DAVIMET™ Multivitamin

Multivitamin Tablet, Chewable

Supplement Facts

Serving Size: 1 Tablet
Servings per container: 30

	Amount Per Serving	%DV
Vitamin A (as Retinyl Acetate)	750 mcg RAE	83%
Vitamin C (as Sodium Ascorbate & Ascorbic Acid)	60 mg	67%
Vitamin D_3 (as Cholecalciferol)	10 mcg	50%
Vitamin E	10 mg	67%
Thiamin (as Thiamine Mononitrate)	1.05 mg	88%
Riboflavin	1.2 mg	92%
Niacin (as Niacinamide)	13.5 mg	84%
Vitamin B_6 (as Pyridoxine Hydrochloride)	1.05 mg	62%
Folate (as Folic Acid)	1667 mcg DFE (1000 mcg folic acid)	417%
Vitamin B_{12} (as Cyanocobalamin)	4.5 mcg	188%

NDC: 59088-0525-54

Other Ingredients:
Aspartame, Croscarmellose Sodium, Grape Flavor, Magnesium Stearate (vegetable source), Microcrystalline Cellulose, Stearic Acid (vegetable source), Sucrose, CI 45410 (Red 27 Lake), CI 42090 (FD&C Blue No. 1 Aluminum Lake).

Phenylketonurics: Phenylalanine 3.0 mg Per Tablet.

INDICATIONS AND USAGE
Davimet™ Multivitamin Chewable Tablets is indicated to provide significant amounts of Vitamins A, C, D, E, thiamine, riboflavin, niacin, vitamin B6, vitamin B12, and folate to supplement the diet, and to help assure that nutritional deficiencies of these vitamins will not develop.

Scan barcode for more information on Davimet™

FOLAMAX Mulitvitamin Dietary Supplement

Prescription Only

FolaMax

Multivitamin Dietary Supplement

Clinical dietary management of folate-deficient patients[1]

NDC: 71741-0086-30

Supplement Facts

Serving size 1 Tablet

Amount per Serving:		%Daily Value
Vitamin A (as Beta-Carotene)	300 mcg RAE	33%
Vitamin C (as Ascorbic Acid)	60 mg	67%
Vitamin D (as Cholecalciferol)	10 mcg	50%
Vitamin E (as dl-Alpha Tocopherol Acetate)	4.5 mg (10 IU)	33%
Vitamin B6 (as Pyridoxine HCl)	26 mg	1529%
Biotin	0.280 mg	933%
Folate	1.67 mg DFE	418%
(from Folic Acid)	0.67 mg DFE	**
(from 5-Methyl Tetrahydrofolate, Calcium Salt)	1 mg DFE	**
Vitamin B12 (as Cyanocobalamin)	0.013 mg	542%
Calcium (as Calcium Carbonate)	80 mg	6%
Magnesium (as Magnesium Oxide)	25 mg	6%
Ferrochel™ Iron (as Ferrous BisGly-cinate Chelate)	20 mg	111%
Iodine (as Potassium Iodide)	0.150 mg	100%

**Daily Value not established

INDICATIONS:
FolaMax is a prescription multivitamin/multimineral dietary supplement formulated for the clinical dietary management of suboptimal nutritional status in patients where advanced folate, vitamin B supplementation, and maintenance of good health is needed.

CONTRAINDICATIONS:
FolaMax is contraindicated in patients with a known hypersensitivity to any of the ingredients.

Scan barcode for more information on Folamax Multivitamin

LIDOGEL™ (Lidocaine 2.8% Gel)

NDC 59088-466-07 Rx Only

DERMACINRx

Lidogel™

Lidocaine HCl 2.8% Gel
Topical Anesthetic
NET WT. 3.5 oz. (100 g)

Specially formulated.
Use under the direction of a medical practitioner.

FOR EXTERNAL USE ONLY. NOT FOR OPHTHALMIC USE.

ACTIVE INGREDIENT: Lidocaine HCl 2.8%
INACTIVE INGREDIENTS: Aloe Barbadensis (Aloe Vera) Leaf Juice, Citric Acid, Hydroxyethylcellulose, Methylparaben, PEG-4, Propylene Glycol, Propylparaben, Purified Water.

INDICATIONS: For temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites and minor skin irritation.

KEEP THIS AND ALL MEDICATIONS OUT OF REACH OF CHILDREN.

USUAL ADULT DOSAGE: Apply two or three times daily or as directed by a physician.

See enclosed insert for complete product information.
Store at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature].
Protect from freezing.

For lot number and expiration date, see crimp of tube and/or carton.

List No. 46607IAA Rev. 38284

Manufactured in the USA by:
PureTek Corporation
Panorama City, CA 91402
For questions or information
call toll-free: **877-921-7873**

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LIDOGEL™ (Lidocaine HCl 2.8% Gel)

NET WT: 3.5 oz. (100g)

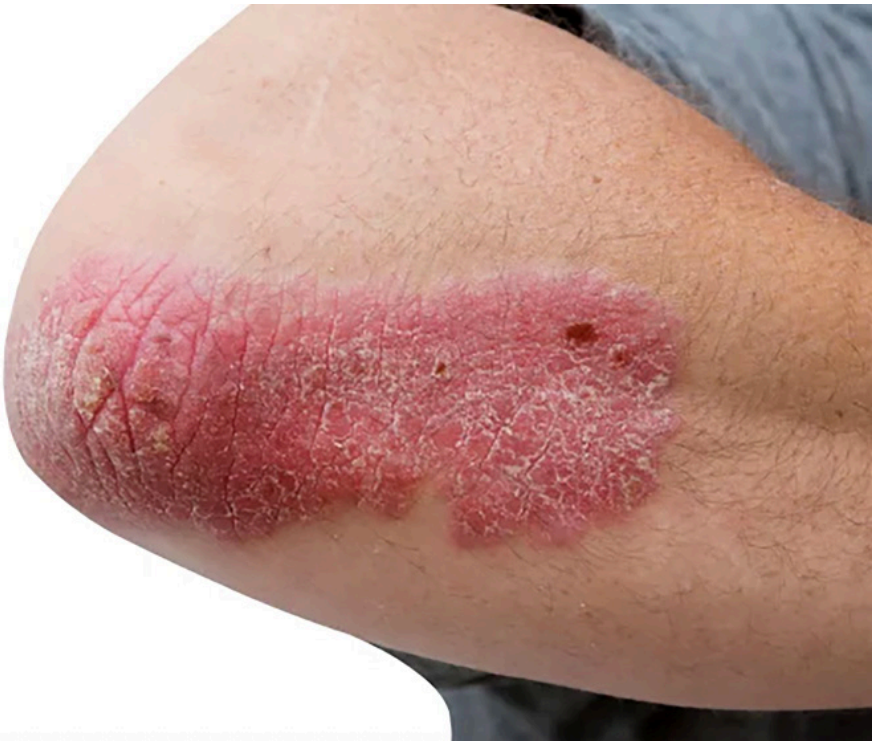
INGREDIENTS: Each gram of Lidogel™ 2.8% Gel contains Lidocaine HCl USP 28 mg.

INACTIVE INGREDIENTS INCLUDE:
Aloe Barbadensis (Aloe Vera) Leaf Juice, Citric Acid, Hydroxyethylcellulose, Methylparaben, PEG-4, Propylene Glycol, Propylparaben, Purified Water.

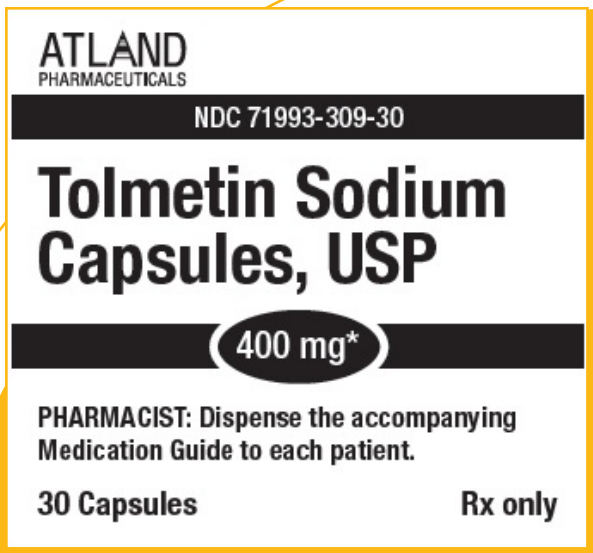
NDC: 59088-0466-07

INDICATIONS:
For temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritation.

Scan barcode for more information on Lidogel™



TOLMETIN SODIUM



Tolmetin Sodium Capsule

Unit Detail: 30 Capsules, 400 mg

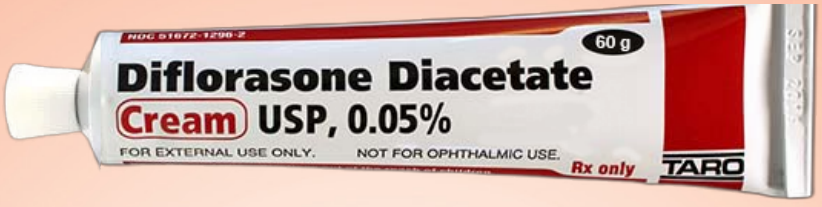
Description:
Each Tolmetin Sodium capsule for oral administration contains 492 mg of tolmetin sodium, USP as the dihydrate in an amount equivalent to 400 mg of Tolmetin.

Tolmetin tablets are indicated for the relief of signs and symptoms of rheumatoid arthritis and osteoarthritis. Tolmetin tablets are indicated in the treatment of acute flares and the long-term management of the chronic disease.

Tolmetin tablets are also indicated for treatment of juvenile rheumatoid arthritis. The safety and effectiveness of Tolmetin tablets have not been established in pediatric patients under 2 years of age.

Scan barcode for more information on Tolmetin Sodium

Diflorasone Diacetate, 0.05 % Cream



Diflorasone Diacetate, 0.05% Cream

Unit Detail: 60g

NDC: 51672-1296-03

Description:
DIFLORASONE reduces swelling, redness, itching, or rashes caused by skin conditions, such as eczema and psoriasis. It works by decreasing inflammation of the skin. It belongs to a group of medications called topical steroids.

Dosage And Administration
Diflorasone diacetate cream USP, 0.05% should be applied to the affected area twice daily.

Scan barcode for more information on Diflorasone Diacetate, 0.05 % Cream

Information for Patients
Patients using topical corticosteroids should receive the following information and instructions:

1. The medication is to be used as directed by the physician. It is for external use only.
Avoid contact with the eyes.

2.The medication should not be used for any disorder other than that for which it was prescribed.

3. The treated skin area should not be bandaged or otherwise covered or wrapped so as to be occlusive unless directed by the physician.

4.Patients should report to their physician any signs of local adverse reactions.

MULTITAM Multivitamin

Multitam Multivitamin

Each Caplet Contains:

Vitamin A (as retinyl acetate)	1500 mcg (5000 IU)
Vitamin C (as ascorbic acid)	120 mg
Vitamin D3 (as cholecalciferol)	20 mcg (800 IU)
Vitamin E (dl-alpha tocopheryl acetate)	30 mg (30 IU)
Thiamin (as thiamine mononitrate)	3 mg
Riboflavin (vitamin B2)	3.4 mg
Niacin (as niacinamide)	20 mg
Vitamin B6 (as pyridoxine hydrochloride)	20 mg
Folate (as folic acid)	1700 mcg DFE (1000 mcg folic acid)
Vitamin B12 (as cyanocobalamin)	8 mcg
Calcium (as calcium carbonate)	200 mg
Magnesium (as magnesium oxide)	200 mg
Zinc (as zinc oxide)	25 mg
Selenium (as selenium amino acid chelate)	55 mcg
Manganese (as manganese sulfate)	2.3 mg
Chromium (as chromium polynicotinate)	35 mcg
Molybdenum (as molybdenum amino acid chelate)	45 mcg

Other Ingredients:

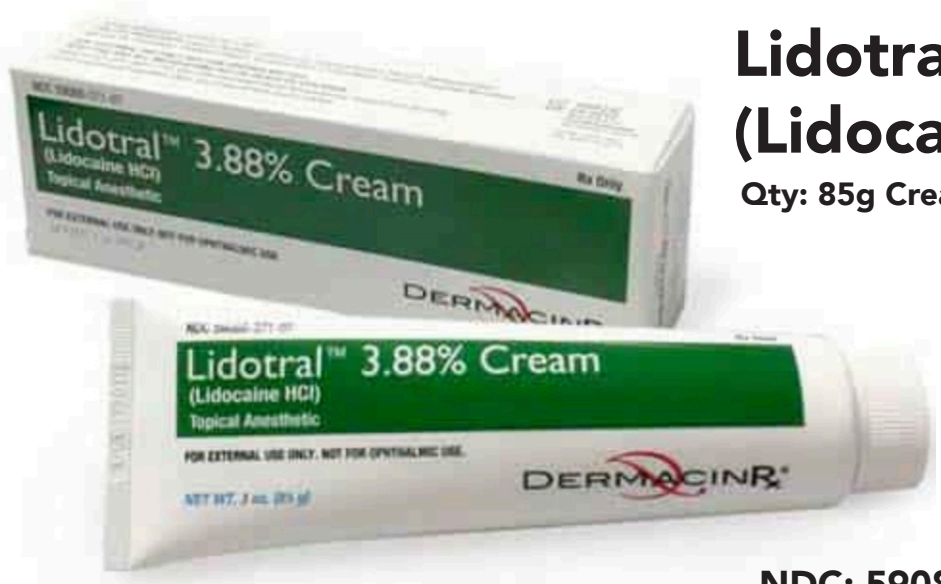
Organic cocoa powder, croscarmellose sodium, crospovidone, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid. Clear coating: (hydroxypropyl methylcellulose, PEG-8).

NDC: 59088-0525-54

Multitam™ is indicated to provide vitamin supplement to men and women. Folic acid is effective in the treatment of megaloblastic anemias due to a deficiency of folic acid (as may be seen in tropical or nontropical sprue) and in anemias of nutritional origin, pregnancy, infancy, or childhood.

Scan barcode for more information on Multitam Multivitamin

LIDOTRAL™ 3.88% Cream (Lidocaine HCl)



Lidotral™ 3.88% Cream (Lidocaine HCl)

Qty: 85g Cream

NDC: 59088-0525-54

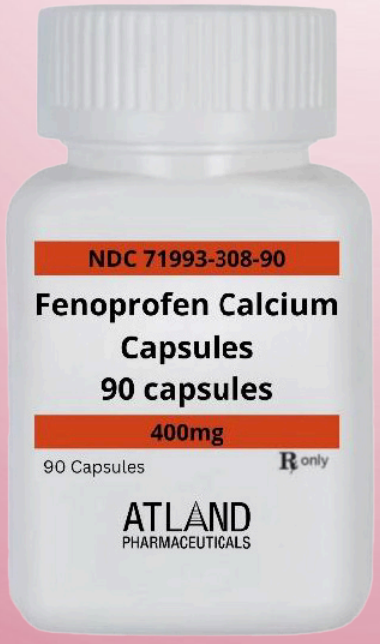
Dosage:
Apply a thin film to the affected area two or three times daily or as directed by a physician.

Indication:
For temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites, pain associated with haemorrhoid and anus fisher and minor skin irritation.

Contraindication:
Tuberculous or fungal lesions of skin vaccinia, varicella and acute herpes simplex and in persons who have shown hypersensitivity to any of its components. Lidocaine is contraindicated in patients with a known history of hypersensitivity to local anesthetics of the amide type.

Scan barcode for more information on Lidotral™

FENOPROFEN 400 mg



Fenoprofen 400 mg

INDICATIONS AND USAGE:
Fenoprofen calcium is a nonsteroidal anti-inflammatory drug indicated for:
- Relief of mild to moderate pain in adults.
- Relief of the signs and symptoms of rheumatoid arthritis.
- Relief of the signs and symptoms of osteoarthritis.

NDC: 71993-0308-90

DOSAGE:
- Use the lowest effective dosage for shortest duration consistent with individual patient treatment goals.
- Analgesia: For the treatment of mild to moderate pain, the recommended dosage is 200 mg given orally every 4 to 6 hours, as needed.
- Rheumatoid Arthritis and Osteoarthritis: For the relief of signs and symptoms of rheumatoid arthritis or osteoarthritis the recommended dose is 400 to 600 mg given orally, 3 or 4 times a day. The dose should be tailored to the needs of the patient and may be increased or decreased depending on the severity of the symptoms. Total daily dosage should not exceed 3,200 mg.

CONTRAINDICATIONS:
- Known hypersensitivity to Fenoprofen or any components of the drug product.
- History of asthma, urticaria, or other allergic-type reactions after taking aspirin or other NSAIDs
- In the setting of CABG surgery.

WARNINGS AND PRECAUTIONS:
- Hepatotoxicity: Inform patients of warning signs and symptoms of hepatotoxicity. Discontinue if abnormal liver tests persist or worsen or if clinical signs and symptoms of liver disease develop.
- Hypertension: Patients taking some antihypertensive medications may have impaired response to these therapies when taking NSAIDs. Monitor blood pressure
- Heart Failure and Edema: Avoid use of Fenoprofen calcium in patients with severe heart failure unless benefits are expected to outweigh risk of worsening heart failure.
- Renal Toxicity: Monitor renal function in patients with renal or hepatic impairment, heart failure, dehydration, or hypovolemia. Avoid use of Fenoprofen calcium in patients with advanced renal disease unless benefits are expected to outweigh risk of worsening renal function.
- Anaphylactic Reactions: Seek emergency help if an anaphylactic reaction occurs.
- Exacerbation of Asthma Related to Aspirin Sensitivity: Fenoprofen calcium is contraindicated in patients with aspirin-sensitive asthma. Monitor patients with preexisting asthma (without aspirin sensitivity).
- Serious Skin Reactions: Discontinue Fenoprofen calcium at first appearance of skin rash or other signs of hypersensitivity.
- Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS): Discontinue and evaluate clinically.
- Fetal Toxicity: Limit use of NSAIDs, including Fenoprofen calcium, between about 20 to 30 weeks in pregnancy due to the risk of oligohydramnios/fetal renal dysfunction. Avoid use of NSAIDs in women at about 30 weeks gestation and later in pregnancy due to the risks of oligohydramnios/fetal renal dysfunction and premature closure of the fetal ductus arteriosus.
- Hematologic Toxicity: Monitor hemoglobin or hematocrit in patients with any signs or symptoms of anemia.

Scan barcode for more information on Fenoprofen.