

**NAPROXEN - naproxen tablet, delayed release**  
**H.J. Harkins Company, Inc.**

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**NAPROXEN DELAYED-RELEASE TABLETS USP**

**1005**

**1006**

**Rx only**

**Cardiovascular Risk**

- NSAIDs\* may cause an increased risk of serious cardiovascular thrombotic events, myocardial infarction, and stroke, which can be fatal. This risk may increase with duration of use. Patients with cardiovascular disease or risk factors for cardiovascular disease may be at greater risk (see **WARNINGS**).
- Naproxen delayed-release tablets are contraindicated for the treatment of peri-operative pain in the setting of coronary artery bypass graft (CABG) surgery (see **WARNINGS**).

**Gastrointestinal Risk**

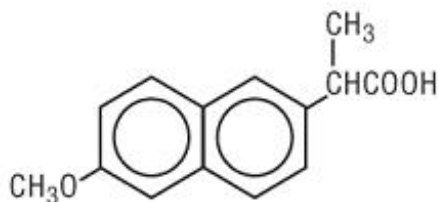
- NSAIDs cause an increased risk of serious gastrointestinal adverse events including bleeding, ulceration, and perforation of the stomach or intestines, which can be fatal. These events can occur at any time during use and without warning symptoms. Elderly patients are at greater risk for serious gastrointestinal events (see **WARNINGS**).

\* Throughout this package insert, the term NSAID refers to a non-aspirin non-steroidal anti-inflammatory drug.

**DESCRIPTION**

Naproxen is a propionic acid derivative related to the arylacetic acid group of non-steroidal anti-inflammatory drugs.

The chemical name for naproxen is (+)-6-methoxy- $\alpha$ -methyl-2-naphthaleneacetic acid. It has the following structural formula:



$C_{14}H_{14}O_3$  M.W. 230.26

Naproxen is a practically odorless, white to off-white crystalline substance. It is lipid-soluble, practically insoluble in water at low pH and freely soluble in water at high pH. The octanol/water partition coefficient of naproxen at pH 7.4 is 1.6 to 1.8.

Naproxen delayed-release tablets USP are available as enteric-coated, white to off-white tablets containing 375 mg or 500 mg of naproxen for oral administration. The inactive ingredients are: corn starch, croscarmellose sodium, FD&C Blue No. 2, magnesium stearate, methacrylic acid copolymer-dispersion, povidone, talc, titanium dioxide, and triethyl citrate. The dissolution of this enteric-coated

naproxen tablet is pH dependent with rapid dissolution above pH 6. There is no dissolution below pH 4.

## CLINICAL PHARMACOLOGY

### Pharmacodynamics

Naproxen is a non-steroidal anti-inflammatory drug (NSAID) with analgesic and antipyretic properties. The sodium salt of naproxen has been developed as a more rapidly absorbed formulation of naproxen for use as an analgesic. The mechanism of action of the naproxen anion, like that of other NSAIDs, is not completely understood but may be related to prostaglandin synthetase inhibition.

### Pharmacokinetics

Naproxen is rapidly and completely absorbed from the gastrointestinal tract with an *in vivo* bioavailability of 95%. The different dosage forms of naproxen are bioequivalent in terms of extent of absorption (AUC) and peak concentration ( $C_{max}$ ); however, the products do differ in their pattern of absorption. These differences between naproxen products are related to both the chemical form of naproxen used and its formulation. Even with the observed differences in pattern of absorption, the elimination half-life of naproxen is unchanged across products ranging from 12 to 17 hours. Steady-state levels of naproxen are reached in 4 to 5 days, and the degree of naproxen accumulation is consistent with this half-life. This suggests that the differences in pattern of release play only a negligible role in the attainment of steady-state plasma levels.

#### Absorption

##### *Delayed release*

Naproxen delayed-release tablets are designed with a pH-sensitive coating to provide a barrier to disintegration in the acidic environment of the stomach and to lose integrity in the more neutral environment of the small intestine. The enteric polymer coating selected for naproxen delayed-release tablets dissolves above pH 6. When naproxen delayed-release tablets were given to fasted subjects, peak plasma levels were attained about 4 to 6 hours following the first dose (range: 2 to 12 hours). An *in vivo* study in man using radiolabeled naproxen delayed-release tablets demonstrated that naproxen delayed-release tablets dissolve primarily in the small intestine rather than in the stomach, so the absorption of the drug is delayed until the stomach is emptied.

When naproxen delayed-release tablets and naproxen tablets were given to fasted subjects (n = 24) in a crossover study following 1 week of dosing, differences in time to peak plasma levels ( $T_{max}$ ) were observed, but there were no differences in total absorption as measured by  $C_{max}$  and AUC:

|   | <b>Naproxen<br/>Delayed-Release<br/>Tablets*<br/>500 mg bid</b> | <b>Naproxen<br/>Tablets*<br/>500 mg bid</b> |
|---|---|---|
| $C_{max}$ (mcg/mL)                      | 94.9 (18%)  | 97.4 (13%)                                  |
| $T_{max}$ (hours)                       | 4 (39%)   | 1.9 (61%)                                   |
| AUC <sub>0-12 hr</sub> (mcg•hr/mL)      | 845 (20%)   | 767 (15%)                                   |
| * Mean value (coefficient of variation) |   |   |

##### *Antacid effects*

When naproxen delayed-release tablets were given as a single dose with antacid (54 mEq buffering capacity), the peak plasma levels of naproxen were unchanged, but the time to peak was reduced (mean  $T_{max}$  fasted 5.6 hours, mean  $T_{max}$  with antacid 5 hours), although not significantly.

### *Food effects*

When naproxen delayed-release tablets were given as a single dose with food, peak plasma levels in most subjects were achieved in about 12 hours (range: 4 to 24 hours). Residence time in the small intestine until disintegration was independent of food intake. The presence of food prolonged the time the tablets remained in the stomach, time to first detectable serum naproxen levels, and time to maximal naproxen levels ( $T_{max}$ ), but did not affect peak naproxen levels ( $C_{max}$ ).

### *Distribution*

Naproxen has a volume of distribution of 0.16 L/kg. At therapeutic levels naproxen is greater than 99% albumin-bound. At doses of naproxen greater than 500 mg/day there is less than proportional increase in plasma levels due to an increase in clearance caused by saturation of plasma protein binding at higher doses (average trough  $C_{ss}$  36.5, 49.2, and 56.4 mg/L with 500, 1000 and 1500 mg daily doses of naproxen, respectively). The naproxen anion has been found in the milk of lactating women at a concentration equivalent to approximately 1% of maximum naproxen concentration in plasma (see **PRECAUTIONS, Nursing Mothers**).

### *Metabolism*

Naproxen is extensively metabolized in the liver to 6-*O*-desmethyl naproxen, and both parent and metabolites do not induce metabolizing enzymes. Both naproxen and 6-*O*-desmethyl naproxen are further metabolized to their respective acylglucuronide conjugated metabolites.

### *Excretion*

The clearance of naproxen is 0.13 mL/min/kg. Approximately 95% of the naproxen from any dose is excreted in the urine, primarily as naproxen (< 1%), 6-*O*-desmethyl naproxen (< 1%) or their conjugates (66% to 92%). The plasma half-life of the naproxen anion in humans ranges from 12 to 17 hours. The corresponding half-lives of both naproxen's metabolites and conjugates are shorter than 12 hours, and their rates of excretion have been found to coincide closely with the rate of naproxen disappearance from the plasma. Small amounts, 3% or less of the administered dose, are excreted in the feces. In patients with renal failure metabolites may accumulate (see **WARNINGS, Renal Effects**).

### *Special Populations*

#### *Pediatric patients*

In pediatric patients aged 5 to 16 years with arthritis, plasma naproxen levels following a 5 mg/kg single dose of naproxen suspension were found to be similar to those found in normal adults following a 500 mg dose. The terminal half-life appears to be similar in pediatric and adult patients. Pharmacokinetic studies of naproxen were not performed in pediatric patients younger than 5 years of age. Pharmacokinetic parameters appear to be similar following administration of naproxen suspension or tablets in pediatric patients. Naproxen delayed-release tablets have not been studied in subjects under the age of 18.

#### *Geriatric patients*

Studies indicate that although total plasma concentration of naproxen is unchanged, the unbound plasma fraction of naproxen is increased in the elderly, although the unbound fraction is < 1% of the total naproxen concentration. Unbound trough naproxen concentrations in elderly subjects have been reported to range from 0.12% to 0.19% of total naproxen concentration, compared with 0.05% to 0.075% in younger subjects. The clinical significance of this finding is unclear, although it is possible that the increase in free naproxen concentration could be associated with an increase in the rate of adverse events per a given dosage in some elderly patients.

### *Race*

Pharmacokinetic differences due to race have not been studied.

#### *Hepatic insufficiency*

Naproxen pharmacokinetics have not been determined in subjects with hepatic insufficiency.

#### *Renal insufficiency*

Naproxen pharmacokinetics have not been determined in subjects with renal insufficiency. Given that naproxen, its metabolites and conjugates are primarily excreted by the kidney, the potential exists for naproxen metabolites to accumulate in the presence of renal insufficiency. Elimination of naproxen is decreased in patients with severe renal impairment. Naproxen-containing products are not recommended for use in patients with moderate to severe and severe renal impairment (creatinine clearance < 30 mL/min) (see **WARNINGS, Renal Effects**).

## **CLINICAL STUDIES**

### **General Information**

Naproxen has been studied in patients with rheumatoid arthritis, osteoarthritis, juvenile arthritis, ankylosing spondylitis, tendonitis and bursitis, and acute gout. Improvement in patients treated for rheumatoid arthritis was demonstrated by a reduction in joint swelling, a reduction in duration of morning stiffness, a reduction in disease activity as assessed by both the investigator and patient, and by increased mobility as demonstrated by a reduction in walking time. Generally, response to naproxen has not been found to be dependent on age, sex, severity or duration of rheumatoid arthritis.

In patients with osteoarthritis, the therapeutic action of naproxen has been shown by a reduction in joint pain or tenderness, an increase in range of motion in knee joints, increased mobility as demonstrated by a reduction in walking time, and improvement in capacity to perform activities of daily living impaired by the disease.

In a clinical trial comparing standard formulations of naproxen 375 mg bid (750 mg a day) vs 750 mg bid (1500 mg/day), 9 patients in the 750 mg group terminated prematurely because of adverse events. Nineteen patients in the 1500 mg group terminated prematurely because of adverse events. Most of these adverse events were gastrointestinal events.

In clinical studies in patients with rheumatoid arthritis, osteoarthritis, and juvenile arthritis, naproxen has been shown to be comparable to aspirin and indomethacin in controlling the aforementioned measures of disease activity, but the frequency and severity of the milder gastrointestinal adverse effects (nausea, dyspepsia, heartburn) and nervous system adverse effects (tinnitus, dizziness, lightheadedness) were less in naproxen-treated patients than in those treated with aspirin or indomethacin.

In patients with ankylosing spondylitis, naproxen has been shown to decrease night pain, morning stiffness and pain at rest. In double-blind studies the drug was shown to be as effective as aspirin, but with fewer side effects.

In patients with acute gout, a favorable response to naproxen was shown by significant clearing of inflammatory changes (e.g., decrease in swelling, heat) within 24 to 48 hours, as well as by relief of pain and tenderness.

Naproxen has been studied in patients with mild to moderate pain secondary to postoperative, orthopedic, postpartum episiotomy and uterine contraction pain and dysmenorrhea. Onset of pain relief can begin within 1 hour in patients taking naproxen and within 30 minutes in patients taking naproxen sodium. Analgesic effect was shown by such measures as reduction of pain intensity scores, increase in pain relief scores, decrease in numbers of patients requiring additional analgesic medication, and delay in time to remedication. The analgesic effect has been found to last for up to 12 hours.

Naproxen may be used safely in combination with gold salts and/or corticosteroids; however, in

controlled clinical trials, when added to the regimen of patients receiving corticosteroids, it did not appear to cause greater improvement over that seen with corticosteroids alone. Whether naproxen has a "steroid-sparing" effect has not been adequately studied. When added to the regimen of patients receiving gold salts, naproxen did result in greater improvement. Its use in combination with salicylates is not recommended because there is evidence that aspirin increases the rate of excretion of naproxen and data are inadequate to demonstrate that naproxen and aspirin produce greater improvement over that achieved with aspirin alone. In addition, as with other NSAIDs, the combination may result in higher frequency of adverse events than demonstrated for either product alone.

In  $^{51}\text{Cr}$  blood loss and gastroscopy studies with normal volunteers, daily administration of 1000 mg of naproxen has been demonstrated to cause statistically significantly less gastric bleeding and erosion than 3250 mg of aspirin.

Three 6 week, double-blind, multicenter studies with naproxen delayed-release tablets (375 or 500 mg bid, n = 385) and naproxen immediate-release tablets (375 or 500 mg bid, n = 279) were conducted comparing naproxen delayed-release tablets with naproxen immediate-release tablets, including 355 rheumatoid arthritis and osteoarthritis patients who had a recent history of NSAID-related GI symptoms. These studies indicated that naproxen delayed-release tablets and naproxen immediate-release tablets showed no significant differences in efficacy or safety and had similar prevalence of minor GI complaints. Individual patients, however, may find one formulation preferable to the other.

Five hundred and fifty-three patients received naproxen delayed-release tablets during long-term open-label trials (mean length of treatment was 159 days). The rates for clinically-diagnosed peptic ulcers and GI bleeds were similar to what has been historically reported for long-term NSAID use.

### **Geriatric Patients**

The hepatic and renal tolerability of long-term naproxen administration was studied in two double-blind clinical trials involving 586 patients. Of the patients studied, 98 patients were age 65 and older and 10 of the 98 patients were age 75 and older. Naproxen was administered at doses of 375 mg twice daily or 750 mg twice daily for up to 6 months. Transient abnormalities of laboratory tests assessing hepatic and renal function were noted in some patients, although there were no differences noted in the occurrence of abnormal values among different age groups.

## **INDICATIONS AND USAGE**

Carefully consider the potential benefits and risks of naproxen delayed-release tablets and other treatment options before deciding to use naproxen delayed-release tablets. Use the lowest effective dose for the shortest duration consistent with individual patient treatment goals (see **WARNINGS**).

Naproxen delayed-release tablets are indicated:

- For the relief of the signs and symptoms of rheumatoid arthritis
- For the relief of the signs and symptoms of osteoarthritis
- For the relief of the signs and symptoms of ankylosing spondylitis
- For the relief of the signs and symptoms of juvenile arthritis

Naproxen as naproxen suspension is recommended for juvenile rheumatoid arthritis in order to obtain the maximum dosage flexibility based on the patient's weight.

Naproxen delayed-release tablets are not recommended for initial treatment of acute pain because the absorption of naproxen is delayed compared to absorption from other naproxen-containing products (see **CLINICAL PHARMACOLOGY** and **DOSAGE AND ADMINISTRATION**).

## **CONTRAINDICATIONS**

Naproxen delayed-release tablets are contraindicated in patients with known hypersensitivity to

naproxen and naproxen sodium.

Naproxen delayed-release tablets should not be given to patients who have experienced asthma, urticaria, or allergic-type reactions after taking aspirin or other NSAIDs. Severe, rarely fatal, anaphylactoid-like reactions to NSAIDs have been reported in such patients (see **WARNINGS, Anaphylactoid Reactions** and **PRECAUTIONS, Preexisting Asthma**).

Naproxen delayed-release tablets are contraindicated for the treatment of peri-operative pain in the setting of coronary artery bypass graft (CABG) surgery (see **WARNINGS**).

## **WARNINGS**

### **CARDIOVASCULAR EFFECTS**

#### Cardiovascular Thrombotic Events

Clinical trials of several COX-2 selective and nonselective NSAIDs of up to three years duration have shown an increased risk of serious cardiovascular (CV) thrombotic events, myocardial infarction, and stroke, which can be fatal. All NSAIDs, both COX-2 selective and nonselective, may have a similar risk. Patients with known CV disease or risk factors for CV disease may be at greater risk. To minimize the potential risk for an adverse CV event in patients treated with an NSAID, the lowest effective dose should be used for the shortest duration possible. Physicians and patients should remain alert for the development of such events, even in the absence of previous CV symptoms. Patients should be informed about the signs and/or symptoms of serious CV events and the steps to take if they occur.

There is no consistent evidence that concurrent use of aspirin mitigates the increased risk of serious CV thrombotic events associated with NSAID use. The concurrent use of aspirin and an NSAID does increase the risk of serious GI events (see **Gastrointestinal Effects – Risk of Ulceration, Bleeding, and Perforation**).

Two large, controlled clinical trials of a COX-2 selective NSAID for the treatment of pain in the first 10 to 14 days following CABG surgery found an increased incidence of myocardial infarction and stroke (see **CONTRAINDICATIONS**).

#### Hypertension

NSAIDs, including naproxen delayed-release tablets, can lead to onset of new hypertension or worsening of preexisting hypertension, either of which may contribute to the increased incidence of CV events. Patients taking thiazides or loop diuretics may have impaired response to these therapies when taking NSAIDs. NSAIDs, including naproxen delayed-release tablets, should be used with caution in patients with hypertension. Blood pressure (BP) should be monitored closely during the initiation of NSAID treatment and throughout the course of therapy.

#### Congestive Heart Failure and Edema

Fluid retention, edema, and peripheral edema have been observed in some patients taking NSAIDs. Naproxen delayed-release tablets should be used with caution in patients with fluid retention, hypertension, or heart failure.

### **Gastrointestinal Effects – Risk of Ulceration, Bleeding, and Perforation**

NSAIDs, including naproxen delayed-release tablets, can cause serious gastrointestinal (GI) adverse events including inflammation, bleeding, ulceration, and perforation of the stomach, small intestine, or large intestine, which can be fatal.

These serious adverse events can occur at any time, with or without warning symptoms, in patients treated with NSAIDs. Only one in five patients, who develop a serious upper GI adverse event on NSAID therapy, is symptomatic. Upper GI ulcers, gross bleeding, or perforation caused by NSAIDs

occur in approximately 1% of patients treated for 3 to 6 months, and in about 2 to 4% of patients treated for one year. These trends continue with longer duration of use, increasing the likelihood of developing a serious GI event at some time during the course of therapy. However, even short-term therapy is not without risk. The utility of periodic laboratory monitoring has not been demonstrated, nor has it been adequately assessed. Only 1 in 5 patients who develop a serious upper GI adverse event on NSAID therapy is symptomatic.

NSAIDs should be prescribed with extreme caution in those with a prior history of ulcer disease or gastrointestinal bleeding. Patients with a prior history of peptic ulcer disease and/or gastrointestinal bleeding who use NSAIDs have a greater than 10 fold increased risk for developing a GI bleed compared to patients with neither of these risk factors. Other factors that increase the risk for GI bleeding in patients treated with NSAIDs include concomitant use of oral corticosteroids or anticoagulants, longer duration of NSAID therapy, smoking, use of alcohol, older age, and poor general health status. Most spontaneous reports of fatal GI events are in elderly or debilitated patients and therefore, special care should be taken in treating this population. To minimize the potential risk for an adverse GI event in patients treated with an NSAID, the lowest effective dose should be used for the shortest possible duration. Patients and physicians should remain alert for signs and symptoms of GI ulceration and bleeding during NSAID therapy and promptly initiate additional evaluation and treatment if a serious GI adverse event is suspected. This should include discontinuation of the NSAID until a serious GI adverse event is ruled out. For high risk patients, alternate therapies that do not involve NSAIDs should be considered.

Epidemiological studies, both of the case-control and cohort design, have demonstrated an association between use of psychotropic drugs that interfere with serotonin reuptake and the occurrence of upper gastrointestinal bleeding. In two studies, concurrent use of an NSAID or aspirin potentiated the risk of bleeding (see **PRECAUTIONS, Drug Interactions**). Although these studies focused on upper gastrointestinal bleeding, there is reason to believe that bleeding at other sites may be similarly potentiated.

NSAIDs should be given with care to patients with a history of inflammatory bowel disease (ulcerative colitis, Crohn's disease) as their condition may be exacerbated.

### **Renal Effects**

Long-term administration of NSAIDs has resulted in renal papillary necrosis and other renal injury. Renal toxicity has also been seen in patients in whom renal prostaglandins have a compensatory role in the maintenance of renal perfusion. In these patients, administration of a non-steroidal anti-inflammatory drug may cause a dose-dependent reduction in prostaglandin formation and, secondarily, in renal blood flow, which may precipitate overt renal decompensation. Patients at greatest risk of this reaction are those with impaired renal function, hypovolemia, heart failure, liver dysfunction, salt depletion, those taking diuretics and ACE- inhibitors, and the elderly. Discontinuation of non-steroidal anti-inflammatory drug therapy is usually followed by recovery to the pretreatment state (see **WARNINGS, Advanced Renal Disease**).

### **Advanced Renal Disease**

No information is available from controlled clinical studies regarding the use of naproxen delayed-release tablets in patients with advanced renal disease. Therefore, treatment with naproxen delayed-release tablets is not recommended in these patients with advanced renal disease. If naproxen delayed-release tablet therapy must be initiated, close monitoring of the patient's renal function is advisable.

### **Anaphylactoid Reactions**

As with other NSAIDs, anaphylactoid reactions may occur in patients without known prior exposure to naproxen delayed-release tablets. Naproxen delayed-release tablets should not be given to patients with the aspirin triad. This symptom complex typically occurs in asthmatic patients who experience rhinitis with or without nasal polyps, or who exhibit severe, potentially fatal bronchospasm after taking aspirin

or other NSAIDs (see **CONTRAINDICATIONS** and **PRECAUTIONS, Preexisting Asthma**). Emergency help should be sought in cases where an anaphylactoid reaction occurs. Anaphylactoid reactions, like anaphylaxis, may have a fatal outcome.

### **Skin Reactions**

NSAIDs, including naproxen delayed-release tablets, can cause serious skin adverse events such as exfoliative dermatitis, Stevens-Johnson syndrome (SJS), and toxic epidermal necrolysis (TEN), which can be fatal. These serious events may occur without warning. Patients should be informed about the signs and symptoms of serious skin manifestations and use of the drug should be discontinued at the first appearance of skin rash or any other sign of hypersensitivity.

### **Pregnancy**

In late pregnancy, as with other NSAIDs, naproxen delayed-release tablets should be avoided because they may cause premature closure of the ductus arteriosus.

## **PRECAUTIONS**

### **General**

**Naproxen-containing products such as naproxen tablets, naproxen delayed-release tablets, naproxen sodium tablets, naproxen suspension, and other naproxen products should not be used concomitantly since they all circulate in the plasma as the naproxen anion.**

Naproxen delayed-release tablets cannot be expected to substitute for corticosteroids or to treat corticosteroid insufficiency. Abrupt discontinuation of corticosteroids may lead to disease exacerbation. Patients on prolonged corticosteroid therapy should have their therapy tapered slowly if a decision is made to discontinue corticosteroids and the patient should be observed closely for any evidence of adverse effects, including adrenal insufficiency and exacerbation of symptoms of arthritis.

Patients with initial hemoglobin values of 10 g or less who are to receive long-term therapy should have hemoglobin values determined periodically.

The pharmacological activity of naproxen delayed-release tablets in reducing fever and inflammation may diminish the utility of these diagnostic signs in detecting complications of presumed noninfectious, noninflammatory painful conditions.

Because of adverse eye findings in animal studies with drugs of this class, it is recommended that ophthalmic studies be carried out if any change or disturbance in vision occurs.

### **Hepatic Effects**

Borderline elevations of one or more liver tests may occur in up to 15% of patients taking NSAIDs including naproxen delayed-release tablets. Hepatic abnormalities may be the result of hypersensitivity rather than direct toxicity. These laboratory abnormalities may progress, may remain essentially unchanged, or may be transient with continued therapy. The SGPT (ALT) test is probably the most sensitive indicator of liver dysfunction. Notable elevations of ALT or AST (approximately three or more times the upper limit of normal) have been reported in approximately 1% of patients in clinical trials with NSAIDs. In addition, rare cases of severe hepatic reactions, including jaundice and fatal fulminant hepatitis, liver necrosis and hepatic failure, some of them with fatal outcomes have been reported.

A patient with symptoms and/or signs suggesting liver dysfunction, or in whom an abnormal liver test has occurred, should be evaluated for evidence of the development of a more severe hepatic reaction while on therapy with naproxen delayed-release tablets.

If clinical signs and symptoms consistent with liver disease develop, or if systemic manifestations occur



(e.g., eosinophilia, rash, etc.), naproxen delayed-release tablets should be discontinued.

Chronic alcoholic liver disease and probably other diseases with decreased or abnormal plasma proteins (albumin) reduce the total plasma concentration of naproxen, but the plasma concentration of unbound naproxen is increased. Caution is advised when high doses are required and some adjustment of dosage may be required in these patients. It is prudent to use the lowest effective dose.

### **Hematological Effects**

Anemia is sometimes seen in patients receiving NSAIDs, including naproxen delayed-release tablets. This may be due to fluid retention, occult or gross GI blood loss, or an incompletely described effect upon erythropoiesis. Patients on long-term treatment with NSAIDs, including naproxen delayed-release tablets, should have their hemoglobin or hematocrit checked if they exhibit any signs or symptoms of anemia.

NSAIDs inhibit platelet aggregation and have been shown to prolong bleeding time in some patients. Unlike aspirin, their effect on platelet function is quantitatively less, of shorter duration, and reversible. Patients receiving naproxen delayed-release tablets who may be adversely affected by alterations in platelet function, such as those with coagulation disorders or patients receiving anticoagulants, should be carefully monitored.

### **Preexisting Asthma**

Patients with asthma may have aspirin-sensitive asthma. The use of aspirin in patients with aspirin-sensitive asthma has been associated with severe bronchospasm, which can be fatal. Since cross reactivity, including bronchospasm, between aspirin and other non-steroidal anti-inflammatory drugs has been reported in such aspirin-sensitive patients, naproxen delayed-release tablets should not be administered to patients with this form of aspirin sensitivity and should be used with caution in patients with preexisting asthma.

### **Information for Patients**

**Patients should be informed of the following information before initiating therapy with an NSAID and periodically during the course of ongoing therapy. Patients should also be encouraged to read the NSAID Medication Guide that accompanies each prescription dispensed.**

1. Naproxen delayed-release tablets, like other NSAIDs, may cause serious CV side effects, such as MI or stroke, which may result in hospitalization and even death. Although serious CV events can occur without warning symptoms, patients should be alert for the signs and symptoms of chest pain, shortness of breath, weakness, slurring of speech, and should ask for medical advice when observing any indicative signs or symptoms. Patients should be apprised of the importance of this follow-up (see **WARNINGS, CARDIOVASCULAR EFFECTS**).
2. Naproxen delayed-release tablets, like other NSAIDs, can cause GI discomfort and, rarely, serious GI side effects, such as ulcers and bleeding, which may result in hospitalization and even death. Although serious GI tract ulcerations and bleeding can occur without warning symptoms, patients should be alert for the signs and symptoms of ulcerations and bleeding, and should ask for medical advice when observing any indicative sign or symptoms including epigastric pain, dyspepsia, melena, and hematemesis. Patients should be apprised of the importance of this follow-up (see **WARNINGS, Gastrointestinal Effects – Risk of Ulceration, Bleeding, and Perforation**).
3. Naproxen delayed-release tablets, like other NSAIDs, can cause serious skin side effects such as exfoliative dermatitis, SJS, and TEN, which may result in hospitalization and even death. Although serious skin reactions may occur without warning, patients should be alert for the signs and symptoms of skin rash and blisters, fever, or other signs of hypersensitivity such as itching, and should ask for medical advice when observing any indicative signs or symptoms. Patients should be advised to stop the drug immediately if they develop any type of rash and contact their physicians as soon as possible.
4. Patients should promptly report signs or symptoms of unexplained weight gain or edema to their

physicians.

5. Patients should be informed of the warning signs and symptoms of hepatotoxicity (e.g., nausea, fatigue, lethargy, pruritus, jaundice, right upper quadrant tenderness, and "flu-like" symptoms). If these occur, patients should be instructed to stop therapy and seek immediate medical therapy.
6. Patients should be informed of the signs of an anaphylactoid reaction (e.g., difficulty breathing, swelling of the face or throat). If these occur, patients should be instructed to seek immediate emergency help (see **WARNINGS**).
7. In late pregnancy, as with other NSAIDs, naproxen delayed-release tablets should be avoided because they may cause premature closure of the ductus arteriosus.
8. Caution should be exercised by patients whose activities require alertness if they experience drowsiness, dizziness, vertigo or depression during therapy with naproxen.

## Laboratory Tests

Because serious GI tract ulcerations and bleeding can occur without warning symptoms, physicians should monitor for signs or symptoms of GI bleeding. Patients on long-term treatment with NSAIDs should have their CBC and a chemistry profile checked periodically. If clinical signs and symptoms consistent with liver or renal disease develop, systemic manifestations occur (e.g., eosinophilia, rash, etc.) or if abnormal liver tests persist or worsen, naproxen delayed-release tablets should be discontinued.

## Drug Interactions

### ACE-Inhibitors

Reports suggest that NSAIDs may diminish the antihypertensive effect of ACE-inhibitors. This interaction should be given consideration in patients taking NSAIDs concomitantly with ACE-inhibitors.

### Antacids and Sucralfate

Concomitant administration of some antacids (magnesium oxide or aluminum hydroxide) and sucralfate can delay the absorption of naproxen.

### Aspirin

When naproxen delayed-release tablets are administered with aspirin, its protein binding is reduced, although the clearance of free naproxen is not altered. The clinical significance of this interaction is not known; however, as with other NSAIDs, concomitant administration of naproxen delayed-release tablets and aspirin is not generally recommended because of the potential of increased adverse effects.

### Cholestyramine

As with other NSAIDs, concomitant administration of cholestyramine can delay the absorption of naproxen.

### Diuretics

Clinical studies, as well as postmarketing observations, have shown that naproxen delayed-release tablets can reduce the natriuretic effect of furosemide and thiazides in some patients. This response has been attributed to inhibition of renal prostaglandin synthesis. During concomitant therapy with NSAIDs, the patient should be observed closely for signs of renal failure (see **WARNINGS, Renal Effects**), as well as to assure diuretic efficacy.

### Lithium

NSAIDs have produced an elevation of plasma lithium levels and a reduction in renal lithium clearance. The mean minimum lithium concentration increased 15% and the renal clearance was decreased by approximately 20%. These effects have been attributed to inhibition of renal prostaglandin synthesis by

the NSAID. Thus, when NSAIDs and lithium are administered concurrently, subjects should be observed carefully for signs of lithium toxicity.

### Methotrexate

NSAIDs have been reported to competitively inhibit methotrexate accumulation in rabbit kidney slices. Naproxen, naproxen sodium, and other non-steroidal anti-inflammatory drugs have been reported to reduce the tubular secretion of methotrexate in an animal model. This may indicate that they could enhance the toxicity of methotrexate. Caution should be used when NSAIDs are administered concomitantly with methotrexate.

### Warfarin

The effects of warfarin and NSAIDs on GI bleeding are synergistic, such that users of both drugs together have a risk of serious GI bleeding higher than users of either drug alone. No significant interactions have been observed in clinical studies with naproxen and coumarin-type anticoagulants. However, caution is advised since interactions have been seen with other non-steroidal agents of this class. The free fraction of warfarin may increase substantially in some subjects and naproxen interferes with platelet function.

### Selective Serotonin Reuptake Inhibitors (SSRIs)

There is an increased risk of gastrointestinal bleeding when selective serotonin reuptake inhibitors (SSRIs) are combined with NSAIDs. Caution should be used when NSAIDs are administered concomitantly with SSRIs.

### Other Information Concerning Drug Interactions

Naproxen is highly bound to plasma albumin; it thus has a theoretical potential for interaction with other albumin-bound drugs such as coumarin-type anticoagulants, sulphonylureas, hydantoins, other NSAIDs, and aspirin. Patients simultaneously receiving naproxen and a hydantoin, sulphonamide or sulphonylurea should be observed for adjustment of dose if required.

Naproxen and other non-steroidal anti-inflammatory drugs can reduce the antihypertensive effect of propranolol and other beta-blockers.

Probenecid given concurrently increases naproxen anion plasma levels and extends its plasma half-life significantly.

Due to the gastric pH elevating effects of H<sub>2</sub>-blockers, sucralfate and intensive antacid therapy, concomitant administration of naproxen delayed-release tablets is not recommended.

### Drug/Laboratory Test Interactions

Naproxen may decrease platelet aggregation and prolong bleeding time. This effect should be kept in mind when bleeding times are determined.

The administration of naproxen may result in increased urinary values for 17-ketogenic steroids because of an interaction between the drug and/or its metabolites with m-di-nitrobenzene used in this assay. Although 17-hydroxy-corticosteroid measurements (Porter-Silber test) do not appear to be artifactually altered, it is suggested that therapy with naproxen be temporarily discontinued 72 hours before adrenal function tests are performed if the Porter-Silber test is to be used.

Naproxen may interfere with some urinary assays of 5-hydroxy indoleacetic acid (5HIAA).

### Carcinogenesis

A 2 year study was performed in rats to evaluate the carcinogenic potential of naproxen at rat doses of 8, 16, and 24 mg/kg/day (50, 100, and 150 mg/m<sup>2</sup>). The maximum dose used was 0.28 times the systemic exposure to humans at the recommended dose. No evidence of tumorigenicity was found.

## **Pregnancy**

### Teratogenic Effects

#### *Pregnancy category C*

Reproduction studies have been performed in rats at 20 mg/kg/day (125 mg/m<sup>2</sup>/day, 0.23 times the human systemic exposure), rabbits at 20 mg/kg/day (220 mg/m<sup>2</sup>/day, 0.27 times the human systemic exposure), and mice at 170 mg/kg/day (510 mg/m<sup>2</sup>/day, 0.28 times the human systemic exposure) with no evidence of impaired fertility or harm to the fetus due to the drug. However, animal reproduction studies are not always predictive of human response. There are no adequate and well-controlled studies in pregnant women. Naproxen delayed-release tablets should be used in pregnancy only if the potential benefit justifies the potential risk to the fetus.

### Nonteratogenic Effects

There is some evidence to suggest that when inhibitors of prostaglandin synthesis are used to delay preterm labor there is an increased risk of neonatal complications such as necrotizing enterocolitis, patent ductus arteriosus and intracranial hemorrhage. Naproxen treatment given in late pregnancy to delay parturition has been associated with persistent pulmonary hypertension, renal dysfunction and abnormal prostaglandin E levels in preterm infants. Because of the known effects of non-steroidal anti-inflammatory drugs on the fetal cardiovascular system (closure of ductus arteriosus), use during pregnancy (particularly late pregnancy) should be avoided.

## **Labor and Delivery**

In rat studies with NSAIDs, as with other drugs known to inhibit prostaglandin synthesis, an increased incidence of dystocia, delayed parturition, and decreased pup survival occurred. Naproxen-containing products are not recommended in labor and delivery because, through its prostaglandin synthesis inhibitory effect, naproxen may adversely affect fetal circulation and inhibit uterine contractions, thus increasing the risk of uterine hemorrhage. The effects of naproxen delayed-release tablets on labor and delivery in pregnant women are unknown.

## **Nursing Mothers**

The naproxen anion has been found in the milk of lactating women at a concentration equivalent to approximately 1% of maximum naproxen concentration in plasma. Because of the possible adverse effects of prostaglandin-inhibiting drugs on neonates, use in nursing mothers should be avoided.

## **Pediatric Use**

Safety and effectiveness in pediatric patients below the age of 2 years have not been established. Pediatric dosing recommendations for juvenile arthritis are based on well-controlled studies (see **DOSAGE AND ADMINISTRATION**). There are no adequate effectiveness or dose-response data for other pediatric conditions, but the experience in juvenile arthritis and other use experience have established that single doses of 2.5 to 5 mg/kg (as naproxen suspension, see **DOSAGE AND ADMINISTRATION**), with total daily dose not exceeding 15 mg/kg/day, are well tolerated in pediatric patients over 2 years of age.

## **Geriatric Use**

Studies indicate that although total plasma concentration of naproxen is unchanged, the unbound plasma fraction of naproxen is increased in the elderly. Caution is advised when high doses are required and some adjustment of dosage may be required in elderly patients. As with other drugs used in the elderly, it is prudent to use the lowest effective dose.

Experience indicates that geriatric patients may be particularly sensitive to certain adverse effects of

non-steroidal anti-inflammatory drugs. Elderly or debilitated patients seem to tolerate peptic ulceration or bleeding less well when these events do occur. Most spontaneous reports of fatal GI events are in the geriatric population (see **WARNINGS**).

Naproxen is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function. Geriatric patients may be at a greater risk for the development of a form of renal toxicity precipitated by reduced prostaglandin formation during administration of non-steroidal anti-inflammatory drugs (see **WARNINGS, Renal Effects**).

## **ADVERSE REACTIONS**

Adverse reactions reported in controlled clinical trials in 960 patients treated for rheumatoid arthritis or osteoarthritis are listed below. In general, reactions in patients treated chronically were reported 2 to 10 times more frequently than they were in short-term studies in the 962 patients treated for mild to moderate pain or for dysmenorrhea. The most frequent complaints reported related to the gastrointestinal tract.

A clinical study found gastrointestinal reactions to be more frequent and more severe in rheumatoid arthritis patients taking daily doses of 1500 mg naproxen compared to those taking 750 mg naproxen (see **CLINICAL PHARMACOLOGY**).

In controlled clinical trials with about 80 pediatric patients and in well-monitored, open-label studies with about 400 pediatric patients with juvenile arthritis treated with naproxen, the incidence of rash and prolonged bleeding times were increased, the incidence of gastrointestinal and central nervous system reactions were about the same, and the incidence of other reactions were lower in pediatric patients than in adults.

In patients taking naproxen in clinical trials, the most frequently reported adverse experiences in approximately 1% to 10% of patients are:

**Gastrointestinal (GI) Experiences, including:** heartburn\*, abdominal pain\*, nausea\*, constipation\*, diarrhea, dyspepsia, stomatitis

**Central Nervous System:** headache\*, dizziness\*, drowsiness\*, lightheadedness, vertigo

**Dermatologic:** pruritus (itching)\*, skin eruptions\*, ecchymoses\*, sweating, purpura

**Special Senses:** tinnitus\*, visual disturbances, hearing disturbances

**Cardiovascular:** edema\*, palpitations

**General:** dyspnea\*, thirst

\* Incidence of reported reaction between 3% and 9%. Those reactions occurring in less than 3% of the patients are unmarked.

In patients taking NSAIDs, the following adverse experiences have also been reported in approximately 1% to 10% of patients.

**Gastrointestinal (GI) Experiences, including:** flatulence, gross bleeding/perforation, GI ulcers (gastric/duodenal), vomiting

**General:** abnormal renal function, anemia, elevated liver enzymes, increased bleeding time, rashes

The following are additional adverse experiences reported in < 1% of patients taking naproxen during clinical trials and through postmarketing reports. Those adverse reactions observed through postmarketing reports are italicized.

**Body as a Whole:** *anaphylactoid reactions, angioneurotic edema, menstrual disorders, pyrexia (chills and fever)*

**Cardiovascular:** *congestive heart failure, vasculitis, hypertension, pulmonary edema*

**Gastrointestinal:** *gastrointestinal bleeding and/or perforation, hematemesis, pancreatitis, vomiting, colitis, exacerbation of inflammatory bowel disease (ulcerative colitis, Crohn's disease), nonpeptic gastrointestinal ulceration, ulcerative stomatitis, esophagitis, peptic ulceration*

**Hepatobiliary:** *jaundice, abnormal liver function tests, hepatitis (some cases have been fatal)*

**Hemic and Lymphatic:** *eosinophilia, leucopenia, melena, thrombocytopenia, agranulocytosis, granulocytopenia, hemolytic anemia, aplastic anemia*

**Metabolic and Nutritional:** *hyperglycemia, hypoglycemia*

**Nervous System:** *inability to concentrate, depression, dream abnormalities, insomnia, malaise, myalgia, muscle weakness, aseptic meningitis, cognitive dysfunction, convulsions*

**Respiratory:** *eosinophilic pneumonitis, asthma*

**Dermatologic:** *alopecia, urticaria, skin rashes, toxic epidermal necrolysis, erythema multiforme, erythema nodosum, fixed drug eruption, lichen planus, pustular reaction, systemic lupus erythematosus, bullous reactions, including Stevens-Johnson syndrome, photosensitive dermatitis, photosensitivity reactions, including rare cases resembling porphyria cutanea tarda (pseudoporphyria) or epidermolysis bullosa. If skin fragility, blistering or other symptoms suggestive of pseudoporphyria occur, treatment should be discontinued and the patient monitored.*

**Special Senses:** *hearing impairment, corneal opacity, papillitis, retrobulbar optic neuritis, papilledema*

**Urogenital:** *glomerular nephritis, hematuria, hyperkalemia, interstitial nephritis, nephrotic syndrome, renal disease, renal failure, renal papillary necrosis, raised serum creatinine*

**Reproduction (female):** *infertility*

In patients taking NSAIDs, the following adverse experiences have also been reported in < 1% of patients.

**Body as a Whole:** *fever, infection, sepsis, anaphylactic reactions, appetite changes, death*

**Cardiovascular:** *hypertension, tachycardia, syncope, arrhythmia, hypotension, myocardial infarction*

**Gastrointestinal:** *dry mouth, esophagitis, gastric/peptic ulcers, gastritis, glossitis, eructation*

**Hepatobiliary:** *hepatitis, liver failure*

**Hemic and Lymphatic:** *rectal bleeding, lymphadenopathy, pancytopenia*

**Metabolic and Nutritional:** *weight changes*

**Nervous System:** *anxiety, asthenia, confusion, nervousness, paresthesia, somnolence, tremors, convulsions, coma, hallucinations*

**Respiratory:** *asthma, respiratory depression, pneumonia*

**Dermatologic:** *exfoliative dermatitis*

**Special Senses:** *blurred vision, conjunctivitis*

**Urogenital:** *cystitis, dysuria, oliguria/polyuria, proteinuria*

## OVERDOSAGE

### Symptoms and Signs

Significant naproxen overdose may be characterized by lethargy, dizziness, drowsiness, epigastric pain, abdominal discomfort, heartburn, indigestion, nausea, transient alterations in liver function, hypoprothrombinemia, renal dysfunction, metabolic acidosis, apnea, disorientation or vomiting.

Gastrointestinal bleeding can occur. Hypertension, acute renal failure, respiratory depression, and coma may occur, but are rare. Anaphylactoid reactions have been reported with therapeutic ingestion of NSAIDs, and may occur following an overdose. A few patients have experienced convulsions, but it is not clear whether or not these were drug-related. It is not known what dose of the drug would be life threatening. The oral LD<sub>50</sub> of the drug is 543 mg/kg in rats, 1234 mg/kg in mice, 4110 mg/kg in hamsters, and greater than 1000 mg/kg in dogs.

### **Treatment**

Patients should be managed by symptomatic and supportive care following a NSAID overdose. There are no specific antidotes. Hemodialysis does not decrease the plasma concentration of naproxen because of the high degree of its protein binding. Emesis and/or activated charcoal (60 to 100 g in adults, 1 to 2 g/kg in children) and/or osmotic cathartic may be indicated in patients seen within 4 hours of ingestion with symptoms or following a large overdose. Forced diuresis, alkalization of urine or hemoperfusion may not be useful due to high protein binding.

### **DOSAGE AND ADMINISTRATION**

Carefully consider the potential benefits and risks of naproxen delayed-release tablets USP and other treatment options before deciding to use naproxen delayed-release tablets USP. Use the lowest effective dose for the shortest duration consistent with individual patient treatment goals (see **WARNINGS**).

After observing the response to initial therapy with naproxen delayed-release tablets USP, the dose and frequency should be adjusted to suit an individual patient's needs.

**Different dose strengths and formulations (i.e., tablets, suspension) of the drug are not necessarily bioequivalent. This difference should be taken into consideration when changing formulation.**

Although naproxen tablets, naproxen suspension, naproxen delayed-release tablets USP, and naproxen sodium tablets all circulate in the plasma as naproxen, they have pharmacokinetic differences that may affect onset of action. Onset of pain relief can begin within 30 minutes in patients taking naproxen sodium and within 1 hour in patients taking naproxen. Because naproxen delayed-release tablets USP dissolve in the small intestine rather than in the stomach, the absorption of the drug is delayed compared to the other naproxen formulations (see **CLINICAL PHARMACOLOGY**).

The recommended strategy for initiating therapy is to choose a formulation and a starting dose likely to be effective for the patient and then adjust the dosage based on observation of benefit and/or adverse events. A lower dose should be considered in patients with renal or hepatic impairment or in elderly patients (see **WARNINGS** and **PRECAUTIONS**).

### **Geriatric Patients**

Studies indicate that although total plasma concentration of naproxen is unchanged, the unbound plasma fraction of naproxen is increased in the elderly. Caution is advised when high doses are required and some adjustment of dosage may be required in elderly patients. As with other drugs used in the elderly, it is prudent to use the lowest effective dose.

### **Patients with Moderate to Severe Renal Impairment**

Naproxen-containing products are not recommended for use in patients with moderate to severe and severe renal impairment (creatinine clearance < 30 mL/min) (see **WARNINGS, Renal Effects**).

### **Rheumatoid Arthritis, Osteoarthritis and Ankylosing Spondylitis**

|                          |        |             |
|--------------------------|--------|-------------|
| Naproxen Delayed-Release | 375 mg | twice daily |
|--------------------------|--------|-------------|

|             |           |             |
|-------------|-----------|-------------|
| Tablets USP | or 500 mg | twice daily |
|-------------|-----------|-------------|

To maintain the integrity of the enteric coating, the naproxen delayed-release tablets USP should not be broken, crushed, or chewed during ingestion.

During long-term administration, the dose of naproxen may be adjusted up or down depending on the clinical response of the patient. A lower daily dose may suffice for long-term administration. The morning and evening doses do not have to be equal in size and the administration of the drug more frequently than twice daily is not necessary.

In patients who tolerate lower doses well, the dose may be increased to naproxen 1500 mg/day for limited periods of up to 6 months when a higher level of anti-inflammatory/analgesic activity is required. When treating such patients with naproxen 1500 mg/day, the physician should observe sufficient increased clinical benefits to offset the potential increased risk. The morning and evening doses do not have to be equal in size and administration of the drug more frequently than twice daily does not generally make a difference in response (see **CLINICAL PHARMACOLOGY**).

### **Juvenile Arthritis**

The recommended total daily dose of naproxen is approximately 10 mg/kg given in 2 divided doses (i.e., 5 mg/kg given twice a day). Naproxen delayed-release tablets USP are not well suited to this dosage so use of naproxen oral suspension is recommended for this indication.

### **HOW SUPPLIED**

Naproxen delayed-release tablets USP, 375 mg are white to off-white, capsule-shaped, enteric-coated, unscored tablets imprinted on one side in blue ink with “93-5”. They are available in bottles of 100 and 500.

Naproxen delayed-release tablets USP, 500 mg are white to off-white, capsule-shaped, enteric-coated, unscored tablets imprinted on one side in blue ink with “93-6”. They are available in bottles of 100 and 500.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].

Dispense in a tight, light-resistant container as defined in the USP, with a child-resistant closure (as required).

Manufactured In Israel By:

**TEVA PHARMACEUTICAL IND. LTD.**

Jerusalem, 91010, Israel

Manufactured For:

**TEVA PHARMACEUTICALS USA**

Sellersville, PA 18960

Rev. H 6/2010

### **Medication Guide for Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)**

**Rx only**

(See the end of this Medication Guide for a list of prescription NSAID medicines.)

**What is the most important information I should know about medicines called Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)?**

**NSAID medicines may increase the chance of a heart attack or stroke that can lead to death.** This chance increases:



- with longer use of NSAID medicines
- in people who have heart disease

**NSAID medicines should never be used right before or after a heart surgery called a “coronary artery bypass graft (CABG).”**

**NSAID medicines can cause ulcers and bleeding in the stomach and intestines at any time during treatment. Ulcers and bleeding:**

- can happen without warning symptoms
- may cause death

**The chance of a person getting an ulcer or bleeding increases with:**

- taking medicines called “corticosteroids” and “anticoagulants”
- longer use
- smoking
- drinking alcohol
- older age
- having poor health

**NSAID medicines should only be used:**

- exactly as prescribed
- at the lowest dose possible for your treatment
- for the shortest time needed

**What are Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)?**

NSAID medicines are used to treat pain and redness, swelling, and heat (inflammation) from medical conditions such as:

- different types of arthritis
- menstrual cramps and other types of short-term pain

**Who should not take a Non-Steroidal Anti-Inflammatory Drug (NSAID)?**

**Do not take an NSAID medicine:**

- if you had an asthma attack, hives, or other allergic reaction with aspirin or any other NSAID medicine
- for pain right before or after heart bypass surgery

**Tell your healthcare provider:**

- about all of your medical conditions.
- about all of the medicines you take. NSAIDs and some other medicines can interact with each other and cause serious side effects. **Keep a list of your medicines to show to your healthcare provider and pharmacist.**
- if you are pregnant. **NSAID medicines should not be used by pregnant women late in their pregnancy.**
- if you are breastfeeding. **Talk to your doctor.**

**What are the possible side effects of Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)?**

| <b>Serious Side effects include:</b> | <b>Other side effects include:</b> |
|--------------------------------------|------------------------------------|
| • heart attack                       | • stomach pain                     |
| • stroke                             | • constipation                     |
| • high blood pressure                | • diarrhea                         |

|  |             |
|--|-------------|
| • heart failure from body swelling (fluid retention) | • gas       |
| • kidney problems including kidney failure           | • heartburn |
| • bleeding and ulcers in the stomach and intestine   | • nausea    |
| • low red blood cells (anemia)                       | • vomiting  |
| • life-threatening skin reactions                    | • dizziness |
| • life-threatening allergic reactions                |             |
| • liver problems including liver failure             |             |
| • asthma attacks in people who have asthma           |             |

**Get emergency help right away if you have any of the following symptoms:**

- shortness of breath or trouble breathing
- chest pain
- weakness in one part or side of your body
- slurred speech
- swelling of the face or throat

**Stop your NSAID medicine and call your healthcare provider right away if you have any of the following symptoms:**

- nausea
- more tired or weaker than usual
- itching
- your skin or eyes look yellow
- stomach pain
- flu-like symptoms
- vomit blood
- there is blood in your bowel movement or it is black and sticky like tar
- unusual weight gain
- skin rash or blisters with fever
- swelling of the arms and legs, hands and feet

These are not all the side effects with NSAID medicines. Talk to your healthcare provider or pharmacist for more information about NSAID medicines.

**Other information about Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)**

- Aspirin is an NSAID medicine but it does not increase the chance of a heart attack. Aspirin can cause bleeding in the brain, stomach, and intestines. Aspirin can also cause ulcers in the stomach and intestines.
- Some of these NSAID medicines are sold in lower doses without a prescription (over-the-counter). Talk to your healthcare provider before using over-the-counter NSAIDs for more than 10 days.

**NSAID medicines that need a prescription**

Generic Name/Tradename:

Celecoxib/Celebrex

Diclofenac/Cataflam, Voltaren, Arthrotec

Difunisal/Dolobid  
Etodolac/Lodine, Lodine XL  
Fenoprofen/Nalfon, Nalfon 200  
Flurbiprofen/Ansaid  
Ibuprofen/Motrin, Tab-Profen, Vicoprofen  
Indomethacin/Indocni, Indocin SR, Indo-Lemmon  
Ketoprofen/Oruvail  
Ketorolac/Toradol  
Mefenamic Acid/Ponstel  
Meloxicam/Mobic  
Nabumetone/Relafen  
Naproxen/Naprosyn, Anaprox, Anaprox DS, Naprelan, Naprapac  
  
Oxaprozin/Daypro  
Piroxicam/Feldene  
Sulindac/Clinoril  
Tolmetin/Toletin, Tolectin DS, Tolectin 600

**Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.**

*This Medication Guide has been approved by the U.S. Food and Drug Administration.*

Manufactured In Israel By:

**TEVA PHARMACEUTICAL IND. LTD.**

Jerusalem, 91010, Israel

Manufactured For:

**TEVA PHARMACEUTICALS USA**

Sellersville, PA 18960

Rev. B 7/2009

Repacked by:

**H.J. Harkins Company, Inc.**

Nipomo, CA 93444

**PRINCIPAL DISPLAY PANEL**

52959-516-30

RX Only: #XXXXXXXX

#XXX

CAUTION: Federal law PROHIBITS the transfer of this drug to anyone other than the person to whom prescribed and prohibits dispensing without a prescription unless OTC. See outsert for add'l RX info KEEP OUT OF REACH OF CHILDREN. Store in a cool dry place 68 to 77 degrees F.

**NAPROXEN 500mg D.R.TABLET(EC)**

Lot #: NPE550TV

Mfg: TEVA PHARM

Exp: 06/09

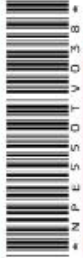
Compare to: Naprosyn E.C.

Mfg Israel

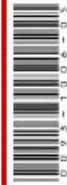
Mfg. NDC: 0093-1006-05

Loc.:

Pill ID: White capsule-shaped tablets



Take as directed by your Doctor or  
See outsert for usual dosage information



NAPROXEN 500mg D.R.TABLET(EC)  
52959-516-30 Qty #30  
06/09 Lot NPE550TV  
Naprosyn E.C. 0093-1006-05

NAPROXEN 500mg D.R.TABLET(EC)  
52959-516-30 Qty #30  
06/09 Lot NPE550TV  
Naprosyn E.C. 0093-1006-05

NAPROXEN 500mg D.R.TABLET(EC)  
52959-516-30 Qty #30  
06/09 Lot NPE550TV  
Naprosyn E.C. 0093-1006-05

NAPROXEN 500mg D.R.TABLET(EC)  
52959-516-30 Qty #30  
06/09 Lot NPE550TV  
Naprosyn E.C. 0093-1006-05

Repack: HJ Harkins Co., Inc. Nipomo, CA 93444  
Dispense in tight, child & light-resistant container per USP

**Naproxen Delayed-Release Tablets USP 500 mg 100s Label Text**

NDC 0093-1006-01

**NAPROXEN**

**DELAYED-RELEASE**

**Tablets USP**

**500 mg**

PHARMACIST: Dispense the Medication Guide  
provided separately to each patient.

**Rx only**

**100 TABLETS**

**TEVA**

**NAPROXEN**

naproxen tablet, delayed release

**Product Information**

|                                |                         |                           |                              |
|--------------------------------|-------------------------|---------------------------|------------------------------|
| <b>Product Type</b>            | HUMAN PRESCRIPTION DRUG | <b>Item Code (Source)</b> | NDC:52959-516(NDC:0093-1006) |
| <b>Route of Administration</b> | ORAL                    |                           |                              |

**Active Ingredient/Active Moiety**

| Ingredient Name  | Basis of Strength | Strength |
|--|-------------------|----------|
| NAPROXEN (UNII: 57Y76R9ATQ) (NAPROXEN - UNII:57Y76R9ATQ) | NAPROXEN          | 500 mg   |

**Inactive Ingredients**

| Ingredient Name   | Strength |
|---|----------|
| STARCH, CORN (UNII: O8232NY3SJ)   |          |
| CROSCARMELOSE SODIUM (UNII: M28OL1HH48)                                     |          |
| FD&C BLUE NO. 2 (UNII: L06K8R7DQK)  |          |
| MAGNESIUM STEARATE (UNII: 70097M6I30)                                       |          |
| METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J) |          |
| POVIDONE K90 (UNII: RDH86HJV5Z)   |          |
| TALC (UNII: 7SEV7J4R1U)   |          |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)   |          |
| TRIETHYL CITRATE (UNII: 8Z96QXD6UM)   |          |

### Product Characteristics

|          |                            |              |          |
|----------|----------------------------|--------------|----------|
| Color    | WHITE (white to off-white) | Score        | no score |
| Shape    | OVAL (capsule-shaped)      | Size         | 16mm     |
| Flavor   |                            | Imprint Code | 93;6     |
| Contains |                            |              |          |

### Packaging

| # | Item Code        | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:52959-516-02 | 120 in 1 BOTTLE     |                      |                    |
| 2 | NDC:52959-516-10 | 10 in 1 BOTTLE      |                      |                    |
| 3 | NDC:52959-516-14 | 14 in 1 BOTTLE      |                      |                    |
| 4 | NDC:52959-516-15 | 15 in 1 BOTTLE      |                      |                    |
| 5 | NDC:52959-516-20 | 20 in 1 BOTTLE      |                      |                    |
| 6 | NDC:52959-516-30 | 30 in 1 BOTTLE      |                      |                    |
| 7 | NDC:52959-516-40 | 40 in 1 BOTTLE      |                      |                    |
| 8 | NDC:52959-516-42 | 42 in 1 BOTTLE      |                      |                    |
| 9 | NDC:52959-516-60 | 60 in 1 BOTTLE      |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| ANDA               | ANDA075227                               | 09/23/2010           |                    |

**Labeler** - H.J. Harkins Company, Inc. (147681894)

### Establishment

| Name                         | Address | ID/FEI    | Business Operations |
|------------------------------|---------|-----------|---------------------|
| Teva Pharmaceuticals USA Inc |         | 118234421 | manufacture         |