sleepy or with any other product containing diphenhydramine, even one that is used on skin.

Ask a doctor before use if you have: a breathing problem such as emphysema or chronic bronchitis, difficulty in urination due to enlargement of the prostate gland, or glaucoma.

Ask a doctor pharmacist before you use if you are taking sedatives or tranquilizers.

When using this product: marked drowsiness may occur, avoid alcoholic beverages, alcohol, sedatives and tranquilizers may increase the drowsiness effect, use caution when driving a motor vehicle or operating machinery, excitability may occur, especially in children.

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

Keep out of reach of Children. In case of overdose, contact a physician or poison control center immediately.

Each caplet may contain calcium 25mg. Store at room temperature 59-86 degrees F (15-30 C), protect from light and use by expiration date on packet. Tamper-evident sealed packets, do not use any opened or torn packets.

Directions - Diphenhydramine

Do not use more than directed.

Adults and Children (12 Years and older) - take 1 to 2 caplets every 4 to 6 hours as needed. Do not take more than 12 caplets in 24 hours, or as directed by a doctor.

Do not give to children under 12 years of age.

Active Ingredients - After Cuts and Scrapes

Lidocaine Hydrochloride 2.5%

Benzalkonium Chloride .13%

Purpose - After Cuts and Scrapes

Anesthetic

Antiseptic

Uses - After Cuts and Scrapes

First aid for the temporary relief of pain and to protect from infection in minor cuts, scrapes and burns.

Warnings and Precautions - After Cuts and Scrapes

For External Use Only.

When using this product: do not get into eyes, do not use in large quantities, particularly over raw surfaces or blistered areas.

Stop use and ask a doctor if rash, redness or swelling occurs or pain increases. In case of deep puncture wounds, animal bites or serious burns consult a doctor.

If swallowed, get medical help or contact a poison control center right away.

If in eyes, flush with water for 15 minutes and call a doctor.

Keep out of Reach of Children.

Directions - After Cuts and Scrapes

Shake Well. Clean affected area. Spray on the area 1 to 3 times daily. Adults and children 2 years and older, apply to affected area. Children under 2 years, ask a doctor.

Active Ingredient - After Burn

Lidocaine Hydrochloride 2.5%

Purpose - After Burn

Topical Analgesic

Use - After Burn

For the temporary relief of pain due to minor burns, sunburn, minor cuts, scrapes or minor skin irritations

Warnings and Precautions - After Burn

For external use only.

When using this product: avoid contact with eyes, do not use in large quantities, particularly over raw surfaces or blistered areas, do not use longer than 1 week unless directed by a doctor, or in case of deep puncture wounds, animal bites or serious burns, consult a doctor.

Stop use and ask a doctor if conditions or symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of Reach of Children.

If swallowed, get medical help or contact a poison control center right away.

If in eyes, flush with water for 15 minutes and call a doctor

Adults and Children 2 years and older, apply small amount to affected area, not more than 3 times daily.

Children under 2 years, consult a doctor.

Inactive Ingredients - After Burn

Aloe Vera Gel, Benzalkonium Chloride, Carbomer 980, glycerol, USP, herbal fragrance, menthol, polysorbate 20, Propylene glycol, purified water, sodium hydroxide (50%)

Package Labels

NEW WITH ALOE VERA

Family Medic

AFTER CUTS & SCRAPES™

Topical Analgesic/Antiseptic
First-Aid Treatment Spray Formula

AFTER CUTS & SCRAPES™

Topical Analgesic/Antiseptic
First-Aid Treatment Spray Formula

- Temporarily relieves pain from minor cuts, scrapes and burns
- Helps prevent infection

Drug Facts

Active Ingredients	Purpose
Lidocaine Hydrochloride 2.50%	Anesthetic
Benzalkonium Chloride 0.13%	Antiseptic

Uses First aid for the temporary relief of pain and to protect from infection in minor cuts, scrapes and burns

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. In case of deep or puncture wounds, animal bites or serious burns consult a doctor.

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

If in eyes flush with water for 15 minutes and call a doctor.

Directions
Shake Well. Clean affected area. Spray on the area 1 to 3 times daily
Adults and children 2 years and older; apply to affected area
Children under 2 years; consult a doctor

Inactive Ingredients
Aloe Vera Gel, Germaben II, Orange Blossom Fragrance, Polysorbate 20, Propylene Glycol, Purified Water, and Sorbitol 70%

Tender CORPORATION
LITTLETON, NH 03561 Made in USA
See a complete line of our products for the prevention and treatment of outdoor ailments at:
www.tendercorp.com

02-124ORO

0 44224 03300 0

- Relieves pain from minor cuts, scrapes and burns
- Helps prevent infection

FOR CUTS • SCRAPES • AND BURNS

Family Medic®

AfterBurn®
Serious Treatment for Burns

NDC 044224-0006-1

COOLS BURNS FAST!

COOLS BURNS FAST!

AfterBurn®
Serious Treatment for Burns

AfterBurn®
Serious Treatment for Burns

ADVANCED PAIN RELIEF
with Lidocaine HCl 2.5%
Clear, Soothing Aloe Gel

BE READY for
Sunburns
Scalds
Stove Burns
and other burns

ADVANCED PAIN RELIEF
with Lidocaine HCl 2.5%
Clear, Soothing Aloe Gel

BE READY for
Sunburns
Scalds
Stove Burns
and other burns

Net wt: 2 oz (57g) Tube

0 44224 05120 2

AfterBurn®
Serious Treatment for Burns

AfterBurn®
Serious Treatment for Burns

AfterBurnRelief.com

CLEAR ALOE GEL SOOTHES SKIN

COOLS BURNS FAST!

This soothing, clear gel provides immediate pain relief from kitchen burns, sunburns or any other burns, scrapes and cuts. **AfterBurn®** contains Lidocaine to relieve pain instantly and Aloe to soothe and moisturize the skin.

Drug Facts

Active Ingredient	Purpose
Lidocaine Hydrochloride 2.50%	Topical Analgesic

Uses For the temporary relief of pain due to minor burns, sunburn, minor cuts, scrapes, or minor skin irritations

Warnings

For external use only

When using this product
- avoid contact with eyes
- do not use in large quantities, particularly over raw surfaces or blistered areas
- do not use longer than 1 week unless directed by a doctor
- In case of deep puncture wounds, animal bites, or serious burns, consult a doctor

Stop use and ask a doctor if
- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days

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

If in eyes flush with water for 15 minutes and call a doctor

Directions
- Clean the affected area
- **Adults and children 2 years of age and older** Apply small amount to affected area not more than 3 times daily
- **Children under 2 years of age** consult a doctor

Inactive Ingredients
Aloe Vera, Benzalkonium Chloride, Carbomer 980, Glycerol USP, Herbal Fragrance, Menthol, Polysorbate 20, Propylene Glycol, Purified Water, and Sodium Hydroxide

0002-1242-3

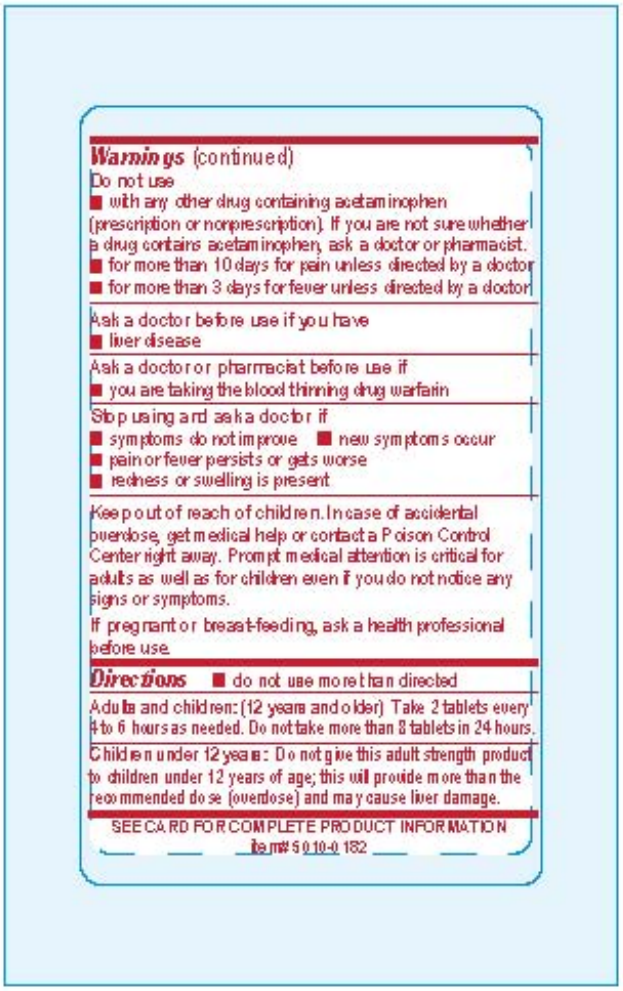
from the makers of AfterBite®

Tender CORPORATION
Littleton, NH 03561
Made in the USA

www.tendercorp.com

Tender Corp.

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



Warnings (continued)

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

Ask a doctor before use if 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 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

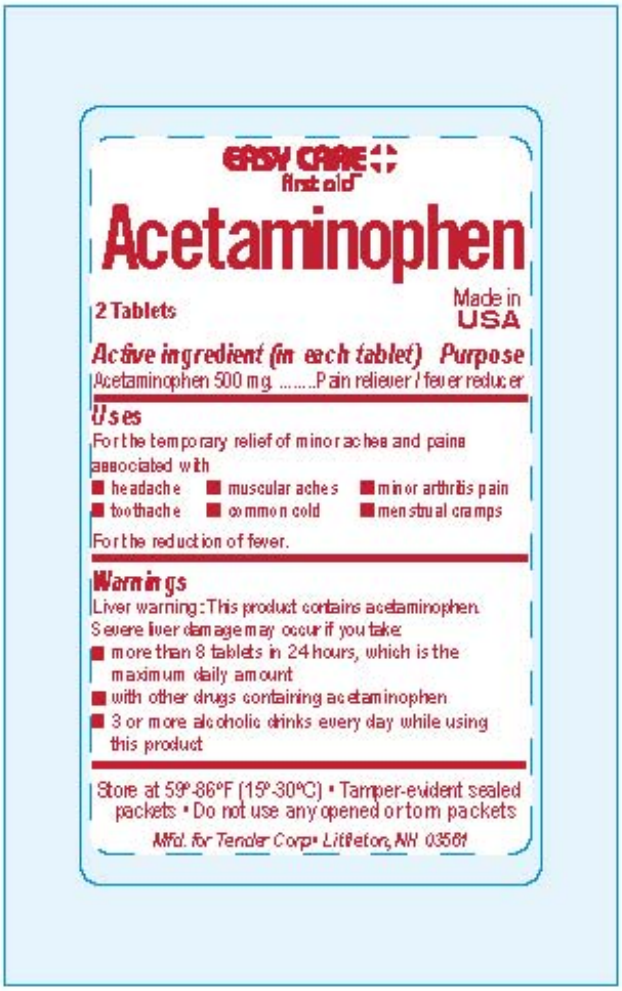
- do not use more than directed

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 this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

SEE CARD FOR COMPLETE PRODUCT INFORMATION

Item # 5010-0182



EASY CARE first aid

Acetaminophen

2 Tablets

Made in USA

Active ingredient (in each tablet) Purpose

Acetaminophen 500 mg .....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. 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 torn packets

Mfd. for Tender Corp • Littleton, NH 03561

PMS 348

AfterBite®

The Itch Eraser®.com

Fast Relief from Insect Bites.

Soulagement rapide des piqûres d'insectes.

Antipruritic / Antiprurigineux

Tender CORPORATION

106 Bonnie Rd, Littleton, NH 03561 USA

NET CONTENTS: 0.037 fl. oz.
CONTAINS/CONTENU: one (1) wipe/serviette
NPN 02229867 NDC 044224-0001-2
Contains Ammonia/Contient Ammoniaque

Imported By:
Trans Canada Distribution Inc.
4-3520 Laird Rd.
Mississauga, ON L5L 5Z7 Canada

DIRECTIONS: Wipe moist towelette 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.

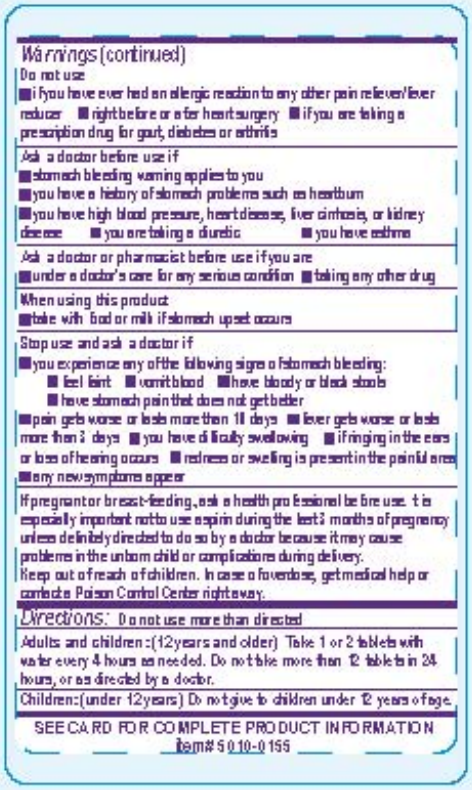
MODE D'EMPLOI: retirer la serviette humide de l'emballage et l'utiliser immédiatement pour nettoyer la piqûre d'un mouvement fluide, sans pression sur la piqûre. Ne pas panser ou couvrir la piqûre encore humide. HORS DE LA PORTÉE DES ENFANTS. ATTENTION: Pour emploi externe seulement. Éviter le contact 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. INGRÉDIENTS: Ammoniaque 3.5% w/v (Actif), Médicinal; Huile Minéral (prévient la dessèchement), l'Alcool Éthoxylate, Méticone, non-médicinal.

0805-3820-2

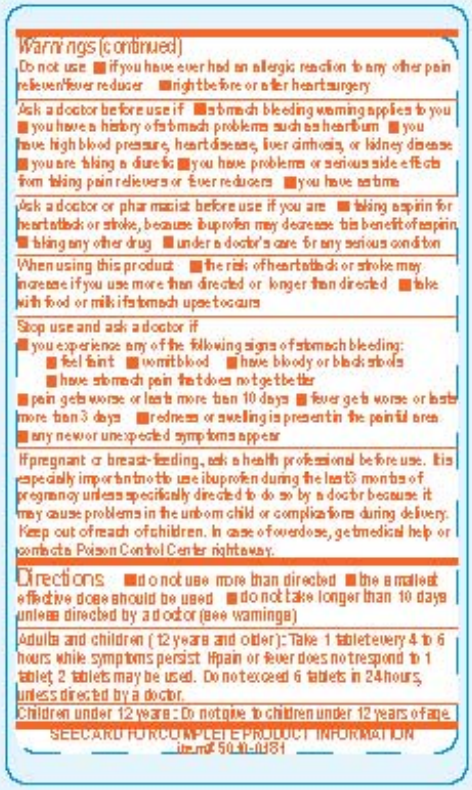
www.tendercorp.com

TEAR HERE DÉCHIRURE ICI

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

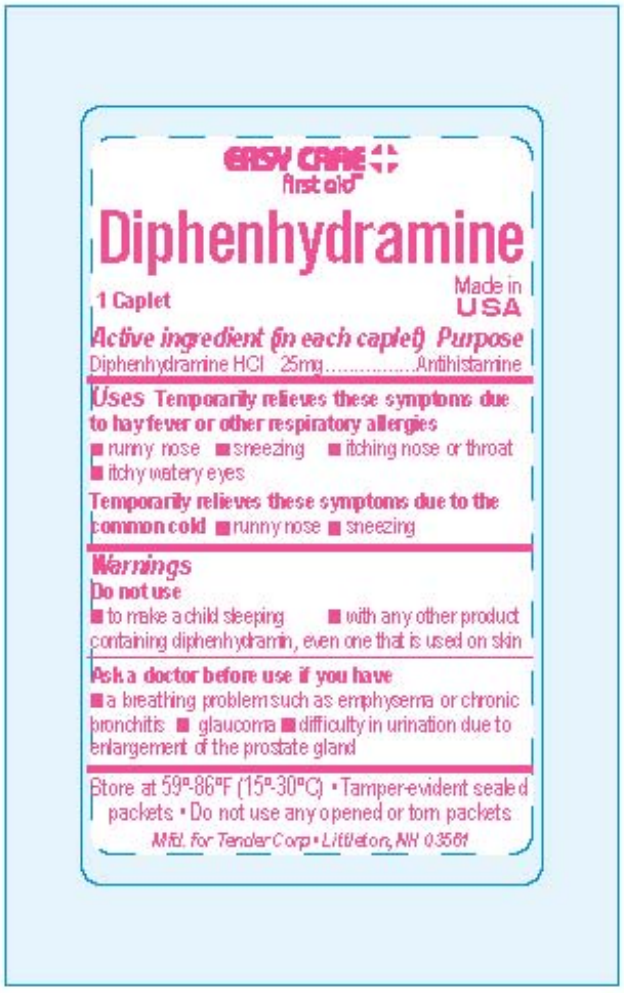


Overall Dimensions: 1.5″ x 2.4″
Cavity Area: 1.125″ x 1.9″
Max Print Area: 1.075″ x 1.75″



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 KIT - COMPREHENSIVE

ammonia, aspirin, diphenhydramine hydrochloride, ibuprofen, acetaminophen, lidocaine hydrochloride, benzalkonium chloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44224-2999

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:44224-2999-0	1 in 1 BOX; Type 0: Not a Combination Product	01/01/2012	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	0 TUBE	1 mL in 59

Part 2	1 PACKET	2
Part 3	1 PACKET	2
Part 4	2 PACKET	2
Part 5	0 BOTTLE	1 mL in 30
Part 6	1 PACKET	2
Part 7	4 PACKAGE	3 mL in .7

Part 1 of 7

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 HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
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:44224-5120-0	59 mL 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 not final	part348	04/12/2012	

Part 2 of 7

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
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)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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 3 of 7

MEDIQUE ASPIRIN

aspirin tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-097
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MINERAL OIL (UNII: T5L8T28FGP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
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 4 of 7

MEDIQUE DIPHEN

diphenhydramine hydrochloride tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-184
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics			
Color	pink (pink)	**Score**	no score
Shape	OVAL (OVAL)	**Size**	11mm
Flavor		**Imprint Code**	061;T
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47682-184-46	1 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	part341	01/01/2012	

Part 5 of 7

AFTER CUTS AND SCRAPES

lidocaine hydrochloride, benzalkonium chloride solution

Product Information	
Item Code (Source)	NDC:44224-3300
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	26 mg in 1 mL
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	25 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SORBITOL (UNII: 506T60A25R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:44224-3300-0	30 mL in 1 BOTTLE; 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	07/11/2003	

Part 6 of 7

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 7 of 7

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

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	01/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2012	

Labeler - Tender Corporation (064437304)

Registrant - Tender Corporation (064437304)

Establishment

Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(44224-2999)

Revised: 3/2017 Tender Corporation