CAMPRAL- acamprosate calcium tablet, delayed release
Carilion Materials Management

HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use CAMPRAL safely and effectively. See full prescribing information for CAMPRAL. CAMPRAL® (acamprosate calcium) Delayed-Release Tablets Initial U.S. Approval: 2004

INDICATIONS AND USAGE

- Campral® is indicated for the maintenance of abstinence from alcohol in patients with alcohol dependence who are abstinent at treatment initiation (,). 114
- Treatment with Campral 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 30-50 mL/min) ( ). 2.1
- Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) (, , , ,). 2.14.25.18.612.3

DOSAGE FORMS AND STRENGTHS

Enteric-coated tablets, 333 mg ( ). 3

CONTRAINDICATIONS

- Campral is contraindicated in patients who previously have exhibited hypersensitivity to acamprosate calcium or any of its components ( ). 4.1
- Campral 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 Campral 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 Forest Laboratories, Inc. at 1-800-678-1605, or FDA at 1-800-FDA-1088 or*www.fda.gov/medwatch***

USE IN SPECIFIC POPULATIONS

- Pregnancy: Campral 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 Campral is administered to a nursing woman ( ). 8.3
- Renal Impairment: Dose reduction required for moderate renal impairment; contraindicated in severe renal impairment (, , , ,) 2.14.25.18.612.3

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 8/2014

FULL PRESCRIBING INFORMATION: CONTENTS*

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

Campral® is indicated for the maintenance of abstinence from alcohol in patients with alcohol dependence who are abstinent at treatment initiation. Treatment with Campral should be part of a comprehensive management program that includes psychosocial support.

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

2 DOSAGE AND ADMINISTRATION

The recommended dose of Campral 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 Campral 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. Campral 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 30-50 mL/min), a starting dose of one 333 mg tablet taken three times daily is recommended. Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) [and]. *see Contraindications ( ), Warnings and Precautions ( ), Use in Specific Populations ( ), 4.25.18.6Clinical Pharmacology ( ) 12.3*

3 DOSAGE FORMS AND STRENGTHS

Campral 333 mg tablets are enteric-coated, white, round, biconvex tablets, identified with “333” debossed on one side.

4 CONTRAINDICATIONS

4.1 Hypersensitivity to Acamprosate Calcium

Campral is contraindicated in patients who previously have exhibited hypersensitivity to acamprosate calcium or any of its components.

4.2 Severe Renal Impairment

Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) [and]. *see Dosage and Administration ( ), Warnings and Precautions ( ), Use in Specific Populations ( ), 2.15.18.6Clinical Pharmacology ( ) 12.3*

5 WARNINGS AND PRECAUTIONS

5.1 Renal Impairment

Treatment with Campral in patients with moderate renal impairment (creatinine clearance of 30-50 mL/min) requires a dose reduction [ ]. Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) [and]. *see Dosage and Administration ( ) 2.1see Dosage and Administration ( ), Contraindications ( ), Use in Specific Populations ( ), 2.14.28.6Clinical Pharmacology ( ) 12.3*

5.2 Suicidality and Depression

In controlled clinical trials of Campral, adverse events of a suicidal nature (suicidal ideation, suicide attempts, completed suicides) were infrequent overall, but were more common in Campral-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 2272 (0.13%) patients in the pooled acamprosate group from all controlled studies and 2 of 1962 patients (0.10%) in the placebo group. Adverse events coded as "depression" were reported at similar rates in Campral-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 Campral, should be monitored for the development of symptoms of depression or suicidal thinking. Families and caregivers of patients being treated with Campral 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 Campral 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 Campral described elsewhere in labeling include suicidality and depression and acute kidney failure [s , and]. *ee Warnings and Precautions ( ) 5.2Adverse Reactions ( ) 6.2*

The adverse event data described below reflect the safety experience in over 7000 patients exposed to Campral for up to one year, including over 2000 Campral-exposed patients who participated in placebo-controlled trials.

Adverse Events Leading to Discontinuation

In placebo-controlled trials of 6 months or less, 8% of Campral-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 Campral-treated patients. Only diarrhea was associated with the discontinuation of more than 1% of patients (2% of Campral-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 Campral-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. shows those events that occurred in any Campral 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

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

Body System/Preferred Term	Number of Patients (%) with Events			
	Campral 1332 mg/day	Campral 1998 mg/day [1]	Campral Pooled [2]	Placebo
Number of patients in Treatment Group	397	1539	2019	1706
Number (%) of patients with an AE	248 (62%)	910 (59%)	1231 (61%)	955 (56%)

Body as a Whole	**121 (30%)**	**513 (33%)**	**685 (34%)**	**517 (30%)**
Accidental Injury*†	17 (4%)	44 (3%)	70 (3%)	52 (3%)
Asthenia	29 (7%)	79 (5%)	114 (6%)	93 (5%)
Pain	6 (2%)	56 (4%)	65 (3%)	55 (3%)
Digestive System	**85 (21%)**	**440 (29%)**	**574 (28%)**	**344 (20%)**
Anorexia	20 (5%)	35 (2%)	57 (3%)	44 (3%)
Diarrhea	39 (10%)	257 (17%)	329 (16%)	166 (10%)
Flatulence	4 (1%)	55 (4%)	63 (3%)	28 (2%)
Nausea	11 (3%)	69 (4%)	87 (4%)	58 (3%)
Nervous System	**150 (38%)**	**417 (27%)**	**598 (30%)**	**500 (29%)**
Anxiety††**	32 (8%)	80 (5%)	118 (6%)	98 (6%)
Depression	33 (8%)	63 (4%)	102 (5%)	87 (5%)
Dizziness	15 (4%)	49 (3%)	67 (3%)	44 (3%)
Dry mouth	13 (3%)	23 (1%)	36 (2%)	28 (2%)
Insomnia	34 (9%)	94 (6%)	137 (7%)	121 (7%)
Paresthesia	11 (3%)	29 (2%)	40 (2%)	34 (2%)
Skin and Appendages	**26 (7%)**	**150 (10%)**	**187 (9%)**	**169 (10%)**
Pruritus	12 (3%)	68 (4%)	82 (4%)	58 (3%)
Sweating	11 (3%)	27 (2%)	40 (2%)	39 (2%)

†*includes events coded as “fracture” by sponsor; ††**includes events coded as “nervousness” by sponsor

includes 258 patients treated with acamprosate calcium 2000 mg/day, using a different dosage strength and regimen. [1]

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. [2]

Concomitant Therapies

In clinical trials, the safety profile in subjects treated with Campral 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 Campral 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 Campral

Following is a list of terms that reflect treatment-emergent adverse events reported by patients treated with Campral in 20 clinical trials (4461 patients treated with Campral, 3526 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 1/1000 patients; rare events are those occurring in fewer than 1/1000 patients.

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

: palpitation, syncope; : hypotension, tachycardia, hemorrhage, angina pectoris, migraine, varicose vein,

myocardial infarct, phlebitis, postural hypotension; : heart failure, mesenteric arterial occlusion, cardiomyopathy, deep thrombophlebitis, shock. **Cardiovascular System –***FrequentInfrequentRare*

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

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

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

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

– myalgia, arthralgia; : leg cramps; rheumatoid arthritis, myopathy. **Musculoskeletal System –***FrequentInfrequentRare:*

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

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

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

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

: impotence; – metrorrhagia, urinary frequency, urinary tract infection, sexual function abnormal, urinary incontinence, vaginitis; kidney calculus, abnormal ejaculation, hematuria, menorrhagia, nocturia, polyuria, urinary urgency. **Urogenital System –***FrequentInfrequentRare:*

6.2 Postmarketing Experience

The following adverse reactions have been identified during post approval use of Campral. 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-US Postmarketing Evaluation of Campral (acamprosate calcium)

The serious adverse event of acute kidney failure has been reported to be temporally associated with Campral treatment in at least 3 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

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/m basis) and in rabbits when given in doses that are approximately 3 times the human dose (on a mg/m 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/m 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/m 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/m 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/m 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. Campral should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. **Teratogenic effects:**[222222]

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/m basis). No effects were observed at a dose of 320 mg/kg/day (approximately one-half the MRHD dose on a mg/m basis). **Nonteratogenic effects:**[22]

8.2 Labor and Delivery

The potential for Campral 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 Campral is administered to a nursing woman.

8.4 Pediatric Use

The safety and efficacy of Campral have not been established in the pediatric population.

8.5 Geriatric Use

Forty-one of the 4234 patients in double-blind, placebo-controlled, clinical trials of Campral 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 and]. *[see Clinical Pharmacology ( ), Adverse Reactions ( ), 12.36.1Dosage and Administration ( ) 2.1*

8.6 Renal Impairment

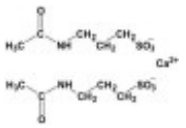
Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) [s and]. *ee Dosage and Administration ( ), Contraindications ( ), Warnings and Precautions ( ), 2.14.25.1Clinical Pharmacology ( ) 12.3*

10 OVERDOSAGE

In all reported cases of acute overdosage with Campral (total reported doses of up to 56 grams of acamprosate calcium), the only symptom that could be reasonably associated with Campral 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

Campral (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 calcium acetylaminopropane sulfonate. Its chemical formula is C H N O S Ca and molecular weight is 400.48. Its structural formula is: 1020282



Acamprosate calcium is a white, odorless or nearly odorless powder. It is freely soluble in water, and practically insoluble in absolute ethanol and dichloromethane.

Each Campral tablet contains acamprosate calcium 333 mg, equivalent to 300 mg of acamprosate. Inactive ingredients in Campral tablets include: crospovidone, microcrystalline cellulose, magnesium silicate, sodium starch glycolate, colloidal anhydrous silica, magnesium stearate, talc, propylene glycol and Eudragit L 30 D or equivalent. 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. and 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. *In vitroin vivo*

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. Campral 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 Campral abuse or dependence.

Campral 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 Campral 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 Campral doses of 2 x 333 mg tablets three times daily average 350 ng/mL and occur at 3-8 hours post-dose. Coadministration of Campral with food decreases bioavailability as measured by C and AUC, by approximately 42% and 23%, respectively. The food effect on absorption is not clinically significant and no adjustment of dose is necessary. $_{max}$

Distribution

The volume of distribution for acamprosate following intravenous administration is estimated to be 72-109 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 Campral, the terminal half-life ranges from approximately 20-33 hours. Following oral administration of Campral, the major route of excretion is via the kidneys as acamprosate.

Special Populations

Campral does not exhibit any significant pharmacokinetic differences between male and female subjects. ***Gender:***

The pharmacokinetics of Campral 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. ***Age:***

The pharmacokinetics of Campral have not been evaluated in a pediatric population. ***Pediatrics:***

: Peak plasma concentrations after administration of a single dose of 2 x 333 mg Campral 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 Campral, three times daily, is recommended in patients with moderate renal impairment (creatinine clearance of 30-50 mL/min, [ ]. ***Renal Impairment****see Use in Specific Populations ( ) 8.6*

Campral is contraindicated in patients with severe renal impairment (creatinine clearance of ≤30 mL/min) [and]. *see Dosage and Administration ( ), Contraindications ( ), Warnings and Precautions ( ), 2.14.25.1Use in Specific Populations ( ) 8.6*

: Acamprosate is not metabolized by the liver and the pharmacokinetics of Campral 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. ***Hepatic Impairment***

: A cross-study comparison of Campral at doses of 2 x 333 mg three times daily indicated similar

pharmacokinetics between alcohol-dependent subjects and healthy subjects. ***Alcohol-dependent subjects***

Drug-Drug Interactions

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

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 bacterial reverse point mutation assay (Ames assay) or an mammalian cell gene mutation test using Chinese Hamster Lung V79 cells. No clastogenicity was observed in an chromosomal aberration assay in human lymphocytes and no chromosomal damage detected in an mouse micronucleus assay. *in vitroin vitroin vitroin vivo*

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 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 basis) had no effect on fertility. [22]

14 CLINICAL STUDIES

The efficacy of Campral 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 Campral 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. Campral 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 Campral 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 Campral over placebo.

16 HOW SUPPLIED/STORAGE AND HANDLING

NDC:68151-4760-0 in a PACKAGE of 1 TABLET, DELAYED RELEASES

Storage and Handling

Store at 25ºC (77ºF); excursions permitted to 15º to 30ºC (59º to 86ºF).

17 PATIENT COUNSELING INFORMATION

17.1 Information for Patients

Physicians are advised to discuss the following issues with patients for whom they prescribe Campral.

Renal Impairment

A lower dose is recommended for patients with moderate renal impairment. Campral is contraindicated in patients with severe renal impairment (creatine clearance of ≤30 mL/min) [ ]. *see Dosage and Administration ( ), Contraindications ( ), Warnings and Precautions ( ) and Use in Specific Populations ( ) 2.14.25.18.6*

Suicidality and Depression

Families and caregivers of patients being treated with Campral 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 Campral 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 Campral therapy as directed, even in the event of relapse and remind them to discuss any renewed drinking with their physicians.
- Advise patients that Campral has been shown to help maintain abstinence only when used as a part of a treatment program that includes counseling and support.

Manufactured by: Merck Santé s.a.s. Subsidiary of Merck KGaA, Darmstadt, Germany 37, rue Saint-Romain 69008 LYON FRANCE

Manufactured for: Forest Pharmaceuticals, Inc. Subsidiary of Forest Laboratories, Inc. St. Louis, MO 63045

Campral

CAMPRAL

acamprosate calcium tablet, delayed release

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:68151-4760(NDC:0456-3330)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
acamprosate calcium (UNII: 59375N1D0U) (acamprosate - UNII:N4K14YGM3J)	acamprosate calcium	333 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CROSPOVIDONE (UNII: 68401960MK)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MAGNESIUM SILICATE (UNII: 9B9691B2N9)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TALC (UNII: 7SEV7J4R1U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)	

Product Characteristics			
Color	WHITE (WHITE)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	10mm
Flavor		**Imprint Code**	333
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:68151-4760-0	1 in 1 PACKAGE; Type 0: Not a Combination Product	01/11/2005	07/01/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021431	01/11/2005	

Labeler - Carilion Materials Management (079239644)

Registrant - Carilion Materials Management (079239644)

Establishment

Name	Address	ID/FEI	Business Operations
Carilion Materials Management		079239644	REPACK(68151-4760)

Revised: 8/2016 Carilion Materials Management