ACAMPROSATE CALCIUM- acamprosate calcium enteric-coated tablet, delayed release
Mylan Pharmaceuticals Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use acamprosate calcium delayed-release tablets safely and effectively. See full prescribing information for acamprosate calcium delayed-release tablets.

ACAMPROSATE calcium delayed-release tablets, for oral use
Initial U.S. Approval: 2004

- INDICATIONS AND USAGE -

• Acamprosate calcium delayed-release tablets are indicated for the maintenance of abstinence from alcohol in patients with alcohol dependence who are abstinent at treatment initiation (1, 14).
• Treatment with acamprosate calcium delayed-release tablets should be part of a comprehensive management program that includes psychosocial support (1).

- DOSAGE AND ADMINISTRATION -

• Recommended dose: 666 mg (two 333 mg tablets) taken three times daily (2).
• Dose reduction to one 333 mg tablet taken three times daily for patients with moderate renal impairment (creatinine clearance 30 to 50 mL/min) (2.1).
• Acamprosate calcium delayed-release tablets are contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) (2.1, 4.2, 5.1, 8.6, 12.3).

- DOSAGE FORMS AND STRENGTHS -
Enteric-coated tablets, 333 mg (3).

- CONTRAINDICATIONS -

• Acamprosate calcium delayed-release tablets are contraindicated in patients who previously have exhibited hypersensitivity to acamprosate calcium or any of its components (4.1).
• Acamprosate calcium is contraindicated in patients with severe renal impairment (4.2).

- WARNINGS AND PRECAUTIONS -

• Dose reduction is required for patients with moderate renal impairment (5.1).
• Monitor patients for depression or suicidal ideation and prompt patients, families, and caregivers to report such symptoms to the health care provider (5.2).

- ADVERSE REACTIONS -

Common adverse events that occurred in any acamprosate calcium treatment group at a rate of 3% or greater and greater than the placebo group in controlled clinical trials with spontaneously reported adverse events are: accidental injury, asthenia, pain, anorexia, diarrhea, flatulence, nausea, anxiety, depression, dizziness, dry mouth, insomnia, paresthesia, pruritus and sweating (6.1).
To report SUSPECTED ADVERSE REACTIONS, contact Mylan Pharmaceuticals Inc. at 1-877-446-3679 (1-877-4-INFO-RX) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

- USE IN SPECIFIC POPULATIONS -

• Pregnancy: Acamprosate calcium should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus (8.1).
• Nursing Mothers: Caution should be exercised when acamprosate calcium is administered to a nursing woman (8.3).
• Renal Impairment: Dose reduction required for moderate renal impairment; contraindicated in severe renal impairment (2.1, 4.2, 5.1, 8.6, 12.3)

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 11/2014
FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE
Acamprosate calcium delayed-release tablets are indicated for the maintenance of abstinence from alcohol in patients with alcohol dependence who are abstinent at treatment initiation. Treatment with acamprosate calcium delayed-release tablets should be part of a comprehensive management program that includes psychosocial support.

The efficacy of acamprosate calcium in promoting abstinence has not been demonstrated in subjects who have not undergone detoxification and not achieved alcohol abstinence prior to beginning acamprosate calcium delayed-release tablet treatment. The efficacy of acamprosate calcium in
promoting abstinence from alcohol in polysubstance abusers has not been adequately assessed.

2 DOSAGE AND ADMINISTRATION
The recommended dose of acamprosate calcium delayed-release tablets is two 333 mg tablets (each dose should total 666 mg) taken three times daily. A lower dose may be effective in some patients.

Although dosing may be done without regard to meals, dosing with meals was employed during clinical trials and is suggested in those patients who regularly eat three meals daily.

Treatment with acamprosate calcium delayed-release tablets should be initiated as soon as possible after the period of alcohol withdrawal, when the patient has achieved abstinence, and should be maintained if the patient relapses. Acamprosate calcium delayed-release tablets should be used as part of a comprehensive psychosocial treatment program.

2.1 Dosage in Renal Impairment
For patients with moderate renal impairment (creatinine clearance of 30 to 50 mL/min), a starting dose of one 333 mg tablet taken three times daily is recommended. Acamprosate calcium delayed-release tablets are contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Contraindications (4.2), Warnings and Precautions (5.1), Use in Specific Populations (8.6) and Clinical Pharmacology (12.3)].

3 DOSAGE FORMS AND STRENGTHS
Acamprosate calcium delayed-release 333 mg tablets are white enteric-coated, round, unscored tablets imprinted with M over AC in black ink on one side of the tablet and blank on the other side.

4 CONTRAINDICATIONS
4.1 Hypersensitivity to Acamprosate Calcium
Acamprosate calcium delayed-release tablets are contraindicated in patients who previously have exhibited hypersensitivity to acamprosate calcium or any of its components.

4.2 Severe Renal Impairment
Acamprosate calcium delayed-release tablets are contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Dosage and Administration (2.1), Warnings and Precautions (5.1), Use in Specific Populations (8.6) and Clinical Pharmacology (12.3)].

5 WARNINGS AND PRECAUTIONS
May contain sodium sulfite, a sulfite that may cause allergic-type reactions including anaphylactic symptoms and life threatening or less severe asthmatic episodes in certain susceptible people. The overall prevalence of sulfite sensitivity in the general population is unknown and probably low. Sulfite sensitivity is seen more frequently in asthmatic than in nonasthmatic people.

5.1 Renal Impairment
Treatment with acamprosate calcium in patients with moderate renal impairment (creatinine clearance of 30 to 50 mL/min) requires a dose reduction [see Dosage and Administration (2.1)]. Acamprosate is contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Dosage and Administration (2.1), Contraindications (4.2), Use in Specific Populations (8.6) and Clinical Pharmacology (12.3)].
5.2 Suicidality and Depression

In controlled clinical trials of acamprosate calcium, adverse events of a suicidal nature (suicidal ideation, suicide attempts, completed suicides) were infrequent overall, but were more common in acamprosate calcium-treated patients than in patients treated with placebo (1.4% vs. 0.5% in studies of 6 months or less; 2.4% vs. 0.8% in year-long studies). Completed suicides occurred in 3 of 2,272 (0.13%) patients in the pooled acamprosate group from all controlled studies and 2 of 1,962 patients (0.10%) in the placebo group. Adverse events coded as "depression" were reported at similar rates in acamprosate calcium-treated and placebo-treated patients. Although many of these events occurred in the context of alcohol relapse, and the interrelationship between alcohol dependence, depression and suicidality is well-recognized and complex, no consistent pattern of relationship between the clinical course of recovery from alcoholism and the emergence of suicidality was identified. Alcohol-dependent patients, including those patients being treated with acamprosate calcium, should be monitored for the development of symptoms of depression or suicidal thinking. Families and caregivers of patients being treated with acamprosate should be alerted to the need to monitor patients for the emergence of symptoms of depression or suicidality, and to report such symptoms to the patient's health care provider.

5.3 Alcohol Withdrawal

Use of acamprosate calcium does not eliminate or diminish withdrawal symptoms.

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

Clinically significant serious adverse reactions associated with acamprosate calcium described elsewhere in labeling include suicidality and depression and acute kidney failure [see Warnings and Precautions (5.2) and Adverse Reactions (6.2)].

The adverse event data described below reflect the safety experience in over 7,000 patients exposed to acamprosate calcium for up to one year, including over 2,000 acamprosate calcium-exposed patients who participated in placebo-controlled trials.

Adverse Events Leading to Discontinuation

In placebo-controlled trials of 6 months or less, 8% of acamprosate calcium-treated patients discontinued treatment due to an adverse event, as compared to 6% of patients treated with placebo. In studies longer than 6 months, the discontinuation rate due to adverse events was 7% in both the placebo-treated and the acamprosate calcium-treated patients. Only diarrhea was associated with the discontinuation of more than 1% of patients (2% of acamprosate calcium-treated vs. 0.7% of placebo-treated patients). Other events, including nausea, depression and anxiety, while accounting for discontinuation in less than 1% of patients, were nevertheless more commonly cited in association with discontinuation in acamprosate calcium-treated patients than in placebo-treated patients.

Common Adverse Events Reported in Controlled Trials

Common adverse events were collected spontaneously in some controlled studies and using a checklist in other studies. The overall profile of adverse events was similar using either method. Table 1 shows those events that occurred in any acamprosate calcium treatment group at a rate of 3% or greater and greater than the placebo group in controlled clinical trials with spontaneously reported adverse events. The reported frequencies of adverse events represent the proportion of individuals who experienced, at least once, a treatment-emergent adverse event of the type listed, without regard to the causal
relationship of the events to the drug.

Table 1. Events Occurring at a Rate of at Least 3% and Greater than Placebo in any Acamprosate Calcium Treatment Group in Controlled Clinical Trials with Spontaneously Reported Adverse Events.

<table>
<thead>
<tr>
<th>Body System/Preferred Term</th>
<th>Number of Patients (%) with Events</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Acamprosate Calcium 1332 mg/day</td>
</tr>
<tr>
<td>Number of patients in Treatment Group</td>
<td>397</td>
</tr>
<tr>
<td>Number (%) of patients with an AE</td>
<td>248 (62%)</td>
</tr>
<tr>
<td>Body as a Whole</td>
<td>121 (30%)</td>
</tr>
<tr>
<td>Accidental Injury‡</td>
<td>17 (4%)</td>
</tr>
<tr>
<td>Asthenia</td>
<td>29 (7%)</td>
</tr>
<tr>
<td>Pain</td>
<td>6 (2%)</td>
</tr>
<tr>
<td>Digestive System</td>
<td>85 (21%)</td>
</tr>
<tr>
<td>Anorexia</td>
<td>20 (5%)</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>39 (10%)</td>
</tr>
<tr>
<td>Flatulence</td>
<td>4 (1%)</td>
</tr>
<tr>
<td>Nausea</td>
<td>11 (3%)</td>
</tr>
<tr>
<td>Nervous System</td>
<td>150 (38%)</td>
</tr>
<tr>
<td>Anxiety§</td>
<td>32 (8%)</td>
</tr>
<tr>
<td>Depression</td>
<td>33 (8%)</td>
</tr>
<tr>
<td>Dizziness</td>
<td>15 (4%)</td>
</tr>
<tr>
<td>Dry mouth</td>
<td>13 (3%)</td>
</tr>
<tr>
<td>Insomnia</td>
<td>34 (9%)</td>
</tr>
<tr>
<td>Paresthesia</td>
<td>11 (3%)</td>
</tr>
<tr>
<td>Skin and Appendages</td>
<td>26 (7%)</td>
</tr>
<tr>
<td>Pruritus</td>
<td>12 (3%)</td>
</tr>
<tr>
<td>Sweating</td>
<td>11 (3%)</td>
</tr>
</tbody>
</table>

* includes 258 patients treated with acamprosate calcium 2000 mg/day, using a different dosage strength and regimen.
† includes all patients in the first two columns as well as 83 patients treated with acamprosate calcium 3000 mg/day, using a different dosage strength and regimen.
‡ includes events coded as “fracture” by sponsor;
§ includes events coded as “nervousness” by sponsor

Concomitant Therapies

In clinical trials, the safety profile in subjects treated with acamprosate calcium concomitantly with anxiolytics, hypnotics and sedatives (including benzodiazepines), or non-opioid analgesics was similar to that of subjects taking placebo with these concomitant medications. Patients taking acamprosate calcium concomitantly with antidepressants more commonly reported both weight gain and weight loss, compared with patients taking either medication alone.

Other Events Observed During the Premarketing Evaluation of Acamprosate Calcium
Following is a list of terms that reflect treatment-emergent adverse events reported by patients treated with acamprosate calcium in 20 clinical trials (4,461 patients treated with acamprosate calcium, 3,526 of whom received the maximum recommended dose of 1998 mg/day for up to one year in duration). This listing does not include those events already listed above; events for which a drug cause was considered remote; event terms which were so general as to be uninformative; and events reported only once which were not likely to be acutely life threatening.

Events are further categorized by body system and listed in order of decreasing frequency according to the following definitions: frequent adverse events are those occurring in at least 1/100 patients (only those not already listed in the summary of adverse events in controlled trials appear in this listing); infrequent adverse events are those occurring in 1/100 to 1/1,000 patients; rare events are those occurring in fewer than 1/1,000 patients.

**Body as a Whole** – **Frequent:** headache, abdominal pain, back pain, infection, flu syndrome, chest pain, chills, suicide attempt; **Infrequent:** fever, intentional overdose, malaise, allergic reaction, abscess, neck pain, hernia, intentional injury; **Rare:** ascites, face edema, photosensitivity reaction, abdomen enlarged, sudden death.

**Cardiovascular System** – **Frequent:** palpitation, syncope; **Infrequent:** hypotension, tachycardia, hemorrhage, angina pectoris, migrane, varicose vein, myocardial infarct, phlebitis, postural hypotension; **Rare:** heart failure, mesenteric arterial occlusion, cardiomyopathy, deep thrombophlebitis, shock.

**Digestive System** – **Frequent:** vomiting, dyspepsia, constipation, increased appetite; **Infrequent:** liver function tests abnormal, gastroenteritis, gastritis, dysphagia, eructation, gastrointestinal hemorrhage, pancreatitis, rectal hemorrhage, liver cirrhosis, esophagitis, hematemesis, nausea and vomiting, hepatitis; **Rare:** melena, stomach ulcer, cholecystitis, colitis, duodenal ulcer, mouth ulceration, carcinoma of liver.

**Endocrine System** – **Rare:** goiter, hypothyroidism.

**Hemic and Lymphatic System** – **Infrequent:** anemia, ecchymosis, eosinophilia, lymphocytosis, thrombocytopenia; **Rare:** leukopenia, lymphadenopathy, monocytosis.

**Metabolic and Nutritional Disorders** – **Frequent:** peripheral edema, weight gain; **Infrequent:** weight loss, hyperglycemia, SGOT increased, SGPT increased, gout, thirst, hyperuricemia, diabetes mellitus, avitaminosis, bilirubinemia; **Rare:** alkaline phosphatase increased, creatinine increased, hyponatremia, lactic dehydrogenase increased.

**Musculoskeletal System** – **Frequent:** myalgia, arthralgia; **Infrequent:** leg cramps; **Rare:** rheumatoid arthritis, myopathy.

**Nervous System** – **Frequent:** somnolence, libido decreased, amnesia, thinking abnormal, tremor, vasodilatation, hypertension; **Infrequent:** convulsion, confusion, libido increased, vertigo, withdrawal syndrome, apathy, suicidal ideation, neuralgia, hostility, agitation, neurosis, abnormal dreams, hallucinations, hypesthesia; **Rare:** alcohol craving, psychosis, hyperkinesia, twitching, depersonalization, increased salivation, paranoid reaction, torticollis, encephalopathy, manic reaction.

**Respiratory System** – **Frequent:** rhinitis, cough increased, dyspnea, pharyngitis, bronchitis; **Infrequent:** asthma, epistaxis, pneumonia; **Rare:** laryngismus, pulmonary embolus.

**Skin and Appendages** – **Frequent:** rash; **Infrequent:** acne, eczema, alopecia, maculopapular rash, dry skin, urticaria, exfoliative dermatitis, vesiculobullous rash; **Rare:** psoriasis.

**Special Senses** – **Frequent:** abnormal vision, taste perversion; **Infrequent:** tinnitus, amblyopia, deafness; **Rare:** ophthalmitis, diplopia, photophobia.

**Urogenital System** – **Frequent:** impotence; **Infrequent:** metrorrhagia, urinary frequency, urinary tract infection, sexual function abnormal, urinary incontinence, vaginitis; **Rare:** kidney calculus, abnormal ejaculation, hematuria, menorrhagia, nocturia, polyuria, urinary urgency.
6.2 Post-marketing Experience

The following adverse reactions have been identified during post-approval use of acamprosate calcium. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Serious Adverse Events Observed During the Non-U.S. Post-marketing Evaluation of Acamprosate Calcium

The serious adverse event of acute kidney failure has been reported to be temporally associated with acamprosate calcium treatment in at least three patients and is not described elsewhere in the labeling.

7 DRUG INTERACTIONS

Acamprosate does not affect the pharmacokinetics of alcohol. The pharmacokinetics of acamprosate are not affected by alcohol, diazepam or disulfiram, and clinically important interactions between naltrexone and acamprosate were not observed [see Clinical Pharmacology (12.3)].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Pregnancy Category C. Teratogenic Effects

Acamprosate calcium has been shown to be teratogenic in rats when given in doses that are approximately equal to the human dose (on a mg/m² basis) and in rabbits when given in doses that are approximately 3 times the human dose (on a mg/m² basis). Acamprosate calcium produced a dose related increase in the number of fetuses with malformations in rats at oral doses of 300 mg/kg/day or greater (approximately equal to the maximum recommended human daily (MRHD) oral dose on a mg/m² basis). The malformations included hydronephrosis, malformed iris, retinal dysplasia and retroesophageal subclavian artery. No findings were observed at an oral dose of 50 mg/kg/day (approximately one-fifth the MRHD oral dose on a mg/m² basis). An increased incidence of hydronephrosis was also noted in Burgundy Tawny rabbits at oral doses of 400 mg/kg/day or greater (approximately 3 times the MRHD oral dose on a mg/m² basis). No developmental effects were observed in New Zealand white rabbits at oral doses up to 1000 mg/kg/day (approximately 8 times the MRHD oral dose on a mg/m² basis). The findings in animals should be considered in relation to known adverse developmental effects of ethyl alcohol, which include the characteristics of fetal alcohol syndrome (craniofacial dysmorphism, intrauterine and postnatal growth retardation, retarded psychomotor and intellectual development) and milder forms of neurological and behavioral disorders in humans. There are no adequate and well controlled studies in pregnant women. Acamprosate calcium should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Nonteratogenic Effects

A study conducted in pregnant mice that were administered acamprosate calcium by the oral route starting on Day 15 of gestation through the end of lactation on postnatal day 28 demonstrated an increased incidence of still-born fetuses at doses of 960 mg/kg/day or greater (approximately 2 times the MRHD oral dose on a mg/m² basis). No effects were observed at a dose of 320 mg/kg/day (approximately one-half the MRHD dose on a mg/m² basis).

8.2 Labor and Delivery

The potential for acamprosate calcium to affect the duration of labor and delivery is unknown.

8.3 Nursing Mothers
In animal studies, acamprosate was excreted in the milk of lactating rats dosed orally with acamprosate calcium. The concentration of acamprosate in milk compared to blood was 1.3:1. It is not known whether acamprosate is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when acamprosate calcium is administered to a nursing woman.

8.4 Pediatric Use
The safety and efficacy of acamprosate calcium have not been established in the pediatric population.

8.5 Geriatric Use
Forty-one of the 4,234 patients in double-blind, placebo-controlled, clinical trials of acamprosate calcium were 65 years of age or older, while none were 75 years of age or over. There were too few patients in the ≥ 65 age group to evaluate any differences in safety or effectiveness for geriatric patients compared to younger patients.

This drug 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 [see Clinical Pharmacology (12.3), Adverse Reactions (6.1) and Dosage and Administration (2.1)].

8.6 Renal Impairment
Acamprosate calcium is contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Dosage and Administration (2.1), Contraindications (4.2), Warnings and Precautions (5.1) and Clinical Pharmacology (12.3)].

10 OVERDOSAGE
In all reported cases of acute overdosage with acamprosate calcium (total reported doses of up to 56 grams of acamprosate calcium), the only symptom that could be reasonably associated with acamprosate calcium was diarrhea. Hypercalcemia has not been reported in cases of acute overdose. A risk of hypercalcemia should be considered in chronic overdosage only. Treatment of overdose should be symptomatic and supportive.

11 DESCRIPTION
Acamprosate calcium is supplied in an enteric-coated tablet for oral administration. Acamprosate calcium is a synthetic compound with a chemical structure similar to that of the endogenous amino acid homotaurine, which is a structural analogue of the amino acid neurotransmitter γ-aminobutyric acid and the amino acid neuromodulator taurine. Its chemical name is 3-(acetylamino)-1-propanesulfonic acid calcium salt (2:1). Its chemical formula is C₁₀H₂₀N₂O₈S₂Ca and molecular weight is 400.48. Its structural formula is:

![Structural formula of acamprosate calcium](image)
Acamprosate calcium is a white odorless crystalline powder. It is freely soluble in water, and practically insoluble in absolute ethanol, dichloromethane and xylene.

Each acamprosate calcium delayed-release tablet contains acamprosate calcium 333 mg, equivalent to 300 mg of acamprosate. Inactive ingredients in acamprosate calcium delayed-release tablets include: ammonium hydroxide, black iron oxide, colloidal silicon dioxide, glyceryl behenate, hypromellose, magnesium stearate, methacrylic acid copolymer, microcrystalline cellulose, propylene glycol, shellac and talc. Sulfites were used in the synthesis of the drug substance and traces of residual sulfites may be present in the drug product.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

The mechanism of action of acamprosate in maintenance of alcohol abstinence is not completely understood. Chronic alcohol exposure is hypothesized to alter the normal balance between neuronal excitation and inhibition. In vitro and in vivo studies in animals have provided evidence to suggest acamprosate may interact with glutamate and GABA neurotransmitter systems centrally, and has led to the hypothesis that acamprosate restores this balance.

12.2 Pharmacodynamics

Pharmacodynamic studies have shown that acamprosate calcium reduces alcohol intake in alcohol-dependent animals in a dose dependent manner and that this effect appears to be specific to alcohol and the mechanisms of alcohol dependence.

Acamprosate calcium has negligible observable central nervous system (CNS) activity in animals outside of its effects on alcohol dependence, exhibiting no anticonvulsant, antidepressant or anxiolytic activity.

The administration of acamprosate calcium is not associated with the development of tolerance or dependence in animal studies. Acamprosate calcium did not produce any evidence of withdrawal symptoms in patients in clinical trials at therapeutic doses. Post-marketing data, collected retrospectively outside the U.S. have provided no evidence of acamprosate calcium abuse or dependence.

Acamprosate calcium is not known to cause alcohol aversion and does not cause a disulfiram-like reaction as a result of ethanol ingestion.

12.3 Pharmacokinetics

Absorption

The absolute bioavailability of acamprosate calcium after oral administration is about 11%. Steady-state plasma concentrations of acamprosate are reached within 5 days of dosing. Steady-state peak plasma concentrations after acamprosate calcium doses of 2 x 333 mg tablets three times daily average 350 ng/mL and occur at 3 to 8 hours post-dose. Coadministration of acamprosate calcium with food decreases bioavailability as measured by C_{\text{max}} and AUC, by approximately 42% and 23%, respectively. The food effect on absorption is not clinically significant and no adjustment of dose is necessary.

Distribution

The volume of distribution for acamprosate following intravenous administration is estimated to be 72 to 109 liters (approximately 1 L/kg). Plasma protein binding of acamprosate is negligible.

Metabolism
Acamprosate does not undergo metabolism.

**Elimination**

After oral dosing of 2 x 333 mg of acamprosate calcium, the terminal half-life ranges from approximately 20 to 33 hours. Following oral administration of acamprosate calcium, the major route of excretion is via the kidneys as acamprosate.

**Special Populations**

**Gender**

Acamprosate calcium does not exhibit any significant pharmacokinetic differences between male and female subjects.

**Age**

The pharmacokinetics of acamprosate calcium have not been evaluated in a geriatric population. However, since renal function diminishes in elderly patients and acamprosate is excreted unchanged in urine, acamprosate plasma concentrations are likely to be higher in the elderly population compared to younger adults.

**Pediatrics**

The pharmacokinetics of acamprosate calcium have not been evaluated in a pediatric population.

**Renal Impairment**

Peak plasma concentrations after administration of a single dose of 2 x 333 mg acamprosate calcium delayed-release tablets to patients with moderate or severe renal impairment were about 2-fold and 4-fold higher, respectively, compared to healthy subjects. Similarly, elimination half-life was about 1.8-fold and 2.6-fold longer, respectively, compared to healthy subjects. There is a linear relationship between creatinine clearance values and total apparent plasma clearance, renal clearance and plasma half-life of acamprosate. A dose of 1 x 333 mg acamprosate calcium, three times daily, is recommended in patients with moderate renal impairment (creatinine clearance of 30 to 50 mL/min [see Use in Specific Populations (8.6)].

Acamprosate calcium is contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Dosage and Administration (2.1), Contraindications (4.2), Warnings and Precautions (5.1) and Use in Specific Populations (8.6)].

**Hepatic Impairment**

Acamprosate is not metabolized by the liver and the pharmacokinetics of acamprosate calcium are not altered in patients with mild to moderate hepatic impairment (groups A and B of the Child-Pugh classification). No adjustment of dosage is recommended in such patients.

**Alcohol-dependent Subjects**

A cross-study comparison of acamprosate calcium at doses of 2 x 333 mg three times daily indicated similar pharmacokinetics between alcohol-dependent subjects and healthy subjects.

**Drug-drug Interactions**

Acamprosate had no inducing potential on the cytochrome CYP1A2 and 3A4 systems, and in vitro inhibition studies suggest that acamprosate does not inhibit in vivo metabolism mediated by cytochrome CYP1A2, 2C9, 2C19, 2D6, 2E1 or 3A4. The pharmacokinetics of acamprosate calcium were unaffected when coadministered with alcohol, disulfiram or diazepam. Similarly, the pharmacokinetics of ethanol, diazepam and nordiazepam, imipramine and desipramine, naltrexone and 6-beta naltrexol were unaffected.
following coadministration with acamprosate calcium. However, coadministration of acamprosate calcium with naltrexone led to a 33% increase in the $C_{\text{max}}$ and a 25% increase in the AUC of acamprosate. No adjustment of dosage is recommended in such patients.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Dietary administration of acamprosate calcium for 2 years to Sprague-Dawley rats at doses of 25, 100 and 400 mg/kg/day (up to 3 times the maximum recommended human daily (MRHD) oral dose on an AUC basis) and CD-1 mice at doses of 400, 1200 and 3600 mg/kg/day (up to 25 times the MRHD on an AUC basis) showed no evidence of increased tumor incidence.

Acamprosate calcium was negative in all genetic toxicology studies conducted. Acamprosate calcium demonstrated no evidence of genotoxicity in an in vitro bacterial reverse point mutation assay (Ames assay) or an in vitro mammalian cell gene mutation test using Chinese Hamster Lung V79 cells. No clastogenicity was observed in an in vitro chromosomal aberration assay in human lymphocytes and no chromosomal damage detected in an in vivo mouse micronucleus assay.

Acamprosate calcium had no effect on fertility after treatment for 70 days prior to mating in male rats and for 14 days prior to mating, throughout mating, gestation and lactation in female rats at doses up to 1000 mg/kg/day (approximately 4 times the MRHD oral dose on a mg/m$^2$ basis). In mice, acamprosate calcium administered orally for 60 days prior to mating and throughout gestation in females at doses up to 2400 mg/kg/day (approximately 5 times the MRHD oral dose on a mg/m$^2$ basis) had no effect on fertility.

14 CLINICAL STUDIES

The efficacy of acamprosate in the maintenance of abstinence was supported by three clinical studies involving a total of 998 patients who were administered at least one dose of acamprosate calcium or placebo as an adjunct to psychosocial therapy. Each study was a double-blind, placebo-controlled trial in alcohol-dependent patients who had undergone inpatient detoxification and were abstinent from alcohol on the day of randomization. Study durations ranged from 90 days to 360 days. Acamprosate calcium proved superior to placebo in maintaining abstinence, as indicated by a greater percentage of subjects being assessed as continuously abstinent throughout treatment.

In a fourth study, the efficacy of acamprosate calcium was evaluated in alcoholics, including patients with a history of polysubstance abuse and patients who had not undergone detoxification and were not required to be abstinent at baseline. This study failed to demonstrate superiority of acamprosate calcium over placebo.

16 HOW SUPPLIED/STORAGE AND HANDLING

Acamprosate Calcium Delayed-release Tablets are available containing 333 mg of acamprosate calcium.

The 333 mg tablet is a white enteric-coated, round, unscored tablet imprinted with M over AC in black ink on one side of the tablet and blank on the other side. They are available as follows:

NDC 0378-6333-80
bottles of 180 tablets

NDC 0378-6333-05
bottles of 500 tablets

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 using a child-resistant closure.

17 PATIENT COUNSELING INFORMATION

17.1 Information for Patients

Physicians are advised to discuss the following issues with patients for whom they prescribe acamprosate calcium delayed-release tablets.

Renal Impairment

A lower dose is recommended for patients with moderate renal impairment. Acamprosate calcium delayed-release tablets are contraindicated in patients with severe renal impairment (creatinine clearance of ≤ 30 mL/min) [see Dosage and Administration (2.1), Contraindications (4.2), Warnings and Precautions (5.1) and Use in Specific Populations (8.6)].

Suicidality and Depression

Families and caregivers of patients being treated with acamprosate calcium delayed-release tablets should be alerted to the need to monitor patients for the emergence of symptoms of depression or suicidality, and to report such symptoms to the patient's health care provider [see Warnings and Precautions (5.2)].

Alcohol Withdrawal

Use of acamprosate calcium delayed-release tablets does not eliminate or diminish withdrawal symptoms [see Warnings and Precautions (5.3)].

Pregnancy and Breast-feeding

• Advise patients to notify their physician if they become pregnant or intend to become pregnant during therapy.
• Advise patients to notify their physician if they are breast-feeding.

Relapse to Drinking

• Advise patients to continue acamprosate calcium delayed-release tablets therapy as directed, even in the event of relapse and remind them to discuss any renewed drinking with their physicians.
• Advise patients that acamprosate calcium delayed-release tablets has been shown to help maintain abstinence only when used as a part of a treatment program that includes counseling and support.
Each enteric-coated tablet contains
333 mg of acamprosate calcium
(equivalent to 300 mg of acamprosate).

Usual Dosage: See accompanying
prescribing information.

Keep this and all medication out of
the reach of children.

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

Manufactured for:
Mylan Pharmaceuticals Inc.
Morgantown, WV 26505 U.S.A.

Made in India

Mylan.com

RMX6333SS3

Dispense in a tight, light-resistant
container as defined in the USP
using a child-resistant closure.

Keep container tightly closed.

Code No.: MH/DRUGS/25/NKD/89

ACAMPROSATE CALCIUM
acamprosate calcium enteric-coated tablet, delayed release
# Product Information

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<thead>
<tr>
<th>Product Type</th>
<th>HUMAN PRESCRIPTION DRUG</th>
<th>Item Code (Source)</th>
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<tr>
<td>Route of Administration</td>
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## Active Ingredient/Active Moiety

<table>
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<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tr>
<td>ACAMPROSATE CALCIUM (UNII: 59375N1D0U) (ACAMPROSATE - UNII:N4K14YGM3J)</td>
<td>ACAMPROSATE CALCIUM</td>
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## Inactive Ingredients

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<td>SILICON DIOXIDE (UNII: ETJ7Z6XBU4)</td>
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<td>GLYCERYL DIBEHENATE (UNII: R8WTH25YS2)</td>
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<td>HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)</td>
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<td>MAGNESIUM STEARATE (UNII: 70097M6E0)</td>
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<td>PROPYLENE GLYCOL (UNII: 6DC9Q167V3)</td>
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<td>TALC (UNII: 7SEV7J4RIU)</td>
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## Product Characteristics

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## Packaging

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<tbody>
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<td>180 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product</td>
<td>09/24/2014</td>
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## Marketing Information

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<tr>
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<th>Application Number or Monograph Citation</th>
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<tr>
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**Labeler** - Mylan Pharmaceuticals Inc. (059295980)