PRODUCT MONOGRAPH

Pr CADUET®

amlodipine besylate and atorvastatin calcium tablets

tablets 5/10 mg, 5/20 mg, 5/40 mg, 5/80 mg and
10/10 mg, 10/20 mg, 10/40 mg, 10/80 mg

Anti-hypertensive/Anti-anginal Agent and Lipid Metabolism Regulator

®C.P, Pharmaceuticals International C.V.
Pfizer Canada Inc., licensee
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CADUET®

amlodipine besylate and atorvastatin calcium tablets

PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	Nonmedicinal Ingredients
oral	Tablets (amlodipine/atorvastatin): 5/10 mg, 5/20 mg, 5/40 mg, 5/80 mg and 10/10 mg, 10/20 mg, 10/40 mg, 10/80 mg	Calcium Carbonate, Croscarmellose Sodium, Microcrystalline Cellulose, Pregelatinized Starch, Polysorbate 80, Hydroxypropyl Cellulose, Purified Water, Colloidal Silicon Dioxide (anhydrous), Magnesium Stearate, Opadry® II White 85F28751 or Opadry® II Blue 85F10919. ®Registered trademark of the Colorcon Company

INDICATIONS AND CLINICAL USE

CADUET (amlodipine /atorvastatin) is indicated in patients for whom treatment with both amlodipine and atorvastatin is appropriate, specifically, patients at cardiovascular risk.

CADUET is not for initial therapy. The dose of CADUET should be determined by the titration of individual components (see DOSAGE AND ADMINISTRATION).

CONTRAINDICATIONS

CADUET (amlodipine/atorvastatin) is contraindicated in patients with hypersensitivity to any component of this medication, the atorvastatin, amlodipine or other dihydropyridines*.
CADUET is contraindicated in patients with severe hypotension (less than 90 mmHg systolic) and in patients with active liver disease or unexplained persistent elevations of serum transaminases exceeding 3 times the upper limit of normal.

* Amlodipine is a dihydropyridine calcium channel blocker

CADUET is also contraindicated in pregnancy and for nursing women: Cholesterol and other products of cholesterol biosynthesis are essential components for fetal development (including synthesis of steroids and cell membranes). CADUET should be administered to women of childbearing age only when such patients are highly unlikely to conceive and have been informed of the possible harm. If the patient becomes pregnant while taking CADUET, the drug should be discontinued immediately and the patient apprised of the potential harm to the fetus. Atherosclerosis being a chronic process, discontinuation of lipid metabolism regulating drugs during pregnancy should have little impact on the outcome of long-term therapy of primary hypercholesterolemia (see WARNINGS AND PRECAUTIONS).

WARNINGS AND PRECAUTIONS

General

Before instituting therapy with CADUET (amlodipine/atorvastatin), an attempt should be made to control elevated serum lipoprotein levels with appropriate diet, exercise, and weight reduction in overweight patients, and to treat other underlying medical problems (see INDICATIONS AND CLINICAL USE). Patients should be advised to inform subsequent physicians of the prior use of atorvastatin or any other lipid-lowering agents.

Pharmacokinetic Interactions

The use of HMG CoA reductase inhibitors like some other lipid-lowering therapies has been associated with severe myopathy, including rhabdomyolysis, which may be more frequent when they are co-administered with drugs that inhibit the cytochrome P450 enzyme system. The atorvastatin component of CADUET is metabolized by cytochrome P450 isoform 3A4 and, as such, may interact with agents that inhibit this enzyme (see Muscle Effects; DRUG INTERACTIONS, CYTOCHROME P450-mediated Interactions).

Muscle Effects

Effects on skeletal muscle such as myalgia, myopathy and rarely, rhabdomyolysis have been reported in patients treated with the atorvastatin component of CADUET.

Rare cases of rhabdomyolysis with acute renal failure secondary to myoglobinuria, have been reported with the atorvastatin component of CADUET and with other HMG-CoA reductase inhibitors.

Myopathy, defined as muscle aching or muscle weakness in conjunction with increases in creatine kinase (CK) values to greater than 10 times the upper limit of normal, should be considered in any patient with diffuse myalgia, muscle tenderness or weakness, and/or marked elevation of CPK. Patients should be advised to report promptly unexplained muscle pain, tenderness or weakness, particularly if accompanied by malaise or fever. Patients who develop any signs or symptoms suggestive of myopathy should have their CK levels measured.

CADUET therapy should be discontinued if markedly elevated CPK levels occur or myopathy is diagnosed or suspected.

Pre-disposing Factors for Myopathy/Rhabdomyolysis: the atorvastatin component of CADUET, as with other HMG-CoA reductase inhibitors, should be prescribed with caution in patients with pre-disposing factors for myopathy/rhabdomyolysis. Such factors include:

- Personal or family history of hereditary muscular disorders
- Previous history of muscle toxicity with another HMG-CoA reductase inhibitor
- Concomitant use of a fibrate, or niacin
- Hypothyroidism
- Alcohol abuse
- Excessive physical exercise
- Age >65 years
- Renal impairment
- Hepatic impairment
- Diabetes with hepatic fatty change
- Surgery and trauma
- Frailty
- Situations where an increase in plasma levels of active ingredient may occur

The risk of myopathy and rhabdomyolysis during treatment with HMG-CoA reductase inhibitors is increased with concurrent administration of drugs that interfere with metabolism of atorvastatin via CYP 3A4, such as cyclosporine, fibric acid derivatives, erythromycin, clarithromycin, niacin (nicotinic acid), azole antifungals, nefazodone, colchicine, hepatitis C protease inhibitors telaprevir, boceprevir , HIV protease inhibitor fosamprenavir and each of the following HIV protease inhibitor combinations: saquinavir plus ritonavir, lopinavir plus ritonavir, tipranavir plus ritonavir, darunavir plus ritonavir and fosamprenavir plus ritonavir. The combined therapy with CADUET and cyclosporine, gemfibrozil, telaprevir or tipranavir plus ritonavir should be avoided. CADUET dose restriction or caution is recommended for combined therapy with other CYP 3A4 inhibitors (see Pharmacokinetic Interactions; DRUG INTERACTIONS, Drug-Drug Interactions; DETAILED PHARMACOLOGY, Human Pharmacokinetics).

Temporary suspension of atorvastatin during fusidic acid therapy is recommended (see DRUG INTERACTIONS, Drug-Drug Interactions).

Although patients with renal impairment are known to be predisposed to the development of rhabdomyolysis with administration of HMG-CoA reductase inhibitors (also known as statins), those with a history of renal impairment may also be predisposed to the development of rhabdomyolysis. Such patients merit close monitoring for skeletal muscle effects.

CADUET therapy should be temporarily withheld or discontinued in any patient with an acute serious condition suggestive of a myopathy or having a risk factor predisposing to the development of renal failure secondary to rhabdomyolysis (such as sepsis, severe acute

infection, hypotension, major surgery, trauma, severe metabolic, endocrine and electrolyte disorders, and uncontrolled seizures).

CADUET therapy should be discontinued if markedly elevated CPK levels occur or myopathy is diagnosed or suspected.

Cardiovascular

Hemorrhagic Stroke in Patients with Recent Stroke or Transient Ischemic Attack (TIA)

A post-hoc analysis of a clinical study in 4,731 patients without coronary heart disease (CHD) who had a stroke or TIA within the preceding six months revealed a higher incidence of hemorrhagic stroke in the atorvastatin 80mg group compared to placebo. Patients with hemorrhagic stroke on entry appeared to be at increased risk for recurrent hemorrhagic stroke. The potential risk of hemorrhagic stroke should be carefully considered before initiating treatment with atorvastatin in patients with recent (1-6 months) stroke or TIA.

Increased Angina and/or Myocardial Infarction

Rarely, patients, particularly those with severe obstructive coronary artery disease, have developed documented increased frequency, duration and/or severity of angina or acute myocardial infarction on starting calcium channel blocker therapy or at the time of dosage increase. The mechanism of this effect has not been elucidated.

Outflow Obstruction (Aortic Stenosis)

CADUET should be used with caution in the presence of fixed left ventricular outflow obstruction (aortic stenosis).

Use in Patients with Congestive Heart Failure

Although generally calcium channel blockers should only be used with caution in patients with heart failure, it has been observed that the amlodipine component of CADUET had no overall deleterious effect on survival and cardiovascular morbidity in both short-term and long-term clinical trials in these patients. While a significant proportion of the patients in these studies had a history of ischemic heart disease, angina or hypertension, the studies were not designed to evaluate the treatment of angina or hypertension in patients with concomitant heart failure.

Hypotension

The amlodipine component of CADUET may occasionally precipitate symptomatic hypotension. Careful monitoring of blood pressure is recommended, especially in patients with a history of cerebrovascular insufficiency, and those taking medications known to lower blood pressure.

Effect on Ubiquinone (CoQ_{10}) Levels

Significant decreases in circulating ubiquinone levels in patients treated with atorvastatin and other statins have been observed. The clinical significance of a potential long-term statin-induced deficiency of ubiquinone has not been established. It has been reported that a decrease in myocardial ubiquinone levels could lead to impaired cardiac function in patients with borderline congestive heart failure (see REFERENCES).

Drug/Laboratory Test Interactions

The atorvastatin component of CADUET may elevate serum transaminase and CPK levels (from skeletal muscle). In the differential diagnosis of chest pain in a patient on therapy with CADUET, cardiac and noncardiac fractions of these enzymes should be determined.

Beta-blocker Withdrawal

The amlodipine component of CADUET gives no protection against the dangers of abrupt beta-blocker withdrawal and such withdrawal should be done by the gradual reduction of the dose of beta-blocker.

Peripheral Edema

Mild to moderate peripheral edema was the most common adverse event in clinical trials with the amlodipine component of CADUET (see ADVERSE REACTIONS). The incidence of peripheral edema was dose-dependent and ranged in frequency from 3.0 to 10.8% in 5 to 10 mg dose range. Care should be taken to differentiate this peripheral edema from the effects of increasing left ventricular dysfunction.

Effect on Lipoprotein (a)

In some patients, the beneficial effect of lowered total cholesterol and LDL-C levels may be partly blunted by a concomitant increase in Lp (a) lipoprotein concentrations. Present knowledge suggests the importance of high Lap (a) levels as an emerging risk factor for coronary heart disease. It is thus desirable to maintain and reinforce lifestyle changes in high risk patients placed on atorvastatin therapy (see REFERENCES).

Patients with Severe Hypercholesterolemia

Higher drug dosages (80 mg/day) required for some patients with severe hypercholesterolemia (including familial hypercholesterolemia) are associated with increased plasma levels of the atorvastatin component of CADUET. **Caution should be exercised in such patients who are also severely renally impaired, elderly, or are concomitantly being administered digoxin or CYP 3A4 inhibitors** (see Pharmacokinetic Interactions, Muscle Effects; DRUG INTERACTIONS; DOSAGE AND ADMINISTRATION).

Endocrine and Metabolism

Endocrine Function

HMG-CoA reductase inhibitors interfere with cholesterol synthesis and, as such, might theoretically blunt adrenal and/or gonadal steroid production. Clinical studies with the atorvastatin component of CADUET and other HMG-CoA reductase inhibitors have suggested that these agents do not reduce plasma cortisol concentration or impair adrenal reserve, and do not reduce basal plasma testosterone concentration. However, the effects of HMG-CoA reductase inhibitors on male fertility have not been studied in adequate numbers of patients. The effects, if any, on the pituitary-gonadal axis in premenopausal women are unknown.

Patients treated with the atorvastatin component of CADUET who develop clinical evidence of endocrine dysfunction should be evaluated appropriately. Caution should be exercised if an HMG-CoA reductase inhibitor or other agent used to lower cholesterol levels is administered to patients receiving other drugs (e.g. ketoconazole, spironolactone or cimetidine) that may decrease the levels of endogenous steroid hormones.

Increases in fasting glucose and HbA1c levels have been reported with inhibitors of HMG-CoA reductase as a class. For some patients, at high risk of diabetes mellitus, hyperglycemia was sufficient to shift them to the diabetes status. The benefit of treatment continues to outweigh the small increased risk. Periodic monitoring of these patients is recommended.

Hepatic/Biliary/Pancreatic

Hepatic Effects

In clinical trials with the atorvastatin component of CADUET, persistent increases in serum transaminases greater than 3 times the upper limit of normal occurred in <1% of patients who received atorvastatin. When the dosage of atorvastatin was reduced, or when drug treatment was interrupted or discontinued, serum transaminase levels returned to pretreatment levels. The increases were generally not associated with jaundice or other clinical signs or symptoms. Most patients continued treatment with a reduced dose of atorvastatin without clinical sequelae. **If increases in alanine aminotransferase (ALT) or aspartate aminotransferase (AST) show evidence of progression, particularly if they rise to greater than 3 times the upper limit of normal and are persistent, the dosage of the atorvastatin component of CADUET should be reduced or the drug discontinued.**

Liver function tests should be performed before the initiation of treatment, and repeated as clinically indicated. **There have been rare postmarketing reports of fatal and non-fatal hepatic failure in patients taking statins, including atorvastatin. If serious liver injury with clinical symptoms and/or hyperbilirubinemia or jaundice occurs during treatment with CADUET, promptly interrupt therapy. If an alternate etiology is not found, do not restart CADUET**.

CADUET, as well as other products containing HMG-CoA reductase inhibitors, should be used with caution in patients who consume substantial quantities of alcohol and/or have a past history of liver disease. Active liver disease or unexplained transaminase elevations are contraindications to the use of the atorvastatin component of CADUET; if such a condition should develop during therapy, CADUET should be discontinued.

There are no adequate studies in patients with liver dysfunction and dosage recommendations have not been established. In a small number of patients with mild to moderate hepatic impairment in which a single dose of 5 mg of the amlodipine component of CADUET was given, half-life has been prolonged (see ACTION AND CLINICAL PHARMACOLOGY, Pharmacokinetics and Metabolism). CADUET should therefore be administered with caution in these patients and careful monitoring should be performed. A lower starting dose of the amlodipine component of CADUET may be required (see DOSAGE AND ADMINISTRATION).

Ophthalmologic

Effect on the Lens

Current long-term data from clinical trials do not indicate an adverse effect of the atorvastatin component of CADUET on the human lens.

Renal

Renal Insufficiency

Plasma concentrations and LDL-C lowering efficacy of the atorvastatin component of CADUET were shown to be similar in patients with moderate renal insufficiency compared with patients with normal renal function. However, since several cases of rhabdomyolysis have been reported in patients with a history of renal insufficiency of unknown severity, as a precautionary measure and pending further experience in renal disease, the lowest dose (10 mg/day) of atorvastatin should be used in these patients. Similar precautions apply in patients with severe renal insufficiency (creatinine clearance <30 mL/min [<0.5 mL/sec]); the lowest dosage should be used and implemented cautiously (see Muscle Effects, DRUG INTERACTIONS; DOSAGE AND ADMINISTRATION).

Hypersensitivity

An apparent hypersensitivity syndrome has been reported with other HMG-CoA reductase inhibitors which has included 1 or more of the following features: anaphylaxis, angioedema, lupus erythematous-like syndrome, polymyalgia rheumatica, vasculitis, purpura, thrombocytopenia, leukopenia, hemolytic anemia, positive ANA, ESR increase, eosinophilia, arthritis, arthralgia, urticaria, asthenia, photosensitivity, fever, chills, flushing, malaise, dyspnea, toxic epidermal necrolysis, erythema multiforme, including Stevens-Johnson syndrome. Although to date hypersensitivity syndrome has not been described as such with CADUET, CADUET should be discontinued if hypersensitivity is suspected.

Concomitant Use with Strong Inhibitors of CYP 3A4
Use of CADUET with drugs that result in strong inhibition of CYP 3A4, such as ketoconazole, clarithromycin, ritonavir, may lead to increased plasma levels of amlodipine and associated serious adverse events (see DRUG INTERACTIONS). Such concomitant use should be avoided.

An observational study demonstrated an increased risk of hospitalization with acute kidney injury when amlodipine was used concomitantly with clarithromycin in elderly patients (>65 years of age) compared to when it was used concomitantly with azithromycin, odds ratio [amlodipine: 1.61 (95% C.I. 1.29-2.02)].

Special Populations

Pregnant Women

The atorvastatin component of CADUET is contraindicated during pregnancy (see CONTRAINDICATIONS).

There are no data on the use of atorvastatin during pregnancy. CADUET should be administered to women of childbearing age only when such patients are highly unlikely to conceive and have been informed of the potential hazards. If the patient becomes pregnant while taking CADUET, the drug should be discontinued and the patient apprised of the potential risk to the fetus.

Although amlodipine was not teratogenic in the rat and rabbit, some dihydropyridine compounds have been found to be teratogenic in animals. In rats, amlodipine has been shown to prolong both the gestation period and the duration of labor. There was no effect on the fertility of rats treated with amlodipine (see TOXICOLOGY, Reproduction and Teratology). There is no clinical experience with amlodipine in pregnant women.

Nursing Women

It is not known whether the amlodipine component of CADUET is excreted in human milk. In rats, milk concentrations of atorvastatin are similar to those in plasma. It is not known whether the atorvastatin component of CADUET is excreted in human milk. Because of the potential for adverse reactions in nursing infants, women taking CADUET should not breast-feed (see CONTRAINDICATIONS).

Pediatrics

There have been no studies conducted to determine the safety or efficacy of amlodipine/atorvastatin (combination product) in pediatric patients. However, there have been studies with pediatrics with amlodipine alone and atorvastatin alone (see below).

Amlodipine
The effect of amlodipine on blood pressure in patients less than 6 years of age is not known. Pediatric safety and efficacy studies beyond 8 weeks of duration have not been conducted. Please refer to the Product Monograph for NORVASC (amlodipine).

Atorvastatin
Safety and effectiveness of atorvastatin in patients 10-17 years of age (N=140) with heterozygous familial hypercholesterolemia have been evaluated in a controlled clinical trial of 6 months duration in adolescent boys and postmenarchal girls. Patients treated with atorvastatin had a safety and tolerability profile generally similar to that of placebo. Doses greater than 20 mg have not been studied in this patient population. Please refer to the Product Monograph for LIPITOR (atorvastatin).
Safety and effectiveness of atorvastatin in pediatric patients has not been determined in the prevention of myocardial infarction. Please refer to the Product Monograph for LIPITOR (atorvastatin).

Geriatrics

Amlodipine
In elderly patients (>65 years), clearance of amlodipine is decreased with a resulting increase in AUC of approximately 40-60%. In general, dose selection of the amlodipine component of CADUET for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy (see ACTION AND CLINICAL PHARMACOLOGY, Pharmacokinetics and Metabolism). In clinical trials, the incidence of adverse reactions in elderly patients was approximately 6% higher than that of younger population (<65 years). Adverse reactions include edema, muscle cramps and dizziness. The amlodipine component of CADUET should be used cautiously in elderly patients. Dosage adjustment is advisable (see DOSAGE AND ADMINISTRATION).

Atorvastatin
Treatment experience in adults 70 years or older (N=221) with doses of atorvastatin up to 80 mg/day has demonstrated that the safety and effectiveness of atorvastatin in this population was similar to that of patients <70 years of age. Pharmacokinetic evaluation of atorvastatin in subjects over the age of 65 years indicates an increased AUC. As a precautionary measure, the lowest dose of the atorvastatin component of CADUET should be administered initially (see DETAILED PHARMACOLOGY, Human Pharmacokinetics; REFERENCES).

Elderly patients may be more susceptible to myopathy (see above, Muscle Effects – Pre-disposing Factors for Myopathy/Rhabdomyolysis).

ADVERSE REACTIONS

Clinical Trial Adverse Drug Reactions

For CADUET, itself

CADUET (amlodipine /atorvastatin) has been evaluated for safety in 1,092 patients in two double-blind, placebo-controlled studies treated for co-morbid hypertension and dyslipidemia. In these studies, no unexpected adverse experiences particular to this combination have been observed. Adverse experiences have been limited to those that were reported previously with amlodipine and/or atorvastatin.

For the most part, adverse experiences with CADUET have been mild or moderate in severity. In these controlled clinical trials, adverse events or laboratory abnormalities leading to discontinuation occurred in 5.1% of patients treated with both amlodipine and atorvastatin compared to 4.0% of patients given placebo. The most common safety-related reasons for discontinuation from these studies in the combination treatment groups were headache and peripheral edema.

In a double-blind, controlled clinical trial of all available CADUET doses (5/10 mg to 10/80 mg amlodipine/atorvastatin respectively), the incidences of treatment-emergent adverse events (all causalities) that occurred in at least 1% of all combination treatment groups, pooled across all the combination doses, are summarized below.

Table 1 - Adverse Events (All Causality) > 1% of Patients taking Concurrent Amlodipine and Atorvastatin

Body System COSTART Preferred Term	Placebo N = 111(%)	AML Only N = 221 (%)	ATO Only N = 443 (%)	AML + ATO N = 885 (%)
Body as a whole /	**16 (14.4)**	**28 (12.7)**	**69 (15.6)**	**137 (15.5)**
Abdominal pain	0 (0.0)	2 (0.9)	10 (2.3)	20 (2.3)
Asthenia	3 (2.7)	4 (1.8)	8 (1.8)	19 (2.1)
Back pain	1 (0.9)	4 (1.8)	5 (1.1)	15 (1.7)
Flu syndrome	1 (0.9)	0 (0.0)	8 (1.8)	9 (1.0)
Headache	11 (9.9)	11 (5.0)	34 (7.7)	47 (5.3)
Cardiovascular /	**8 (7.2)**	**16 (7.2)**	**26 (5.9)**	**67 (7.6)**
Palpitation	2 (1.8)	4 (1.8)	4 (0.9)	17 (1.9)
Vasodilatation	3 (2.7)	2 (0.9)	3 (0.7)	18 (2.0)
Digestive /	**10 (9.0)**	**16 (7.2)**	**39 (8.8)**	**77 (8.7)**
Constipation	1 (0.9)	3 (1.4)	2 (0.5)	15 (1.7)
Diarrhea	2 (1.8)	2 (0.9)	5 (1.1)	17 (1.9)
GGT increased	0 (0.0)	1 (0.5)	6 (1.4)	16 (1.8)
Nausea	3 (2.7)	3 (1.4)	7 (1.6)	9 (1.0)
Metabolic and nutritional /	**6 (5.4)**	**32 (14.5)**	**21 (4.7)**	**133 (15.0)**
Alkaline phosphatase increased	0 (0.0)	0 (0.0)	2 (0.5)	10 (1.1)
Hyperglycemia	0 (0.0)	1 (0.5)	4 (0.9)	10 (1.1)
Peripheral edema	3 (2.7)	27 (12.2)	5 (1.1)	88 (9.9)
SGOT increased	1 (0.9)	1 (0.5)	3 (0.7)	13 (1.5)
SGPT increased	0 (0.0)	1 (0.5)	5 (1.1)	15 (1.7)
Musculoskeletal /	**7 (6.3)**	**12 (5.4)**	**25 (5.6)**	**35 (4.0)**

Body System COSTART Preferred Term	Placebo N = 111(%)	AML Only N = 221 (%)	ATO Only N = 443 (%)	AML + ATO N = 885 (%)
Arthralgia	4 (3.6)	3 (1.4)	4 (0.9)	10 (1.1)
Myalgia	2 (1.8)	3 (1.4)	8 (1.8)	14 (1.6)
Nervous	**9 (8.1)**	**12 (5.4)**	**25 (5.6)**	**47 (5.3)**
Dizziness	3 (2.7)	7 (3.2)	5 (1.1)	21 (2.4)
Respiratory /	**9 (8.1)**	**12 (5.4)**	**28 (6.3)**	**69 (7.8)**
Pharyngitis	1 (0.9)	1 (0.5)	3 (0.7)	9 (1.0)
Respiratory tract infection	5 (4.5)	7 (3.2)	17 (3.8)	43 (4.9)
Skin and appendages /	**4 (3.6)**	**4 (1.8)**	**6 (1.4)**	**32 (3.6)**
Rash	1 (0.9)	1 (0.5)	3 (0.7)	15 (1.7)

AML = amlodipine
ATO = atorvastatin

The incidence (%) of dose-related adverse events was consistent with those seen for amlodipine and/or atorvastatin.

In this clinical trial, the most frequently reported adverse events among patients who took concurrent amlodipine and atorvastatin were peripheral edema (9.9%), headache (5.3%), respiratory tract infection (4.9%), dizziness (2.4%), abdominal pain (2.3%), asthenia (2.1%), and vasodilatation (2.0%).

In this controlled clinical trial, similar percentages of patients who took concurrent amlodipine and atorvastatin (5.6%) versus patients who took placebo (4.5%), amlodipine only (5.4%), or atorvastatin only (4.1%) discontinued due to adverse safety experiences. Only 1 subject discontinued due to laboratory abnormalities. The most common safety-related reasons for discontinuation from the study in the combination treatment groups were peripheral edema (1.5%) and headache (1.4%), but these events led to the discontinuation of subjects in the combination treatment groups no more frequently than they did among subjects treated with either amlodipine alone or atorvastatin alone within this study.

The following information is based on the clinical experience with the parent compounds, NORVASC (amlodipine) and LIPITOR (atorvastatin).

Amlodipine

Amlodipine besylate has been administered to 1,714 patients (805 hypertensive and 909 angina patients) in controlled clinical trials, when compared to placebo alone or active comparators. Most adverse reactions reported during therapy were of mild to moderate severity.

Hypertension

In the 805 hypertensive patients treated with amlodipine in controlled clinical trials, adverse effects were reported in 29.9% of patients and required discontinuation of therapy due to side effects in 1.9% of patients. The most common adverse reactions in controlled clinical trials were: oedema (8.9%), and headaches (8.3%).

The following adverse reactions were reported with an incidence of >0.5% in the controlled clinical trials program (n=805):

Cardiovascular: oedema (8.9%), palpitations (2.0%), tachycardia (0.7%), postural dizziness (0.5%).

Skin and Appendages : pruritus (0.7%).

Musculoskeletal: muscle cramps (0.5%).

Central and Peripheral Nervous System : headaches (8.3%), dizziness (3.0%), paresthesia (0.5%)

Autonomic Nervous System: flushing (3.1%), increased sweating (0.9%), dry mouth (0.7%).

Psychiatric: somnolence (1.4%).

Gastrointestinal: nausea (2.4%), abdominal pain (1.1%), dyspepsia (0.6%), constipation (0.5%).

General: fatigue (4.1%), pain (0.5%).

Angina

In the controlled clinical trials in 909 angina patients treated with amlodipine, adverse effects were reported in 30.5% of patients and required discontinuation of therapy due to side effects in 0.6% of patients. The most common adverse reactions reported in controlled clinical trials were: oedema (9.9%) and headaches (7.8%).

The following adverse reactions occurred at an incidence of >0.5% in the controlled clinical trials program (n=909):

Cardiovascular: oedema (9.9%), palpitations (2.0%), postural dizziness (0.6%).

Skin and Appendages: rash (1.0%), pruritus (0.8%).

Musculoskeletal: muscle cramps (1.0%).

Central and Peripheral Nervous System: headaches (7.8%), dizziness (4.5%), paraesthesia

(1.0%), hypoaesthesia (0.9%)

Autonomic Nervous System : flushing (1.9%).

Psychiatric: somnolence (1.2%), insomnia (0.9%), nervousness (0.7%).

Gastrointestinal : nausea (4.2%), abdominal pain (2.2%), dyspepsia (1.4%), diarrhea (1.1%), flatulence (1.0%), constipation (0.9%).

Respiratory System: dyspnoea (1.1%).

Special Senses : visual impairment (1.3%), tinnitus (0.6%).

General : fatigue (4.8%), pain (1.0%), asthenia (1.0%).

Less Common Clinical Trial Adverse Drug Reactions

Amlodipine

Amlodipine has been evaluated for safety in about 11,000 patients with hypertension and angina. The following events occurred in <1% but >0.1% of patients in comparative clinical trials (double-blind comparative vs placebo or active agents; n = 2,615) or under conditions of open trials or marketing experience where a causal relationship is uncertain.

Cardiovascular: arrhythmia (including ventricular tachycardia and atrial fibrillation), bradycardia, myocardial infarction, hypotension, peripheral ischemia, syncope, tachycardia, postural dizziness, postural hypotension, vasculitis, chest pain.

Central and Peripheral Nervous System: hypoaesthesia/paraesthesia, neuropathy peripheral, tremor, vertigo.

Gastrointestinal: anorexia, constipation, dysphagia, vomiting, gingival hyperplasia, change in bowel habits, dyspepsia.

General : allergic reaction, asthenia*, back pain, pain, hot flushes, malaise, rigors, weight increased/weight decreased.

Musculoskeletal System: arthralgia, arthrosis, myalgia, muscle cramps.

Psychiatric : sexual dysfunction (male* and female), insomnia, nervousness, depression, abnormal dreams, anxiety, depersonalization, mood altered.

Respiratory System: dyspnoea, epistaxis.

Skin and Appendages : pruritus*, rash erythematous, rash maculopapular, erythema multiforme.

Special Senses: conjunctivitis, diplopia, eye pain, visual impairment, tinnitus.

Urinary System : pollakiuria, micturition disorder, nocturia.

Autonomic Nervous System: dry mouth, hyperhidrosis.

Metabolic and Nutritional : hyperglycaemia, thirst.

Hemopoietic: leukopenia, purpura, thrombocytopenia.

Reproductive system and breast disorders: gynecomastia, erectile dysfunction

*These events occurred in less than 1% in placebo-controlled trials, but the incidence of these side effects was between 1% and 2% in all multiple dose studies.

The following events occurred in <0.1% of patients: cardiac failure, skin discoloration*, urticaria*, skin dryness, Stevens-Johnson syndrome, alopecia*, twitching, ataxia, hypertonia*, migraine, apathy, amnesia, gastritis*, pancreatitis*, increased appetite, coughing*, rhinitis*, parosmia, taste perversion*, and xerophthalmia.
* these events were observed in marketing experience as well.

Isolated cases of angioedema have been reported. Angioedema may be accompanied by breathing difficulty.

Atorvastatin

Dyslipidemia

Adverse reactions with atorvastatin have usually been mild and transient. In the atorvastatin placebo-controlled clinical trial database of 16,066 (8,755 LIPITOR versus 7,311 placebo) patients treated for a median period of 53 weeks , 5.2% of patients on atorvastatin discontinued due to adverse reactions compared to 4.0% of the patients on placebo.

Adverse experiences occurring at an incidence ≥1% in patients participating in placebo-controlled clinical studies of atorvastatin and reported to be possibly, probably or definitely drug related are shown in Table 2 below:

Table 2 - Associated Adverse Events Reported in ≥1% of Patients in Placebo-Controlled Clinical Trials

	Atorvastatin % (n=8755)	Placebo % (n=7311)
Gastrointestinal disorders:		
Diarrhea	6.8	6.3
Dyspepsia	4.6	4.3
Nausea	4.0	3.5
Constipation	3.9	4.3
Flatulence	1.2	1.0
General disorders and administration site conditions:		
Asthenia	1.1	1.1
Infections and Infestations:		
Nasopharyngitis	8.3	8.2
Metabolism and nutrition disorders:		
Liver function test abnormal*	4.1	2.0
Blood creatine phosphokinase increased	1.9	1.8
Hyperglycemia	5.9	5.5
Musculoskeletal and connective tissue disorders:		
Arthralgia	6.9	6.5
Pain in extremity	6.0	5.9
Musculoskeletal pain	3.8	3.6
Muscle spasms	3.6	3.0
Myalgia	3.5	3.1
Joint swelling	1.3	1.2
Nervous system disorders		

	Atorvastatin % (n=8755)	Placebo % (n=7311)
Headache	6.5	6.7
Respiratory, thoracic and mediastinal disorders:		
Pharyngolaryngeal pain	2.3	2.1
Epistaxis	1.2	1.1

*alanine aminotransferase increased, aspartate aminotransferase increased, blood bilirubin increased, hepatic enzyme increased, liver function test abnormal and transaminases increased.

Less Common Clinical Trial Adverse Drug Reactions

Atorvastatin

The following additional adverse events were reported in placebo-controlled clinical trials during atorvastatin therapy: Muscle cramps, myositis, muscle fatigue, myopathy, paresthesia, peripheral neuropathy, pancreatitis, hepatitis, cholestatic jaundice, cholestasis, anorexia, vomiting, abdominal discomfort, alopecia, pruritus, rash, urticaria, impotence, nightmare, vision blurred, tinnitus, eructation, neck pain, malaise, pyrexia and white blood cells urine positive.

In summary, the adverse events occurring at a frequency <1% are listed below:

General disorders and administration site conditions: malaise; pyrexia
Gastrointestinal disorders: abdominal discomfort, eructation
Hepatobiliary disorders: hepatitis, cholestasis
Musculoskeletal and connective tissue disorders: muscle fatigue, neck pain
Psychiatric disorders: nightmare
Skin and subcutaneous tissue disorders: urticaria
Eye disorders: vision blurred
Ear and labyrinth disorders: tinnitus
Investigations: white blood cells urine positive

Abnormal Hematologic and Clinical Chemistry Findings

Atorvastatin: Laboratory Tests: Increases in serum transaminase levels and serum glucose have been noted in clinical trials (see WARNINGS AND PRECAUTIONS, ADVERSE REACTIONS, Atorvastatin).

Post-Market Adverse Drug Reactions

Amlodipine

In post-marketing experience, jaundice and hepatic enzyme elevations (mostly consistent with cholestasis or hepatitis), in some cases severe enough to require hospitalization, have been reported in association with the use of amlodipine. Postmarketing reporting has also revealed cases of extrapyramidal disorders induced by amlodipine.

Atorvastatin

The following adverse events have also been reported during post-marketing experience with the atorvastatin component of CADUET, regardless of causality assessment:

Rare reports: severe myopathy with or without rhabdomyolysis (see WARNINGS AND PRECAUTIONS, Muscle Effects, Renal Insufficiency, DRUG INTERACTIONS).

There have been rare reports of immune-mediated necrotizing myopathy (IMNM), an autoimmune myopathy, associated with statin use. IMNM is characterized by: proximal muscle weakness and elevated serum creatine kinase, which persist despite discontinuation of statin treatment; muscle biopsy showing necrotizing myopathy without significant inflammation; improvement with immunosuppressive agents.

Isolated reports: Gynecomastia, thrombocytopenia, arthralgia and allergic reactions including urticaria, angioneurotic edema, anaphylaxis and bullous rashes (including erytheme multiforme, Stevens-Johnson syndrome and toxic epidermal necrolysis) and fatigue, back pain, chest pain, malaise, dizziness, amnesia, peripheral edema, weight gain, abdominal pain, insomnia, hypoesthesia, tinnitus, tendon rupture, pancreatitis and dysgeusia.

Ophthalmologic observations: see WARNINGS AND PRECAUTIONS, Ophthalmologic.

Cases of erectile dysfunction have been reported in association with the use of statins.

The following adverse events have been reported with some statins:
- Sleep disturbances, including insomnia and nightmares;
- Mood related disorders, including depression;
- Very rare cases of interstitial lung disease, especially with long term therapy. If it is suspected a patient has developed interstitial lung disease, statin therapy should be discontinued.

Endocrine disorders: Increases in fasting glucose and HbA1c levels have been reported with CADUET.

There have been rare post-marketing reports of cognitive impairment (e.g. memory loss, forgetfulness, amnesia, memory impairment, confusion) associated with statin use. These cognitive issues have been reported for all statins. The reports are generally non-serious and reversible upon statin discontinuation, with variable times to symptom onset (1 day to years) and symptom resolution (median of 3 weeks).

DRUG INTERACTIONS

Pharmacokinetic interaction studies conducted with drugs in healthy subjects may not detect the possibility of a potential drug interaction in some patients due to differences in underlying diseases and use of concomitant medications (see also WARNINGS AND PRECAUTIONS, Geriatric Use, Renal Insufficiency, Patients with Severe Hypercholesterolemia).

Data from a drug-drug interaction study involving 10 mg of amlodipine and 80 mg of atorvastatin in healthy subjects indicate that the pharmacokinetics of amlodipine are not altered when the drugs are coadministered. The effect of amlodipine on the pharmacokinetics of atorvastatin showed no effect on the C_{max} but the AUC of atorvastatin increased by 18% (90% confidence interval: 109 to 127%) in the presence of amlodipine.

No drug interaction studies have been conducted with CADUET (amlodipine/atorvastatin) and other drugs, although studies have been conducted in the individual amlodipine and atorvastatin components, as described below:

Cytochrome P-450 Mediated Interactions

Drugs known to be inhibitors of the cytochrome P450 system include: azole antifungals, cimetidine, cyclosporine, erythromycin, quinidine, warfarin, diltiazem.

Drugs known to be inducers of the cytochrome P450 system include: phenobarbital, phenytoin, rifampin, hypericum perforatum (St John's wort).

Drugs known to be biotransformed via the cytochrome P450 system include: benzodiazepines, flecainide, imipramine, propafenone, and theophylline.

Amlodipine: As with all drugs, care should be exercised when treating patients with multiple medications. Dihydropyridine calcium channel blockers undergo biotransformation by the cytochrome P450 system, mainly via CYP 3A4 isoenzyme. Coadministration of the amlodipine component of CADUET with other drugs which follow the same route of biotransformation may result in altered bioavailability of amlodipine or these drugs. Dosages of similarly metabolized drugs, particularly those of low therapeutic ratio, and especially in patients with renal and/or hepatic impairment, may require adjustment when starting or stopping concomitantly administered amlodipine to maintain optimum therapeutic blood levels.

Co-administration of a 180 mg daily dose of diltiazem with 5 mg amlodipine in elderly hypertensive patients (69 to 87 years of age) resulted in a 57% increase in amlodipine systemic exposure. Erythromycin co-administration in healthy volunteers (18 to 43 years of age) increased the systemic exposure of amlodipine by 22%. These pharmacokinetic changes may be more pronounced in the elderly. Close monitoring and dose adjustments may be required. Strong inhibitors of CYP3A4 (e.g., ketoconazole, itraconazole, clarithromycin, ritonavir) may increase the plasma concentrations of amlodipine to a greater extent than diltiazem. Due to the amlodipine component of CADUET, CADUET should be used with caution together with CYP3A4 inhibitors. Monitoring of therapy is required.

There is no data available regarding the effect of CYP3A4 inducers on amlodipine. The concomitant use of CYP3A4 inducers may give a lower plasma concentration of amlodipine which in turn can result in decreased blood pressure lowering effects. Due to the amlodipine component of CADUET, CADUET should be used with caution together with CYP3A4 inducers and dose adjustment may be necessary to maintain efficacy. Hence, monitoring of therapy is required.

The amlodipine component of CADUET has a low (rate of first-pass) hepatic clearance and consequent high bioavailability, and thus, may be expected to have a low potential for clinically relevant effects associated with elevation of amlodipine plasma levels when used concomitantly with drugs that compete for or inhibit the cytochrome P450 system.

In clinical trials, the amlodipine component of CADUET has been safely administered with thiazide diuretics, beta blockers, angiotensin converting enzyme inhibitors, long acting nitrates, sublingual nitroglycerin, digoxin, warfarin, non steroidal anti-inflammatory drugs, antibiotics, and oral hypoglycemic drugs.

Atorvastatin**:** The atorvastatin component of CADUET is metabolized by the cytochrome P450 isoenzyme, CYP 3A4. Erythromycin, a CYP 3A4 inhibitor, increased atorvastatin plasma levels by 40%. Coadministration of CYP 3A4 inhibitors, such as grapefruit juice, some macrolide antibiotics (i.e. erythromycin, clarithromycin), immunosuppressants (cyclosporine), azole antifungal agents (i.e. itraconazole, ketoconazole), protease inhibitors, or the antidepressant, nefazodone, have the potential to increase plasma concentrations of HMG CoA reductase inhibitors, including atorvastatin (see Drug-Drug Interactions, REFERENCES). Concomitant use of atorvastatin with cyclosporine, gemfibrozil, telaprevir or the combination of tipranavir with ritonavir should be avoided. Other CYP 3A4 inhibitors should be used with atorvastatin dose restriction or caution, and patients should be monitored closely for musculoskeletal effects (see WARNINGS AND PRECAUTIONS, Pharmacokinetic Interactions, Muscle Effects, Renal Insufficiency and Endocrine Function; Drug-Drug Interactions, Table 3 – Established or Predicted Drug-Drug Interactions; REFERENCES).

Pharmacokinetic drug interactions that result in increased systemic concentration of atorvastatin have been noted with HIV protease inhibitors (lopinavir plus ritonavir, saquinavir plus ritonavir, darunavir plus ritonavir, fosamprenavir, fosamprenavir plus ritonavir and nelfinavir), Hepatitis C protease inhibitor (boceprevir), clarithromycin and itraconazole. Caution should be used when co-prescribing atorvastatin and appropriate clinical assessment is recommended to ensure that the lowest dose necessary of atorvastatin is employed (see Drug-Drug Interactions, Table 3 - Established or Potential Drug-Drug Interactions).

Inducers of cytochrome P450 3A4

Concomitant administration of atorvastatin with inducers of cytochrome P450 3A4 (e.g. efavirenz, rifampin) can lead to variable reductions in plasma concentrations of atorvastatin. Due to the dual interaction mechanism of rifampin, (cytochrome P450 3A4 induction and inhibition of hepatocyte uptake transporter OATP1B1), simultaneous co-administration of atorvastatin with rifampin resulted in a mean increase in Cmax and AUC of atorvastatin of 12 and 190%, respectively. In contrast, a delayed administration of atorvastatin after

administration of rifampin has been associated with a significant reduction (approximately 80%) in atorvastatin plasma concentrations.

Transporter Inhibitors

Atorvastatin and atorvastatin metabolites are substrates of the OATP1B1 transporter. Inhibitors of the OATP1B1 (e.g. cyclosporine) can increase the bioavailability of atorvastatin (see DETAILED PHARMACOLOGY, Human Pharmacokinetics).

Concomitant Therapy with Other Lipid Metabolism Regulators

Based on post-marketing surveillance, gemfibrozil, fenofibrate, other fibrates, and lipid-modifying doses of niacin (nicotinic acid) may increase the risk of myopathy when given concomitantly with HMG-CoA reductase inhibitors, probably because they can produce myopathy when given alone (see WARNINGS – Muscle Effects). The concomitant use of CADUET with gemfibrozil should be avoided. The combined therapy with other fibrates and niacin should be used with caution; lower starting and maintenance doses of the atorvastatin component of CADUET should be considered**.**

Drug-Drug Interactions

The drugs listed in this table are based on either drug interaction case reports or studies, or predicted interactions due to the expected magnitude and seriousness of the interaction (i.e. those identified as contraindicated).

Table 3 - Established or Predicted Drug-Drug Interactions*

Effect			**Clinical comment**
	Amlodipine	**Atorvastatin**	
Amlodipine		↔ In healthy subjects, atorvastatin PK were not altered by the coadministration of LIPITOR 80 mg and amlodipine 10 mg at steady state. No apparent changes in BP or HR. In healthy volunteers, co-administration of multiple 10 mg doses of amlodipine with 80 mg of atorvastatin resulted in no clinical significant change in the AUC (average of 18% increase) or C_{max} or T_{max} of atorvastatin.	See PHARMACOLOGY, Human Pharmacokinetics. Close monitoring is required.
Antacids (aluminum- and magnesium-based)	↔ on the disposition of amlodipine	↓ in plasma concentrations of atorvastatin by ~ 35% ↔in LDL-C reduction - triglyceride-lowering effect may be affected	

	Effect		Clinical comment
	Amlodipine	**Atorvastatin**	
Antipyrine		↔ in the PK of antipyrine	Antipyrine was used as a non-specific model for drugs metabolized by the microsomal hepatic enzyme system (cytochrome P450 system). Interactions with other drugs metabolized via the same cytochrome isozymes are not expected.
Beta-blockers	blood pressure lowering effect of beta-blockers may be ↑by amlodipine		Patients should be carefully monitored
Bile Acid Sequestrants		↓ in plasma concentration of atorvastatin by ~ 26%	See ACTIONS AND CLINICAL PHARMACOLOGY When atorvastatin is used concurrently with colestipol or any other resin, an interval of at least 2 hours should be maintained between the two drugs, since the absorption of atorvastatin may be impaired by the resin.
Cimetidine	↔ in the PK of amlodipine	↔ in plasma concentration of atorvastatin ↔ in LDL-C reduction ↓ triglyceride lowering effect from 34% to 26%	

	Effect		Clinical comment
	Amlodipine	**Atorvastatin**	
Cyclosporine	No drug interaction studies have been conducted with cyclosporine and amlodipine in healthy volunteers or other populations, with the exception of renal transplant patients. A prospective study in hypertensive renal transplant patients (N=11) showed on an average increase of 40% in trough cyclosporine levels when concomitantly treated with amlodipine	↑ Concomitant administration of atorvastatin 10 mg and cyclosporine 5.2 mg/kg/day resulted in a 7.7 fold increase in exposure to atorvastatin.	Concomitant use should be avoided. See WARNINGS and PRECAUTIONS, Muscle Effects; DOSAGE AND ADMINISTRATION, Concomitant Therapy DETAILED PHARMACOLOGY, Human Pharmacokinetics
Itraconazole		↑ Concomitant administration of atorvastatin 20 to 40 mg and itraconazole 200 mg daily resulted in 2.5 – 3.3-fold increase in atorvastatin AUC.	The dose of the atorvastatin component of CADUET used in combination with itraconazole should not exceed 20 mg daily (see DETAILED PHARMACOLOGY, Human Pharmacokinetics).

	Effect		Clinical comment
	Amlodipine	**Atorvastatin**	
Strong inhibitors of CYP3A4 (e,g, ketaconazole, itraconazole, ritonavir, clarithromycin)	May significantly increase the plasma concentrations of amlodipine to a greater extent than diltiazem.		Amlodipine should be used with caution together with CYP3A4 inhibitors and monitoring of therapy is required. Appropriate dosage adjustement of amlodipine may be necessary when used with CYP3A4 inhibitors. Patients should be advised to seek medical attention if they experience edema or swelling of the lower extremities; sudden, unexplained weight gain; difficulty breathing; chest pain or tightness; or hypotension as indicated by dizziness, fainting, or orthostasis. Avoid concomitant administration of amlodipine with strong CYP3A4 inhibitors.
Clarithromycin	In elderly patients (>65 years of age), concomitant use of amlodipine with clarithromycin was associated with increased risk of hospitalization with acute kidney injury.		Avoid concomitant use.
Diltiazem Hydrochloride	In elderly patients, the plasma concentration of amlodipine increased by 50 %	Steady-state diltiazem increases the exposure, based on AUC_{LASTs}, of a single dose of atorvastatin by approximately 50%.	
Digoxin	↔ in serum digoxin levels or digoxin renal clearance	↔ in digoxin PK by coadministration with atorvastatin 10 mg daily ↑ in digoxin concentrations ~ 20% following coadministration with atorvastatin 80 mg daily	See PHARMACOLOGY-Human Pharmacokinetics Patients taking digoxin should be monitored appropriately.

Effect			Clinical comment
	Amlodipine	**Atorvastatin**	
Fibric Acid Derivatives (gemfibrozil, fenofibrate, bezafibrate) and Niacin (nicotinic acid):		↑ in the risk of myopathy during treatment with other drugs in this class, including atorvastatin	The concomitant therapy with CADUET and gemfibrozil should be avoided. The benefits and risks of combined therapy with atorvastatin and fenofibrate, bezafibrate and niacin should be carefully considered. See WARNINGS, Muscle Effects and REFERENCES
Macrolide antibiotics	In young patients the plasma concentration of amlodipine increased by 22% with concomitant use of erythromycin	↑ in atorvastatin plasma levels by ~ 40% with erythromycin and ~ 80% with clarithromycin ↔ in atorvastatin plasma levels with azithromycin	See WARNINGS, Muscle Effects
Oral Contraceptives and Hormone Replacement Therapy		↑ in AUC of norethindrone by ~ 30% and ethinyl estradiol by ~ 20%	These increases should be considered when selecting an oral contraceptive. - In clinical studies, atorvastatin was used concomitantly with estrogen replacement therapy without evidence to date of clinically significant adverse interactions.

Effect			Clinical comment
	Amlodipine	**Atorvastatin**	
Protease Inhibitor		↑ in AUC by 74% and C_{max} by 122% of atorvastatin by nelfinavir mesylate 1250 mg BID, 14 days	The dose of the atorvastatin component of CADUET used in combination with nelfinavir should not exceed 40 mg daily.
		↑ in AUC by 5.9 fold and C_{max} by 4.7 fold with atorvastatin 20mg daily and Lopinavir 400mg / Ritonavir 100mg twice daily***	The concomitant therapy with CADUET and the combination of lopinavir plus ritonavir should be used with caution and lowest atorvastatin dose necessary. (See Warnings and Precautions, Muscle Effects)
		↑ in AUC by 2.9 fold and C_{max} by 3.3 fold with atorvastatin 40mg daily, for 4 days, and Ritonavir 400mg, BID,15 days / Saquinavir 400mg twice daily***†	† The dose of saquinavir plus ritonavir in this study is not the clinically used dose. The increase in atorvastatin exposure when used clinically is likely to be higher than what was observed in this study. Therefore caution should be applied and the lowest dose necessary should be used.
		↑ AUC by 8.4 fold and ↑ C_{max} by 7.6 fold with atorvastatin 10mg SD and Tipranavir 500mg BID / Ritonavir 200mg BID, 7 days. Atorvastatin 10 mg SD had no effect on the PK of Tripanavir 500mg BID / Ritonavir 200 mg BID, 7 days*	The concomitant therapy with CADUET and the combination of tipranavir plus ritonavir or CADUET and telaprevir should be avoided.
		↑ AUC by 6.9 fold and ↑ C_{max} by 9.6 fold with atorvastatin 20mg SD and Telaprevir 750mg q8h, 10 days*	

	Effect		Clinical comment
	Amlodipine	**Atorvastatin**	
		↑ AUC by 2.30 fold and ↑ C_{max} by 2.66 fold with atorvastatin 40mg SD and Boceprevir 800 mg TID, 7 days ↑ AUC by 2.4 fold and ↑ C_{max} by 1.3 fold with atorvastatin 10mg QD for 4 days and Darunavir 300mg BID/ Ritonavir 100 mg BID, 9 days* ↑ AUC by 1.5 fold and ↑ C_{max} by 1.8 fold with atorvastatin 10mg QD for 4 days and Fosamprenavir 700 mg BID/ritonavir 100mg BID,14 days* ↑ AUC by 1.3 fold and ↑ C_{max} by 3.0 fold with atorvastatin 10mg QD for 4 days and Fosamprenavir 1400 mg BID, 14 days*. Atorvastatin 10mg QD for 4 days had the following effect on the PK of Fosamprenavir 1400 mg BID, 14 days: ↓ AUC by 0.27 fold and ↓ C_{max} by 0.18 fold* Atorvastatin 10mg QD, 4 days had no effect on the PK of Fosamprenavir 700mg BID/ Ritonavir 100 mg BID, 14 days*	The dose of the atorvastatin component of CADUET should be restricted to 20 mg daily when used in combination with boceprevir, saquinavir plus ritonavir, darunavir plus ritonavir, fosamprenavir alone or fosamprenavir plus ritonavir
Quinapril		↔ in PK profile of atorvastatin	See PHARMACOLOGY, Human Pharmacokinetics
Sildenafil	↔ in AUC or Cmax of amlodipine - mean additional reduction of supine systolic and diastolic blood pressure was 8 mmHg and 7 mmHg, respectively		
Warfarin	↔ in warfarin-induced prothrombin response time	↔ in warfarin-induced prothrombin response time	

Effect			Clinical comment
	Amlodipine	**Atorvastatin**	
Efavirenz		↓ in AUC by 41% and C_{max} by 1% with atorvastatin 10mg and Efavirenz 600mg daily***	
Rifampin		Co-administration***: ↑ in AUC by 30% and C_{max} by 2.7 fold co-administered atorvastatin 40mg single dose and Rifampin 600mg daily Separate administration*** ↓ in AUC by 80% and C_{max} by 40 % with atorvastatin 40mg single dose and Rifampin 600mg daily (doses separated)	Due to the dual interaction mechanism of rifampin, simultaneous co administration of atorvastatin with rifampin is recommended, as delayed administration of atorvastatin after administration of rifampin has been associated with a significant reduction in atorvastatin plasma concentrations.
Fusidic Acid		Although interaction studies with the atorvastatin component of CADUET and fusidic acid have not been conducted, severe muscle problems such as rhabdomyolysis have been reported in post-marketing experience with this combination.	Temporary suspension of atorvastatin treatment should be considered (see WARNINGS AND PRECAUTIONS, Muscle Effects).
Colchicine		Although interaction studies with atorvastatin and colchicine have not been conducted, cases of myopathy have been reported with atorvastatin co-administrated with colchicine.	Caution should be exercised when prescribing atorvastatin with colchicine. (See Warnings and Precautions, Muscle Effect)
Tacrolimus	There is a risk of increased tacrolimus blood levels when co-administered with amlodipine.		In order to avoid toxicity of tacrolimus, administration of amlodipine in a patient treated with tacrolimus requires monitoring of tacrolimus blood levels and dose adjustments of tacrolimus when appropriate.

* For more detailed drug interaction information please refer to individual Product Monographs for NORVASC and LIPITOR.

** Legend: ↔ = no change; ↑ = increase; ↓ = decrease; ~ approximately; AUC = area under the curve; Cmax = maximal concentrations; LDL-C = low density lipoprotein cholesterol; PK = pharmacokinetics; Tmax = time to maximal concentrations

*** Data given as x-fold change represent a simple ratio between co-administration and atorvastatin alone (i.e., 1-fold = no change). Data given as % change represent % difference relative to atorvastatin alone (i.e., 0% = no change).

Drug-Food Interactions

Grapefruit Juice

Because of the potential effects of grapefruit juice on both the amlodipine and atorvastatin components of CADUET, administration is not recommended.

Amlodipine: Published data indicate that through inhibition of the cytochrome P450 system, grapefruit juice can increase plasma levels and augment pharmacodynamic effects of some dihydropyridine calcium channel blockers.
Co-administration of 240 mL of grapefruit juice with a single oral dose of amlodipine 10 mg in 20 healthy volunteers had no significant effect on the pharmacokinetics of amlodipine. The study did not allow examination of the effect of genetic polymorphism in CYP3A4, the primary enzyme responsible for metabolism of amlodipine, therefore administration of amlodipine with grapfruit or grapefruit juice is not recommended as bioavailability may be increased in some patients resulting in increased blood pressure lowering effects.

Atorvastatin: Coadministration of grapefruit juice has the potential to increase plasma concentrations of HMG CoA reductase inhibitors including LIPITOR. The equivalent of 1.2 litres per day resulted in a 2.5 fold increase in AUC of atorvastatin.

DOSAGE AND ADMINISTRATION

CADUET is a combination product containing amlodipine besylate and atorvastatin calcium.

CADUET is not intended for initial therapy.

The dosage of CADUET must be individualized on the basis of both effectiveness and tolerance for each component which should be determined by titration as described below.

CADUET can be administered once daily, at any time of the day, with or without food.

Amlodipine

Use in Adults

For both hypertension and angina, the recommended initial dose of amlodipine besylate is 5 mg once daily. If necessary, dose can be increased after 1-2 weeks to a maximum dose of 10 mg once daily.

Use in the Elderly or in Patients with Impaired Renal Function

The recommended initial dose in patients over 65 years of age or patients with impaired renal function is 5 mg once daily. If required, increasing in the dose should be done gradually and with caution (see WARNINGS AND PRECAUTIONS, Renal Insufficiency; Special

Populations, Geriatrics).

Use in Patients with Impaired Hepatic Function

Dosage requirements have not been established in patients with impaired hepatic function. When amlodipine is used in these patients, the dosage should be carefully and gradually adjusted depending on patients tolerance and response. A lower starting dose of 2.5 mg once daily should be considered (see WARNINGS AND PRECAUTIONS, Hepatic Effects).

Use in Children

There have been no studies conducted to determine the safety or efficacy of CADUET in pediatric patients.

The effective antihypertensive oral dose in pediatric patients ages 6-17 years is 2.5 mg to 5 mg once daily. Doses in excess of 5 mg daily have not been studied; dose should be determined based upon the medical need of the patients (See WARNINGS AND PRECAUTIONS Special Populations, Pediatrics).

Atorvastatin

Patients should be placed on a standard cholesterol-lowering diet before receiving LIPITOR, and should continue on this diet during treatment with LIPITOR. If appropriate, a program of weight control and physical exercise should be implemented.

Prior to initiating therapy with LIPITOR, secondary causes for elevations in plasma lipid levels should be excluded. A lipid profile should also be performed.

Primary Hypercholesterolemia and Combined (Mixed) Dyslipidemia, Including Familial Combined Hyperlipidemia

The recommended starting dose of LIPITOR is 10 or 20 mg once daily, depending on patient's LDL-C reduction required. Patients who require a large reduction in LDL-C (more than 45%) may be started at 40 mg once daily. The dosage range of LIPITOR is 10 to 80 mg once daily. Doses can be given at any time of the day with or without food, and should preferably be given in the evening. A significant therapeutic response is evident within 2 weeks, and the maximum response is usually achieved within 2-4 weeks. The response is maintained during chronic therapy. Adjustments of dosage, if necessary, should be made at intervals of 2 to 4 weeks. The maximum dose is 80 mg/day.

The dosage of LIPITOR should be individualized according the baseline LDL-C, total-C/HDL-C ratio and/or TG levels to achieve the recommended desired lipid values at the lowest dose needed to achieve LDL-C desired level. Lipid levels should be monitored periodically and, if necessary, the dose of LIPITOR adjusted based on desired lipid levels recommended by guidelines.

Severe Dyslipidemias

In patients with severe dyslipidemias, including homozygous and heterozygous familial hypercholesterolemia and dysbetalipoproteinemia (Type III), higher dosages (up to 80 mg/day) may be required (see WARNINGS AND PRECAUTIONS, Pharmacokinetic Interactions, Muscle Effects; DRUG INTERACTIONS).

Heterozygous Familial Hypercholesterolemia in Pediatric Patients (10-17 years of age)

There have been no studies conducted to determine the safety or efficacy of CADUET in pediatric patients.

In this population, the recommended starting dose of LIPITOR is 10 mg/day; the maximum recommended dose is 20 mg/day (doses greater than 20 mg/day have not been studied in this patient population). Doses should be individualized according to the recommended goal of therapy (see WARNINGS AND PRECAUTIONS, Special Populations, Pediatrics). Adjustments should be made at intervals of 4 weeks or more.

Prevention of Cardiovascular Disease

Clinical trials conducted that evaluated atorvastatin in the primary prevention of myocardial infarction used a dose of 10 mg atorvastatin once daily.

For secondary prevention of myocardial infarction, optimal dosing may range from 10 mg to 80 mg atorvastatin once daily, to be given at the discretion of the prescriber, taking into account the expected benefit and safety considerations relevant to the patient to be treated.

Concomitant Therapy

See DRUG INTERACTIONS

Patients with Renal Insufficiency

See WARNINGS AND PRECAUTIONS, Renal.

OVERDOSAGE

There is no information on overdosage with CADUET in humans.

For management of a suspected drug overdose, contact your regional Poison Control Center.

Amlodipine

Symptoms
Overdosage can cause excessive peripheral vasodilation with marked and probably prolonged hypotension and possibly a reflex tachycardia. In humans, experience with overdosage of the amlodipine component of CADUET is limited. When amlodipine was ingested at doses of 105-250 mg some patients remained normotensive with or without gastric lavage while another patient experienced hypotension (90/50 mmHg) which normalized following plasma expansion. A patient who took 70 mg of amlodipine with benzodiazepine developed shock which was refractory to treatment and died. In a 19-month old child who ingested 30 mg of amlodipine (about 2 mg/kg) there was no evidence of hypotension but tachycardia (180 bpm) was observed. Ipecac was administered 3.5 hrs after ingestion and on subsequent observation (overnight) no sequelae were noted.

Treatment
Clinically significant hypotension due to overdosage requires active cardiovascular support, including frequent monitoring of cardiac and respiratory function, elevation of extremities, and attention to circulating fluid volume and urine output. A vasoconstrictor (such as norepinephrine) may be helpful in restoring vascular tone and blood pressure, provided that there is no contraindication to its use. As amlodipine is highly protein bound, hemodialysis is not likely to be of benefit. Intravenous calcium gluconate may be beneficial in reversing the effects of calcium channel blockade. Clearance of amlodipine is prolonged in elderly patients and in patients with impaired liver function. Since amlodipine absorption is slow, gastric lavage may be worthwhile in some cases.

Atorvastatin

There is no specific treatment for the atorvastatin component of CADUET overdosage. Should an overdose occur, the patient should be treated symptomatically and supportive measures instituted as required. Due to extensive drug binding to plasma proteins, hemodialysis is not expected to significantly enhance atorvastatin clearance.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action:

CADUET (amlodipine besylate/atorvastatin calcium), is a combination tablet which combines 2 mechanisms of action: the dihydropyridine calcium antagonist (calcium entry blocker or calcium ion antagonist) action of amlodipine and the HMG-CoA reductase inhibition of atorvastatin. The amlodipine component of CADUET inhibits the transmembrane influx of calcium ions into vascular smooth muscle and cardiac muscle. The atorvastatin component of CADUET is a selective, competitive inhibitor of HMG-CoA reductase, the rate-limiting enzyme that converts 3-hydroxy-3-methylglutaryl-coenzyme A to mevalonate, a precursor of sterols, including cholesterol.

The antihypertensive/antianginal action of CADUET:

Experimental data suggest that amlodipine binds to both dihydropyridine and nondihydropyridine binding sites. The contractile processes of cardiac and vascular smooth muscle tissues are dependent upon the movement of extracellular calcium ions into these cells through specific ion channels. Amlodipine inhibits calcium ion influx across cell membranes selectively, with a greater effect on vascular smooth muscle cells than on cardiac muscle cells. Serum calcium concentration is not affected by amlodipine. Within the physiologic pH range, amlodipine is an ionized compound and its kinetic interaction with the calcium channel receptor is characterized by the gradual association and dissociation with the receptor binding site.

- Hypertension: The mechanism by which amlodipine reduces arterial blood pressure involves direct peripheral arterial vasodilation and reduction in peripheral vascular resistance.

- Angina: The precise mechanism by which amlodipine relieves angina has not been fully delineated. Amlodipine is a dilator of peripheral arteries and arterioles which reduces the total peripheral resistance and, therefore, reduces the workload of the heart (afterload). The unloading of the heart is thought to decrease ischemia and relieve effort angina by reducing myocardial energy oxygen consumption and oxygen requirements.

The antidyslipidemic action of CADUET:

Atorvastatin lowers plasma cholesterol and lipoprotein levels by inhibiting HMG-CoA reductase and cholesterol synthesis in the liver and by increasing the number of hepatic LDL receptors on the cell-surface for enhanced uptake and catabolism of LDL.

Atorvastatin reduces LDL-C and the number of LDL particles. Atorvastatin also reduces VLDL-C, serum TG and IDL, as well as the number of apo B containing particles, but increases HDL-C. Elevated serum cholesterol due to elevated LDL-C is a major risk factor for the development of cardiovascular disease. Low serum concentration of HDL-C is an independent risk factor. Elevated plasma TG is also a risk factor for cardiovascular disease, particularly if due to increased IDL, or associated with decreased HDL-C or increased LDL-C.

Epidemiologic, clinical and experimental studies have established that high LDL-C, low HDL-C and high plasma TG promote human atherosclerosis, and are risk factors for developing cardiovascular disease. Some studies have also shown that the total (TC):HDL-C ratio (TC:HDL-C) is the best predictor of coronary artery disease. In contrast, increased levels of HDL-C are associated with decreased cardiovascular risk. Drug therapies that reduce levels of LDL-C or decrease TG while simultaneously increasing HDL-C have demonstrated reductions in rates of cardiovascular mortality and morbidity.

Pharmacodynamics:

CADUET

Studies have been conducted in which placebo, amlodipine alone, atorvastatin alone, and the 8 dose combinations of amlodipine and atorvastatin have been administered once daily, in patients with co- morbid dyslipidemia and hypertension. Analyses of changes in systolic blood pressure demonstrated that there was no overall modification of amlodipine's effect on systolic blood pressure when the drug was taken in combination with atorvastatin compared to amlodipine alone. Analyses of changes in LDL-C demonstrated that there was no overall modification of atorvastatin's effect on LDL-C when the drug was taken in combination with amlodipine compared with atorvastatin alone (see CLINICAL TRIALS).

Amlodipine

Hemodynamics

Following administration of recommended doses to patients with hypertension, amlodipine produces vasodilation resulting in a reduction of supine and standing blood pressures. These decreases in blood pressure are not accompanied by any significant change in heart rate or plasma catecholamine levels with chronic dosing. With chronic once daily oral administration (5 and 10 mg once daily), antihypertensive effectiveness is maintained throughout the 24-hour dose interval with minimal peak to trough differences in plasma concentration. Since the vasodilation induced by amlodipine is gradual in onset, acute hypotension has rarely been reported after oral administration of amlodipine. In normotensive patients with angina, amlodipine has not been associated with any clinically significant reductions in blood pressure or changes in heart rate.

Negative inotropic effects have not been observed when amlodipine was administered at the recommended doses to man, but has been demonstrated in animal models. Hemodynamic measurements of cardiac function at rest and during exercise (or pacing) in angina patients with normal ventricular function have generally demonstrated a small increase in cardiac index without significant influence on dP/dt or on left ventricular end diastolic pressure or volume.

In hypertensive patients with normal renal function, therapeutic doses of amlodipine resulted in a decrease in renal vascular resistance and an increase in glomerular filtration rate and effective renal plasma flow without change in filtration fraction.

Electrophysiologic Effects:

Amlodipine does not change sinoatrial nodal function or atrioventricular conduction in intact animals, or man. In patients with chronic stable angina, intravenous administration of 10 mg of amlodipine and a further 10 mg of amlodipine after a 30-minute interval produced peripheral vasodilation and afterload reduction, but did not significantly alter A-H and H-V conduction and sinus node recovery time after pacing. Similar results were obtained in patients receiving amlodipine and concomitant beta-blockers. In clinical studies in which amlodipine was administered in combination with beta-blockers to patients with either hypertension or angina, no

adverse effects on electrocardiographic parameters were observed. In clinical trials with angina patients, amlodipine as monotherapy did not alter electrocardiographic intervals.

Atorvastatin

Human Pharmacology

The lowering of total cholesterol, LDL-C and apo B have been shown to reduce the risk of cardiovascular events and mortality.

Atorvastatin is a selective, competitive inhibitor of HMG-CoA reductase. In both subjects and in patients with homozygous and heterozygous familial hypercholesterolemia, nonfamilial forms of hypercholesterolemia, mixed dyslipidemia, hypertriglyceridemia, and dysbetalipoproteinemia, atorvastatin has been shown to reduce levels of total-C, LDL-C, apo B and total TG, and raises HDL-C levels.

Epidemiologic and clinical studies have associated the risk of coronary artery disease (CAD) with elevated levels of total-C, LDL-C and decreased levels of HDL-C. These abnormalities of lipoprotein metabolism are considered as major contributors to the development of the disease. Like LDL, cholesterol-enriched lipoproteins, including VLDL, IDL and remnants can also promote atherosclerosis. Elevated plasma triglycerides are frequently found in a triad with low HDL-C levels and small LDL particles, as well as in association with non-lipid metabolic risk factors for coronary heart disease (metabolic syndrome). Clinical studies have also shown that serum triglycerides can be an independent risk factor for CAD. CAD risk is especially increased if the hypertriglyceridemia is due to increased intermediate density lipoproteins (IDL) or associated with decreased HDL or increased LDL-C. In addition, high TG levels are associated with an increased risk of pancreatitis. Although epidemiological and preliminary clinical evidence link low HDL-C levels and high triglyceride levels with coronary artery disease and atherosclerosis, the independent effect of raising HDL or lowering TG on the risk of coronary and cerebrovascular morbidity and mortality has not been demonstrated in prospective, well-controlled outcome studies. Other factors, e.g. interactions between lipids/lipoproteins and endothelium, platelets and macrophages, have also been incriminated in the development of human atherosclerosis and of its complications. Regardless of the intervention used (low-fat/low-cholesterol diet, partial ileal bypass surgery or pharmacologic therapy), effective treatment of hypercholesterolemia/dyslipidemia has consistently been shown to reduce the risk of CAD.

Atorvastatin reduces LDL-C and the number of LDL particles, lowersVLDL-C and serum TG, reduces the number of apo B containing particles, and also increases HDL-C. Atorvastatin is effective in reducing LDL-C in patients with homozygous familial hypercholesterolemia, a condition that rarely responds to any other lipid-lowering medication. In addition to the above effects, atorvastatin reduces IDL-C and apolipoprotein E (apo E) in patients with dysbetalipoproteinemia (Type III).

In patients with Type II dyslipidemia, atorvastatin improved endothelial dysfunction. Atorvastatin significantly improved flow-mediated endothelium-dependent dilatation induced by reactive hyperemia, as assessed by brachial ultrasound ($p<0.01$).

Pharmacokinetics

Absorption

CADUET

Following oral administration of therapeutic doses of CADUET tablets, 2 distinct peak plasma concentrations are observed. The first peak is attributable to atorvastatin and occurs within 1 to 2 hours after dosing. The second peak is attributable to amlodipine and occurs between 6 and 12 hours after dosing. The rate and extent of absorption (bioavailability) of both amlodipine and atorvastatin from CADUET combination tablet are not significantly different from those observed during coadministration of separate amlodipine and atorvastatin tablets, as assessed by Cmax: 101% (90% CI: 98, 104) and AUC: 100% (90% CI: 97, 103) for the amlodipine component and Cmax: 94% (90% CI: 85, 104) and AUC: 105% (90% CI: 99, 111) for the atorvastatin component, respectively.

The bioavailability of amlodipine from the CADUET tablet was not affected under the fed state as assessed by Cmax and AUC. Food decreases the rate and extent of absorption of atorvastatin from the CADUET tablets by approximately 32% and 11%, respectively. Similar reductions in plasma concentrations were observed with atorvastatin in the fed state without a reduction in LDL-C effect.

Amlodipine:
After oral administration of therapeutic doses of amlodipine, absorption occurs gradually with peak plasma concentration reached between 6 and 12 hours. Absolute bioavailability has been estimated to be between 64 and 90%. The bioavailability of amlodipine is not altered by the presence of food.

Atorvastatin:
Atorvastatin is rapidly absorbed after oral administration; maximal plasma concentrations occur within 1 to 2 hours. Extent of absorption and plasma atorvastatin concentrations increase in proportion to atorvastatin dose. The absolute bioavailability (parent drug) of atorvastatin is approximately 12% and the systemic availability of HMG-CoA reductase inhibitory activity is approximately 30%. The low systemic availability is attributed to presystemic clearance in gastrointestinal mucosa and/or first-pass metabolism in the liver. Although food decreases the rate and extent of drug absorption by approximately 25% and 9%, as assessed by Cmax and AUC respectively, LDL-C reduction and HDL-C elevation are similar when atorvastatin is given with and without food. Plasma atorvastatin concentrations are lower (approximately 30% for Cmax and AUC) following drug administration in the evening compared with morning dosing. However, LDL-C reduction and HDL-C elevation are the same regardless of the time of drug administration.

Distribution

Amlodipine:
Ex vivo studies have shown that approximately 93% of the circulating drug is bound to plasma proteins in hypertensive patients. Steady state plasma levels of amlodipine are reached after 7 to 8 days of consecutive daily dosing.

Atorvastatin:
Mean volume of distribution of atorvastatin is approximately 381 liters. Atorvastatin is ≥98% bound to plasma proteins. A blood/plasma ratio of approximately 0.25 indicates poor drug penetration into red blood cells. Based on observations in rats, atorvastatin is likely to be secreted in human milk (see CONTRAINDICATIONS, WARNINGS AND PRECAUTIONS).

Metabolism

Amlodipine:
Amlodipine is metabolized through the cytochrome P450 system, mainly via CYP 3A4 isoenzyme. Amlodipine is extensively (about 90%) converted to inactive metabolites (via hepatic metabolism).

Atorvastatin:
Atorvastatin is extensively metabolized to ortho- and para-hydroxylated derivatives by cytochrome P450 system via the CYP 3A4 isoenzyme and to various beta-oxidation products. In vitro, inhibition of HMG-CoA reductase by ortho- and para-hydroxylated metabolites is equivalent to that of atorvastatin. Approximately 70% of circulating inhibitory activity for HMG-CoA reductase is attributed to active metabolites. In animals, the ortho-hydroxy metabolite undergoes further glucuronidation. Atorvastatin and its metabolites are eliminated by biliary excretion.

Excretion

Amlodipine:
Elimination from the plasma is biphasic with a terminal elimination half-life of about 35-50 hours. Ten percent (10%) of the parent compound and 60% of the metabolites are excreted in the urine.

Atorvastatin:
Atorvastatin is eliminated primarily in bile following hepatic and/or extrahepatic metabolism; however, the drug does not appear to undergo significant enterohepatic recirculation. Mean plasma elimination half-life of atorvastatin in humans is approximately 14 hours, but the half-life for inhibitory activity for HMG-CoA reductase is 20 to 30 hours due to the contribution of longer-lived active metabolites. Less than 2% of a dose of atorvastatin is recovered in urine following oral administration.

Special Populations and Conditions

Pediatric

Pharmacokinetic data in the pediatric population are not available.

Geriatrics

Amlodipine:
In elderly hypertensive patients (mean age 69 years) there was a decrease in clearance of amlodipine from plasma as compared to young volunteers (mean age 36 years) with a resulting increase in the area under the curve (AUC) of about 60%.

Atorvastatin:
Plasma concentrations of atorvastatin are higher (approximately 40% for Cmax and 30% for AUC) in healthy elderly subjects (age 65 years or older) compared with younger individuals. LDL-C reduction, however, is comparable to that seen in younger patient populations.

Gender

Atorvastatin:
Plasma concentrations of atorvastatin in women differ (approximately 20% higher for Cmax and 10% lower for AUC) from those in men; however, there is no clinically significant difference in LDL-C reduction between men and women**.**

Race

Atorvastatin:
Plasma concentrations of atorvastatin are similar in black and white subjects.

Hepatic Insufficiency

Amlodipine:
Following single oral administration of 5 mg of amlodipine, patients with chronic mild-moderate hepatic insufficiency showed about 40% increase in AUC of amlodipine as compared to normal volunteers. This was presumably due to a reduction in clearance of amlodipine as the terminal elimination half-life was prolonged from 34 hrs in young normal subjects to 56 hrs in the elderly patients with hepatic insufficiency.

Atorvastatin:
Plasma concentrations of atorvastatin are markedly increased (approximately 16-fold in Cmax and 11-fold in AUC) in patients with chronic alcoholic liver disease (Childs-Pugh B).

Renal Insufficiency

Amlodipine:
The pharmacokinetics of amlodipine are not significantly influenced by renal impairment. Plasma concentrations in the patients with moderate to severe renal failure were higher than in the normal subjects. Accumulation and mean elimination half-life in all patients were within the range of those observed in other pharmacokinetic studies with amlodipine in normal subjects.

Atorvastatin:
Plasma concentrations and LDL-C lowering efficacy of atorvastatin are similar in patients with moderate renal insufficiency compared with patients with normal renal function. However, since several cases of rhabdomyolysis have been reported in patients with a history of renal insufficiency of unknown severity, as a precautionary measure and pending further experience in renal disease, the lowest dose (10 mg/day) of atorvastatin should be used in these patients. Similar precautions apply in patients with severe renal insufficiency (creatinine clearance <30 mL/min [<0.5 mL/sec]); the lowest dosage should be used and implemented cautiously (see WARNINGS AND PRECAUTIONS, Muscle Effects, DRUG INTERACTIONS; DOSAGE AND ADMINISTRATION).

STORAGE AND STABILITY

Store at 25°C (77°F); excursions permitted to 15-30°C (59-86°F).

SPECIAL HANDLING INSTRUCTIONS

Not Applicable.

DOSAGE FORMS, COMPOSITION AND PACKAGING

CADUET (amlodipine besylate/atorvastatin calcium) tablets are formulated for oral administration to contain amlodipine besylate and atorvastatin calcium and are available in tablet doses of 5mg/10mg, 5mg/20mg, 5mg/40mg, 5mg/80mg, 10mg/10mg, 10mg/20mg, 10mg/40mg, and 10mg/80mg, respectively.

CADUET tablets are differentiated by tablet color/size and are engraved with “Pfizer” on one side and a unique number on the other side. CADUET tablets are supplied for oral administration in 5mg/10mg (white), 5mg/20mg (white), 5mg/40mg (white), 5mg/80mg (white), 10mg/10mg (blue), 10mg/20mg (blue), 10mg/40mg (blue) and 10mg/80mg (blue) tablets.

CADUET tablets are available in high-density polyethylene (HDPE) bottles, containing desiccant, in packs of 90 tablets, with child-resistant closure.

CADUET Tablets		
Package Configuration	**Tablet Strength (amlodipine besylate/atorvastatin calcium) mg**	**Engraving**
Bottle of 90	**5mg/40mg**	**CDT 054**
Bottle of 90	**5mg/80mg**	**CDT 058**
Bottle of 90	**10mg/40mg**	**CDT 104**
Bottle of 90	**10mg/80mg**	**CDT 108**
Bottle of 90	**5mg/10mg**	**CDT 051**
Bottle of 90	**5mg/20mg**	**CDT 052**
Bottle of 90	**10mg/10mg**	**CDT 101**
Bottle of 90	**10mg/20mg**	**CDT 102**

Each tablet contains the following non-medicinal ingredients: Calcium Carbonate, Croscarmellose Sodium, Microcrystalline Cellulose, Pregelatinized Starch, Polysorbate 80, Hydroxypropyl Cellulose, Purified Water, Colloidal Silicon Dioxide (anhydrous), Magnesium Stearate, Opadry® II White 85F28751 or Opadry® II Blue 85F10919.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

CADUET

Proper Name:

amlodipine besylate/atorvastatin calcium

Physical Form:

CADUET **i**s a white to off-white crystalline powder, containing amlodipine besylate with a molecular weight of 567.11 and atorvastatin calcium with a molecular weight of 1209.422.

Drug Substance

Amlodipine component of CADUET

Proper Name:

amlodipine besylate

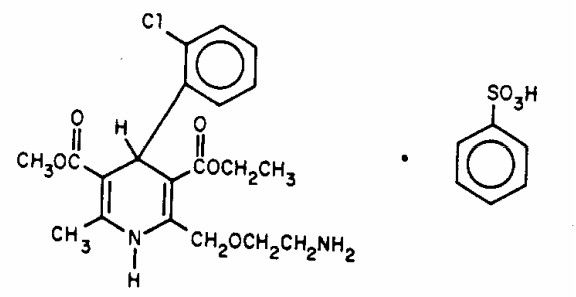
Chemical Name:

3-Ethyl-5-methyl-2-(2-aminoethoxymethyl)-4-(2-chlorophenyl)-1,4-dihydro-6-methyl-3,5-pyridinedicarboxylate benzenesulphonate.

Molecular Formula:

$C_{20}H_{25}ClN_2O_5.C_6H_6O_3S$

Structural Formula:



Molecular Weight:

567.1

Physical Form:

Amlodipine besylate is a white crystalline substance.

Solubility:

Amlodipine besylate is slightly soluble in water and sparingly soluble in ethanol, M.P.= 203ºC with decomposition. pKa = 9.02 at 23.5ºC.

Atorvastatin component of CADUET

Proper Name:

atorvastatin calcium

Chemical Name:

[R-(R*,R*)]-2-(4-fluorophenyl)-ß,∗-dihydroxy-5-(1-methylethyl)-3-phenyl-4-[(phenylamino)-carbonyl]-1H-pyrrole-1-heptanoic acid, calcium salt (2:1) trihydrate

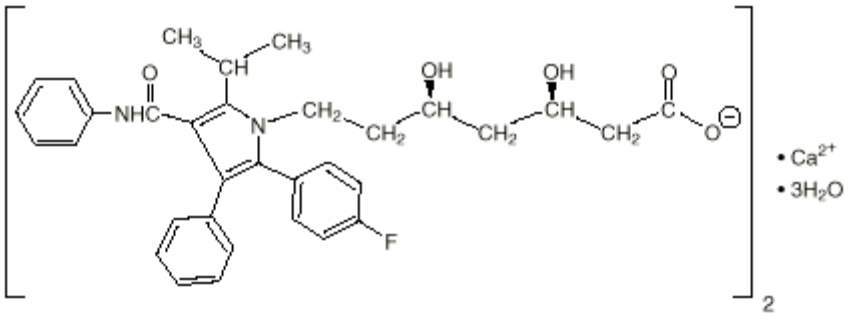
Empirical Formula:

$(C_{33}H_{34}FN_2O_5)_2CaX3H_2O$

Molecular Weight:

1209.42

Structural Formula:



Physical Form:

Atorvastatin calcium is a white to off-white crystalline powder.

Solubility:

Atorvastatin calcium is practically insoluble in aqueous solutions of pH 4 and below. Atorvastatin calcium is very slightly soluble in distilled water, pH 7.4 phosphate buffer, and acetonitrile, slightly soluble in ethanol, and freely soluble in methanol.

CLINICAL TRIALS

CADUET

Clinical studies in patients with hypertension and dyslipidemia

In a double-blind, placebo-controlled study, a total of 1660 patients with co-morbid hypertension and dyslipidemia received once daily treatment with 8 dose combinations of amlodipine besylate and atorvastatin calcium (5/10 mg, 5/20 mg, 5/40 mg, 5/80 mg, 10/10 mg, 10/20 mg, 10/40 mg, 10/80 mg), amlodipine alone (5 mg and 10 mg), atorvastatin alone (10 mg, 20 mg, 40 mg, 80 mg) or placebo. At 8 weeks, all 8-combination treatment groups of amlodipine and atorvastatin demonstrated statistically significant dose-related reductions in systolic blood pressure and LDL-C compared to placebo, with no overall modification of effect of either component on SBP and LDL-C (Table 4).

Table 4 - Primary Efficacy Analysis: Efficacy of the Combined Treatments in Reducing SBP and LDL-C

Efficacy of the Combined Treatments in Reducing Systolic BP						
Parameter / Analysis		**Placebo**	**ATO 10 mg**	**ATO 20 mg**	**ATO 40 mg**	**ATO 80 mg**
Placebo	LS mean change mmHg	-2.9	-4.3	-6.1	-6.2	-6.6
AML 5 mg	LS mean change mmHg	-12.6	-13.6	-15.3	-12.8	-12.6
	95% CIs		-12.3/ -6.3	-12.2/ -6.2	-9.7/ -3.6	-9.0/ -3.0
AML 10 mg	LS mean change mmHg	-16.5	-15.9	-16	-16.5	-17.5
	95% CIs		-14.6/ -8.5	-12.9/ -6.8	-13.3/ -7.2	-14.0/ -7.9

Efficacy of the Combined Treatments in Reducing LDL-C						
Parameter / Analysis		**Placebo**	**ATO 10 mg**	**ATO 20 mg**	**ATO 40 mg**	**ATO 80 mg**
Placebo	LS mean % chg	-1.2	-33.5	-39.5	-43.1	-47
AML 5 mg	LS mean % chg	-0.1	-39	-42.2	-44.9	-48.2
	95% CIs		-42.9/ -34.9	-46.2/ -38.2	-48.8/ -40.8	-52.2/ -44.2
AML 10 mg	LS mean % chg	-2.6	-36.6	-38.6	-43.2	-49.2
	95% CIs		-38.1/ -30.0	-40.0/ -32.0	-44.6/ -36.7	-50.6/ -42.6

ATO: Atorvastatin AML: Amlodipine LDL-C: Low density lipoprotein cholesterol SBP: Systolic Blood Pressure
Comparisons described above were between each individual combination treatment group and the corresponding amlodipine treatment group. BASELINE LDL-C= 182.0mg/dL SBP=148.4mmHg

In a double-blind, placebo-controlled study, a total of 847 patients with co-morbid hypertension and dyslipidemia received once daily placebo, 5 mg amlodipine, 10 mg of atorvastatin or the combination of 5 mg amlodipine and 10 mg atorvastatin. The primary objective of the study was the percentage of patients on the combination of amlodipine and atorvastatin reaching JNC VI and NCEP III goals compared to atorvastatin, amlodipine and placebo alone. The results following 8 weeks of treatment are summarized in Table 5. Significantly more patients treated with the combination (45.5%) reached both their BP and LDL-C goals compared to amlodipine or atorvastatin alone.

Table 5 - Results of efficacy end-points in placebo-controlled study of amlodipine/atorvastatin in patients with hypertension and dyslipidemia

	Placebo N = 239	ATO 10 mg N = 200	AML 5 mg N = 201	ATO 10 mg & AML 5 mg N= 207
JNC VI* Blood Pressure goals	29.7%	32.3%	54%	51%†
NCEP ATP III LDL-C goals	6.6%	78.2%	12.4%	82.1%**
Both JNC VI and NCEP ATP III* goals	3.5%	28.6%	8.3%	45.5%*†
Change in BP mmHg	-5.4/-3.3	-5.9/-4.2	-14.3/ -8.9	-12.7/ -8.2 +
Change in LDL-C -%	0.2	-33.9	-1.8	-37.2 a

ATO: Atorvastatin AML: Amlodipine LDL-C: Low density lipoprotein cholesterol SBP: Systolic Blood Pressure
** *P*<0.001 versus amlodipine
† *P*<0.001 versus atorvastatin
+ p< 0.001 vs. atorvastatin and NS vs. amlodipine
a p=0.07 vs Atorvastatin & <0.001 vs amlodipine
BASELINE LDL-C = 163.5 mg/dL, SBP = 146.9mmHg
***** BP goals in JNC VII for this population are consistent with JNC VI BP goals

Amlodipine

Effects in Hypertension: The antihypertensive efficacy of amlodipine has been demonstrated in a total of 15 double-blind, placebo-controlled, randomized studies involving 800 patients on amlodipine and 538 on placebo. Once daily administration produced statistically significant placebo-corrected reductions in supine and standing blood pressures at 24 hours postdose, averaging about 12/6 mmHg in the standing position and 13/7 mmHg in the supine position in patients with mild to moderate hypertension. Maintenance of the blood pressure effect over the 24-hour dosing interval was observed, with little difference in peak and trough effect. Tolerance was not demonstrated in patients studied for up to 1 year. The 3 parallel, fixed-dose, dose response studies showed that the reduction in supine and standing blood pressures was dose related within the recommended dosing range. Effects on diastolic pressure were similar in young and older patients. The effect on systolic pressure was greater in older patients, perhaps because of greater baseline systolic pressure. Effects were similar in black patients and in white patients.

Effects in Chronic Stable Angina: The effectiveness of 5-10 mg/day of amlodipine in exercise-induced angina has been evaluated in 8 placebo-controlled, double-blind clinical trials of up to 6 weeks duration involving 1038 patients (684 amlodipine, 354 placebo) with chronic stable angina. In 5 of the 8 studies significant increases in exercise time (bicycle or treadmill) were seen with the 10 mg dose. Increases in symptom limited exercise time averaged 12.8% (63 sec) for amlodipine, 10 mg, and averaged 7.9% (38 sec) for amlodipine, 5 mg. Amlodipine, 10 mg also increased time to 1 mm ST segment deviation in several studies and decreased angina attack rate. The sustained efficacy of amlodipine in angina patients has been demonstrated over long-term dosing. In patients with angina, there were no clinically significant reductions in blood pressures (4/1 mmHg) or changes in heart rate (+0.3 bpm).

Atorvastatin

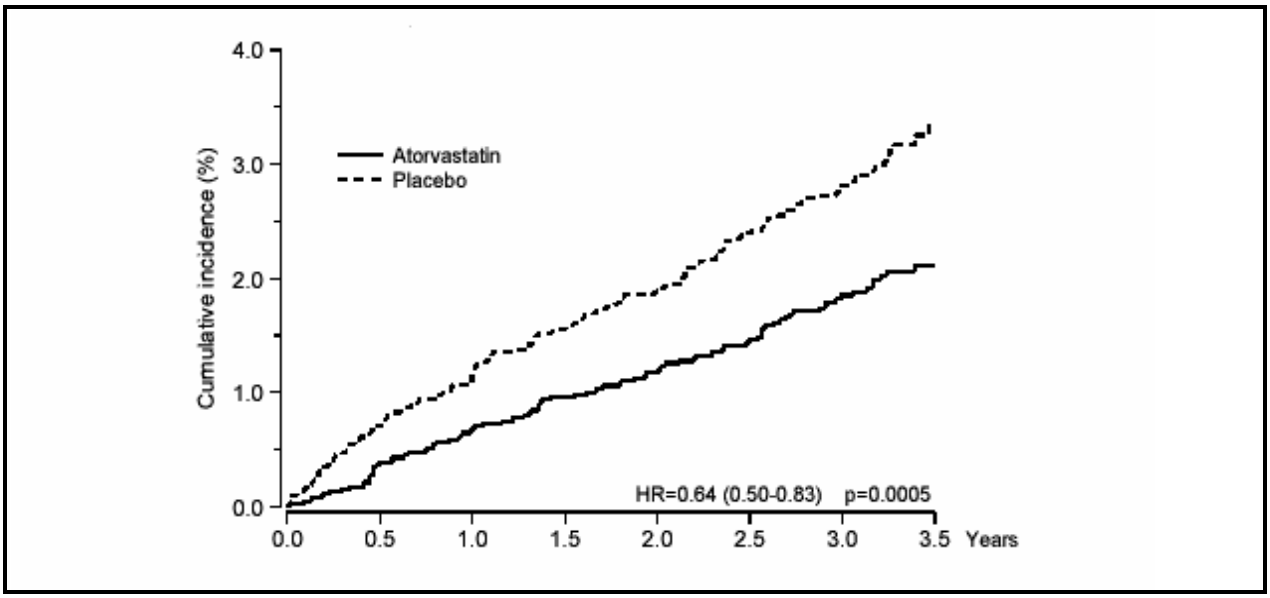
Prevention of Cardiovascular Disease

In the Anglo-Scandinavian Cardiac Outcomes Trial (ASCOT), the effect of atorvastatin on fatal and non-fatal coronary heart disease was assessed in 10,305 hypertensive patients 40-80 years of age (mean of 63 years), without a previous myocardial infarction and with TC levels ≤6.5 mmol/L. Additionally all patients had at least 3 of the following cardiovascular risk factors: male gender (81.1%), age ≥55 years (84.5%), smoking (33.2%), diabetes (24.3%), history of CHD in a first-degree relative (26%), TC:HDL ≥6 (14.3%), peripheral vascular disease (5.1%), left ventricular hypertrophy (14.4%), prior cerebrovascular event (9.8%), specific ECG abnormality (14.3%), proteinuria/albuminuria (62.4%). In this double-blind, placebo-controlled study, patients were treated with anti-hypertensive therapy (Goal BP <140/90 mm Hg for non-diabetic patients, <130/80 mm Hg for diabetic patients) and allocated to either atorvastatin 10 mg daily (n=5168) or placebo (n=5137), using a covariate adaptive method which took into account the distribution of nine baseline characteristics of patients already enrolled and minimized the imbalance of those characteristics across the groups. Patients were followed for a median duration of 3.3 years.

The effect of 10 mg/day of atorvastatin on lipid levels was similar to that seen in previous clinical trials.

Atorvastatin significantly reduced the rate of coronary events [either fatal coronary heart disease (46 events in the placebo group vs 40 events in the LIPITOR group) or nonfatal MI (108 events in the placebo group vs 60 events in the LIPITOR group)] with an absolute risk reduction of 1.1% and a relative risk reduction of 36% (based on incidences of 1.9% for atorvastatin vs 3.0% for placebo), p=0.0005 (see figure 1)]. This risk reduction yields a Number Needed to Treat of 311 patients per year. The risk reduction was consistent regardless of age, smoking status, obesity or presence of renal dysfunction. The effect of atorvastatin was seen regardless of baseline LDL levels. Due to the small number of events, results for women were inconclusive.

Figure 1: Effect of atorvastatin 10 mg/day on Cumulative Incidence of Nonfatal Myocardial Infarction or Coronary Heart Disease Death (in ASCOT-LLA)



Hypercholesterolemia

Atorvastatin has been shown to significantly improve lipid profiles in a variety of dyslipidemic conditions. Atorvastatin has been shown to be highly effective in reducing total and LDL-cholesterol, and triglycerides and apolipoprotein B in patients with primary hypercholesterolemia, familial and non-familial hypercholesterolemia, and mixed dyslipidemia, including familial combined dyslipidemia and patients with non-insulin dependent diabetes mellitus (NIDDM).

In 2 multicenter, placebo-controlled, double-blind, dose-response studies in patients with mild to moderate hypercholesterolemia (Fredrickson Types IIa and IIb), atorvastatin given as a single daily dose over 6 weeks reduced total-C, LDL-C, apo B, and TG; HDL was increased (Table 6). A therapeutic response was evident within 2 weeks, and the maximum response was usually achieved within 2-4 weeks.

Table 6 - Dose-Response in Patients With Mild to Moderate Hypercholesterolemia (Fredrickson Types IIa and IIb) (Mean Percent Change From Baseline)[a]

atorvastatin Dose (mg/day)	N	Total-C	LDL-C	Apo B	TG	HDL-C
Placebo	21	+4	+4	+3	+10	-3
10	22	-29	-39	-32	-19	+6
20	20	-33	-43	-35	-26	+9
40	21	-37	-50	-42	-29	+6
80	23	-45	-60	-50	-37	+5

[a] Results are pooled from 2 dose-response studies

In a pooled data set from 24 controlled clinical trials in patients with primary hypercholesterolemia (Type IIa) and mixed (combined) dyslipidemia (Type IIb), atorvastatin increased HDL-C by 5% to 8% from baseline at each dose tested (10, 20, 40, and 80 mg QD) (Table 7). In patients with HDL-C < 0.9 mmol/L (a condition often observed in persons with the metabolic syndrome) (see INDICATIONS AND CLINICAL USE), atorvastatin raised HDL-C 7% to 14%. These changes were independent of the dose administered. Atorvastatin also decreased total-C/HDL-C, LDL-C/HDL-C, and non-HDL-C/HDL-C ratios from baseline in a dose-dependent manner (Table 4). Atorvastatin (10, 20, 40 and 80 mg QD) increased HDL-C levels from baseline for both men and women.

Table 7 – Adjusted[a] Mean Percent Changes from Baseline in HDL-C, Total-C/HDL-C, LDL-C/HDL-C, Non-HDL-C/HDL-C, and HDL-C 0.9 mmol/L for Patients[b] With Mild to Moderate Hypercholesterolemia (Fredrickson Types IIa and IIb)

atorvastatin Dose (mg/day)	N (all patients)	HDL-C	Total-C/ HDL-C	LDL-C/ HDL-C	Non HDL-C/ HDL-C	HDL-C (baseline # 0.9 mmol/L) (N)
Placebo	250	+0.2‡	+2.8‡	+3.8‡	+3.5‡	+6.2* (17)
10	1871	+6.4	-29.3†	-37.0†	-35.5†	+13.8 (248)
20	147	+7.8	-36.0†	-44.1†	-43.0†	+8.3 (20)
40	115	+7.1	-38.9†	-49.6†	-47.1†	+8.6 (8)
80	318	+5.0	-43.5†	-55.3†	-52.4†	+7.1 (58)

[a] Least squares means from ANCOVA model with study, treatment and baseline
[b] Data pooled from 24 controlled studies
†significant linear dose trend
‡ significantly different from atorvastatin 10 mg ($p<0.01$)
* signficantly different from atorvastatin 10 mg ($p<0.05$)

In another multicenter, placebo-controlled, double-blind trial in patients with hypertriglyceridemia, atorvastatin lowered triglycerides in a dose-related manner, without causing a redistribution of triglycerides into various lipoprotein fractions (Table 8).

Table 8 - Efficacy in Patients With Hypertriglyceridemia (Mean Percent Change From Baseline)

atorvastatin Dose (mg/day)	N	VLDL-C	Total-C	VLDL-TG	LDL-C	TG	HDL-C	Apo B
Placebo	12	-2	+0.3	-6.6	+1.4	-5.3	+2.4	+2.7
5	11	-34.0*	-19.9*	-28.7	-12.7*	-27.3	+7.1	-15.4*
20	12	-46.0*	-33.1*	-35.7*	-31.1*	-33.7*	+10.6	-32.7*
80	11	-54-2*	-41.3*	-43.6*	-36.1*	-42.4*	+11.8*	-38.7*
* Significantly different from placebo, $p<0.05$								

Comparison of pooled data by Fredrickson types shows similar reductions for Type IIa and IIb patients in total-C, LDL-C and apo B; however, Type IIb patients, and Types IV patients experience a greater percent decrease in VLDL-C and TG levels (Table 9).

Table 9 - Efficacy in Patients by Fredrickson Type[a] (Mean Percent Change from Baseline)

	atorvastatin 10 mg/day		
Lipid Parameter	Type IIa (N = 935)	Type IIb (N = 550)	Type IV (N = 29)
LDL-C	-36	_35	-26
Apo B	-28	-28	-25
Total-Cl	-27	-27	-25
TG	-14	-24	-29
VLDL-C	-15	-28	-41
HDL-C	+6	+10	+13
Apo B/HDL-C	-31	-34	-33
Non-HDL-C/HDL-C	-37	-38	-38
[a] Pooled dataset			

A comparison of results in patients with heterozygous familial and non-familial hypercholesterolemia shows similar magnitudes of reductions in LDL-C, apo B and non-HDL-C/HDL-C ratio, in both patient populations (Table 10).

Table 10 - Efficacy in Heterozygous FH and Non FH Patients† (Mean Percent Change from Baseline)

Lipid Parameter	Phenotype	atorvastatin	
		10 mg/day	80 mg/day
LDL-C	Heterozygous FH	-36 (N=140)	-53 (N=154)
	Non FH	-36 (N=1215)	-52 (N=166)
Apo B	Heterozygous FH	-27 (N=134)	-46 (N=153)
	Non FH	-28 (N=1149)	-46 (N=144)
Non HDL-C/HDL-C Ratio	Heterozygous FH	-37 (N=140)	-53 (N=132)
	Non FH	-37 (N=1215)	-54 (N=166)

† Data from several studies

Comparison of results in patients with and without familial combined dyslipidemia (FCH) demonstrated that atorvastatin lowered LDL-C, apo B, total-C, VLDL-C, TG, and the non HDL-C/HDL-C ratio to a similar extent in both patient populations (Table 11).

Table 11 - Efficacy in Patients With and Without FCH†,[a] (Mean Percent Change from Baseline)

Lipid Parameter	atorvastatin 10 mg/day	
	FCH (N = 78-84)	Non-FCH (N = 1084-1224)
Total-C	-26%	-27%
LDL-C	-34%	-36%
TG	-21%	-17%
HDL-C	+8%	+7%
Apo B	-26%	-28%
VLDL-C	-25%	-18%
Non HDL-C/HDL-C Ratio	-36%	-37%
LDL-C/Apo B ratio	-9%	-11%

† Data from several studies

[a] The following criteria were used to define patients with FCH: first degree relative with lipid disorder, TG >250 mg/dL (>2.8 mmol/L), VLDL >45 mg/dL (>1.16 mmol/L), HDL <35 mg/dL (<0.9 mmol/L) (men) or <45 mg/dL (<1.16 mmol/L) (women).

In 3, double-blind, multicenter studies in patients with mild to moderate hypercholesterolemia, the number of patients meeting NCEP target LDL-C levels on atorvastatin was assessed over a 1-year period. After 16 weeks, between 46-74% of patients receiving 10 mg/day atorvastatin reached target LDL-C levels. The efficacy of atorvastatin (10 or 20 mg/day) was maintained over 52 weeks, with between 50-78% of patients achieving their LDL-C target levels.

The effect of atorvastatin was evaluated in comparative clinical trials with lovastatin, simvastatin and pravastatin. **For information on these results please refer to "REFERENCES".**

For more detailed clinical trial information please refer to the individual Product Monographs for **NORVASC** and **LIPITOR**.

DETAILED PHARMACOLOGY

Atorvastatin

Human Pharmacokinetics

Pharmacokinetic interaction studies have been conducted in healthy subjects with 3 macrolide antibiotics: erythromycin and clarithromycin (both of which inhibit CYP 3A4), and with azithromycin. Coadministration of atorvastatin with erythromycin or clarithromycin, resulted in moderately increased atorvastatin plasma levels but atorvastatin plasma levels were not altered by azithromycin. Twelve (12) healthy subjects were administered atorvastatin 10 mg on Days 1 and 15; erythromycin 500 mg QID was administered from days 8 to 19. Erythromycin increased atorvastatin C_{max} and AUC approximately 40%. In a second study, atorvastatin 10 mg was administered daily for 8 days; clarithromycin (500 mg BID) or azithromycin (500 mg QD) was coadministered from Days 6 - 8 (N=12/treatment). Coadministration with clarithromycin increased atorvastatin AUC ~80% and C_{max} ~50%, but atorvastatin plasma levels were not significantly altered by coadministration with azithromycin.

Steady-state, open-label, pharmacokinetic studies with digoxin have been performed in healthy subjects with both low and high doses of atorvastatin. Atorvastatin (10 mg or 80 mg QD; N=11 and N=12, respectively), was administered from days 1 - 20 and digoxin (0.25 mg QD) from Days 11 - 20. At steady-state, atorvastatin 10 mg daily had no significant effect on steady-state digoxin pharmacokinetics. However, following co-administration with atorvastatin 80 mg QD, the mean steady-state digoxin AUC and C_{max} increased 15% and 20%, respectively. Patients taking digoxin should be monitored appropriately.

The effect of amlodipine on the pharmacokinetics of atorvastatin was assessed at steady-state in a randomised, open-label, placebo-controlled, crossover study in healthy male subjects (N=16). Atorvastatin (80 mg QD) was administered with amlodipine (10 mg QD) or placebo from Days 1-8. Following a 14 day washout, the alternate combination was administered from Days 22 - 29. At steady-state, the coadministration of maximum doses of atorvastatin and amlodipine did not significantly alter the pharmacokinetics of atorvastatin and there were no apparent changes in blood pressure or heart rate.

The effect of quinapril on the pharmacokinetics of atorvastatin was assessed in a randomized, open-label study in healthy volunteers (N=22). Single doses of atorvastatin (10 mg) were administered on Days 1 to 14, and single doses of quinapril (80 mg) were administered on days 1 to 7 or Days 8 to 14. The mean T_{max} value for atorvastatin during steady state quinapril administration was shortened by 1.25 hours compared to that of atorvastatin administered alone, but with no change in absorption/AUC or C_{max}. No significant changes in blood pressure or heart rates were observed.

Concomitant administration of atorvastatin 20 mg to 40 mg and itraconazole 200mg daily resulted in a 2.5 – 3.3-fold increase in atorvastatin's AUC.

Concomitant administration of atorvastatin 10 mg and cyclosporine 5.2 mg/kg/day resulted in a 7.7 fold increase in exposure to atorvastatin.

For more detailed pharmacology information please refer to the individual Product Monographs for **NORVASC** and **LIPITOR**.

TOXICOLOGY

Amlodipine

ACUTE TOXICITY - Amlodipine (as maleate unless otherwise indicated)

			LD_{50}	Range of Lethal Doses (mg/kg)	
SPECIES	**SEX**	**ROUTE**	**base/mg/kg**	**No Deaths**	**All Dead**
Mice	M	p.o.	N.D.	10	40
	F	p.o.	N.D.	10	40
	M	i.v.	N.D.	2.5	10
	F	i.v.	N.D.	2.5	10
Rats	M	p.o.	150	2/10 at 100	400
	F	p.o.	140	2/10 at 100	250
	M	i.v.	N.D.	1	10
	F	i.v.	N.D.	1	10
Rats*	M	p.o.	393**		
	F	p.o.	686**		

* Sprague Dawley Rats from Shizouka Lab Animal Centre, Hamamatsu, Japan
** Besylate Salt
\+ Dogs from Interfauna, France
++ Dogs from Japan
N.D. Not Determined: The result did not permit calculations of LD50 values. Thus, range of lethal doses is given.

The main clinical signs in the oral studies were somnolence, decreased spontaneous movement and for rats salivation, dyspnea, ptosis, lacrimation, blanching, cyanosis, rough coat, abdominal distension, and eventually coma. After i.v. injection, the animals died rapidly showing only somnolence, tachypnea or ptosis.

SPECIES	ROUTE	DOSE base mg/kg/day	ANIMAL PER DOSE LEVEL	DURATION	FINDINGS
MAXIMUM TOLERATED DOSE (SINGLE)					
Dog	Oral (gavage)	4 8 16	2 M	Single Dose	At all dose levels: Vasodilation and increases in plasma aldosterone levels. At 4 mg/kg: Compensatory tachycardia. At 8 mg/kg: In 1 of 2 dogs vomiting, sedation, respiratory distress and diarrhea 48 hr post-dose; normal at day 5. Compensatory tachycardia. At 16 mg/kg: Moribund with hyperthermia within 24 hours; low blood pressure returned to normal over 2-6 days; transient raise in heart rate. Histological examination showed congestion, edema and hemorrhage of the right atrial wall in the 2 dogs at 16 mg/kg. The hemorrhage in the right atrial wall corresponds to the right atrial lesions seen in long-term studies with amlodipine and other vasodilators (see long-term toxicity). One of 2 dogs at each dose showed fibrosis of the left ventricle in the subendocardial region and the posterior papillary muscle. The maximum tolerated dose was not determined.
Dog (Japanese Study)	Oral	3.5 7	1 M 1 F	Single Dose	Mortality: 1 male dog at 7 mg/kg. Decreased spontaneous movement and flushing of palpebral conjunctiva and buccal cavity. At 7 mg/kg: 1 female vomiting; 1 male hypothermia, lying prone. Hematology/Clinical Chemistry: Increase in WBC and BUN at 10 and 5 mg/kg (males). The maximum tolerated dose was not determined.

SUBACUTE AND CHRONIC TOXICITY

SPECIES	ROUTE	DOSE base mg/kg/day	ANIMAL PER DOSE LEVEL	DURATION	FINDINGS
Mouse	Oral (diet)	0 2.5 5 10	10 M 10 F	2 Months	At 10 mg/kg/day: Mice died during week 2 of the study. At 5 mg/kg/day (males and females) and 2.5 mg/kg/day (males): Increase in water consumption. At 5 mg/kg/day - Pathology: Drug-related increases in heart and liver weights.
Rat (Japanese Study)	Oral (gavage)	0 4 16 32 64	12 M 12 F	1 Month	At 64 mg/kg/day: All rats died within 9 days. At 32 mg/kg/day: 12/24 rats died; decreased food consumption, growth inhibition, ptosis, decreased spontaneous movement. At 16 and 32 mg/kg/day: The pattern of results on heart weights, increased urinary volume, effect on electrolyte balance and the adrenals was similar to that of the 6 month study below; increase in BUN at 16 mg/kg (males) and at 32 mg/kg (males and females).

SPECIES	ROUTE	DOSE base mg/kg/day	ANIMAL PER DOSE LEVEL	DURATION	FINDINGS
Rat (Japanese Study)	Oral (gavage)	0 2 7 21	16 M 16 F	3 Months followed by 1 Month drug withdrawal	21 mg/kg/day: Salivation, growth inhibition, increased BUN, increased urinary volume, effect on electrolyte balance and adrenals was similar to that of the 6 month study below. Also post-mortem dilation of small intestine without morphological lesions. At 7 mg/kg/day: Alterations in urinary electrolytes excretion. No drug related effects at the end of 1 month drug withdrawal phase.
Rat	Oral (gavage)	0 2.5 5 10	20 M 20 F	6 Months	At all dose levels: Renal effects: increased urinary volume and/or Na/K/Cl excretion, decreased plasma Na/K and/or Ca/Cl and increased urea; Post-mortem: Increase in heart weights. At 10 mg/kg/day: Renal effects: increased kidney weight. Histopathology: Thickening of zona glomerulosa at 5 and 10 mg/kg/day.
Rat (Japanese Study)	Oral (gavage)	1.4 7 18	30 M 30 F	12 Months	(interim sacrifice 5/sex/group after 6 months) Mortality: 3 rats (2 males and 1 female) at 18 mg/kg/day. At 18 mg/kg/day: Salivation, growth inhibition; Renal effects: increase in urinary volume with increased electrolytes excretion and decreased serum electrolytes; increase in BUN. At 7 mg/kg/day: Growth inhibition (males); Renal effects: increases of urinary volume and electrolyte excretion. Post-mortem: Increases of adrenal weights (at 18/mg/kg), increases of relative heart weight (18 and 7 mg/kg), dilated small intestines without morphological change (18 mg/kg). Histopathology - Main Finding: Enlargement of the zona glomerulosa of the adrenals (18 and 7 mg/kg).
Dog	Oral (gavage)	0.5 to 4	2 M 2 F	10 Days	Supplementary Dose Escalation Study (0.5 mg/kg/day) At 4 mg/kg: Death of all (4/4) dogs preceded in 3 dogs by low systolic blood pressure, bradycardia, disturbances of heart rhythm and conduction. Clinical signs included pale skin, hypothermia and prostration. Histopathology: Showed foci of myocyte necrosis and sarcoplasmic vacuolation in the left ventricle, papillary muscle and left and right atria. Congestion and/or edema in several organs (i.e. gastrointestinal tract/gall bladder wall and surrounding tissues as well as the connective tissue surrounding both kidneys).
Dog	Oral	0 0.25 0.5 1	3 M 3 F	6 Months	At all dose levels: Increase in urinary volume and urinary excretion of electrolytes (not dose-related). Reduction in blood pressure and increases in heart rate. At 1 mg/kg/day - Pathology: Increase in relative heart weights in 4/6 dogs, inflammatory lesion of the right atrial wall was seen which was considered to be consequence of excessive hemodynamic changes.
Dog	Oral	0 0.125 0.25 0.5	4 M 4 F	12 Months	At 0.5 mg/kg/day: Reduction in blood pressure and increases in heart rate; increase in urinary volume and urinary excretion of electrolytes (females). At 0.5 mg/kg/day - Pathology: Showed inflammatory lesions of the right atrial wall in 1/8 dogs, similar to that of the 6 month study above, and diffuse gingival hyperplasia.

MUTAGENICITY

Study	Test Organism	Dose	Route	Major Findings
Ames Test (modified) Quantitative Plate Assay (QAP) and Metabolic Activation (MA) with Hepatic Microsomes	salmonella typhimurium: Strains TA 1535, TA 1537, TA 98 and TA 100	10-0.02 mg/plate (QAP) 0.2-0.0005 mg/plate (MA)	In-vitro	No evidence of mutation frequency.
In-vivo Cytogenetic Tests	mouse bone marrow	20 mg/kg single dose 10 mg/kg/day for 5 days	In-vivo p.o. s.c.	No indication of chromosome breakage or mutagenicity observed.
In vitro Cytogenetic Tests with or without metabolic activation [rat liver microsomal enzymes (S-9)]	human lymphocytes	Without metabolic activation: 0.01 to 1000 :g/Ml of culture medium With metabolic activation: 1.0 to 25 :g/Ml of culture medium.	In-vitro	Non-activation: No evidence of induced chromosome breakage observed at levels of 1.0 :g/mL and below. At levels higher that 1.0 :g/mL, compound produced mitotic inhibition. Activation: No drug induced clastogenic activity observed at levels up to 10 :g/mL. Higher levels produced mitotic inhibition.
Quantitative Plate Assay (QAP) of Mouse Urine	Salmonella typhimurium Strains: TA 1535, TA 1537, TA 98 and TA 100.	0, 1, 10 and 20 mg/kg	In-vivo p.o.	No incidence of an excreted mutagen.
L 5178Y/TK +/- Gene Mutation Assay with and without liver S-9 fraction	mouse lymphoma cells	1.2 - 38 :g/mL	In-vitro	No evidence of gene mutational activity.

CARCINOGENICITY

There was no evidence of a carcinogenic effect when amlodipine was administered in the diet for up to 24 months to rats up to 2.5 mg/kg/day. Amlodipine was also administered for up to 24 months of dietary administration to mice at doses up to 2.5 mg/kg/day and no evidence of carcinogenicity was observed.

REPRODUCTION AND TERATOLOGY

Species	Route	Dose base/mg/kg/day	Animal per Dose Level	Duration	Findings
Fertility					
Rat (SD) (Japanese Study)	Oral (gavage)	0 1.4 7 18	24 M + 24 F	Males 71 days prior to and during mating. Females 14 days prior to and during mating and up to 7 days of gestation.	At 18 mg/kg: Impairment of body weight gain (females). There were no effects of the drug on copulation or pregnancy rates, nor any evidence of embryotoxicity or teratogenicity.
Teratology					
Rat (Charles River CD/SD)	Oral (gavage)	0 2 5 10	20 F	Days 6-15 post insemination. Hysterectomies on day 20 of gestation.	No effects were observed.
Rat (SD) Japanese Study	Oral (gavage)	3 7 18	34 F	Days 7-17 post-insemination. b of dams sacrificed on day 21 of gestation. F_1 generation followed.	No effects were observed except in the dams. At 18 mg/kg: Reduction in food intake and body weight gain.
Rabbit (Japanese White) Japanese Study	Oral	3 7 18	18 or 19 F	Day 6 to day 18 of gestation.	At 18 and 7 mg/kg: Decrease in maternal body weight (18 mg/kg) decrease in food consumption (18 and 7 mg/kg). No evidence of drug induced fetotoxicity or teratogenicity.
Peri- and Post-Natal					
Rat (SD) Japanese Study	Oral (gavage)	0 1.4 2.8 7.0	25 F	Day 17 of gestation to day 21 post-partum.	As in the combined Fertility/Perinatal Study above; at the high dose level (7.0 mg/kg/day) adverse effects were observed on parturition and number of viable pups at birth and day 4 post-partum.

Atorvastatin

Acute Toxicity

The acute toxicity of atorvastatin following single doses was evaluated in mice, rats and dogs by oral and intravenous routes, and the results are summarized below:

Table 12 - Acute Oral and Intravenous Toxicity Studies with Atorvastatin

Species	Sex	Route	Dose Range (mg/kg)	Results
Mouse	Male/Female	Oral	200-5000	No Deaths
Mouse	Male/Female	IV	0.4 - 4	No Deaths
Rat	Male/Female	Oral	200-5000	No Deaths
Rat	Male/Female	IV	0.4 - 4	No Deaths
Dog	Male/Female	Oral	10 - 400	No Deaths
Dog	Male/Female	IV	0.4 - 4	No Deaths

The acute toxicity of atorvastatin in rodents and dogs is low. Oral median lethal doses in mice and rats are greater than 5000 mg/kg.

Subacute and Chronic Toxicity Studies

The target organs affected by atorvastatin in multiple dose toxicity studies in rats (2 weeks to 52 weeks), and dogs (2 weeks to 104 weeks) are summarized in the table below. The spectrum of effects observed is not unexpected in view of the magnitude of the dose levels used, potency of atorvastatin in inhibiting mevalonate synthesis and the essential role of HMG-CoA reductase in maintaining cellular homeostasis.

Table 13 - Atorvastatin: Target Organs Affected in Animal Studies

Rat		Dog
Liver		Liver
Stomach (non-glandular)		Gallbladder
Skeletal Muscle		Skeletal Muscle
		Intestine
		Brain/Optic Nerve*

* Occurred after administration of high, intolerable doses (280 mg/kg)

The following table summarizes the significant adverse changes observed during long-term toxicology studies in rats (52 weeks) and dogs (104 weeks):

Table 14 - Atorvastatin: Significant Adverse Changes in Chronic Studies

Species/Results	**Minimal Toxic Dose (mg/kg/day)**		**No-Effect Dose (mg/kg/day)**
RAT			
Hepatocellular atypia	70		5
Bile Duct hyperplasia[1]	125		70
Nonglandular stomach acanthosis	125		70
DOG			
Death[2]	120		40
Hepatocellular granulomata[3]	10		ND
Hepatocellular necrosis[3]	120		40
Gallbladder edema/hemorrhage[3]	120		40
Bile duct hyperplasia[3]	120		10
Intestinal ulcers and single cell necrosis[3]	120		40
Skeletal muscle (tongue) necrosis[2]	120		40

1 Present only at Week 26; not observed at Week 52.
2 Findings occurred in Week 7 or 9.
3 Findings occurred at Week 52 or in moribund dogs, were less pronounced after a 12-week withdrawal period (Week 64), and were not observed after 104 weeks of dosing.
ND = Not determined

The results of the long-term toxicology studies with atorvastatin indicated that similar to other HMG-CoA reductase inhibitors, the liver is the primary target organ. This is expected since the liver is the primary site of the pharmacologic action of atorvastatin and it is subject to the greatest drug exposure following oral administration. In both the rat and dog studies, the hepatic changes diminished with time (i.e. effects were less pronounced at the end of the 52-week and 104-week studies) suggesting an adaptive response.

Brain hemorrhage, optic nerve degeneration, lenticular opacities and testicular degeneration were not seen in dogs treated for 104-weeks with atorvastatin up to 120 mg/kg/day.

Carcinogenicity and Genotoxicity Studies

Atorvastatin was not carcinogenic in rats given 10, 30 or 100 mg/kg/day for 2 years. The 100 mg/kg dose is 63-fold higher than the maximum recommended human dose of 80 mg (1.6 mg/kg, based on a 50 kg human) and AUC (0-24 hr) values were 8- to 16-fold higher.

In a 2-year study in mice given 100, 200 or 400 mg/kg/day, incidences of hepatocellular adenoma in males and hepatocellular carcinoma in females were increased at 400 mg/kg. This dose is 250 times the maximum recommended human dose on a mg/kg basis and systemic exposure based on AUC (0-24 hr) was 6 to 11 times higher. There was no evidence of

treatment-related increases in tumor incidences at the lower doses of 100 and 200 mg/kg/day (i.e. up to 125 times the maximum recommended human dose on a mg/kg basis and systemic exposures of 3 times higher based on AUC (0-24 hr).

Atorvastatin did not demonstrate mutagenic or clastogenic potential in four in vitro tests with and without metabolic activation or in one in vivo assay. It was negative in the Ames test with *Salmonella typhimurium* and *Escherichia coli*, and in the in vitro HGPRT forward mutation assay in Chinese hamster lung cells. Atorvastatin did not produce significant increases in chromosomal aberrations in the in vitro Chinese hamster lung cell assay and was negative in the in vivo mouse micronucleus test.

Reproductive and Teratogenicity Studies

No adverse effects on fertility or reproduction were observed in male rats given doses of atorvastatin up to 175/mg/kg/day or in female rats given doses up to 225 mg/kg/day. These doses are 100 to 140 times the maximum recommended human dose on a mg/kg basis.

Atorvastatin did not cause any adverse effects on sperm or semen parameters, or in reproductive organ histopathology in dogs given doses of 10, 40 or 120 mg/kg for 2 years. Atorvastatin was not teratogenic in either rats or rabbits.

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Amlodipine

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PART III: CONSUMER INFORMATION
PrCADUET®
(amlodipine besylate and atorvastatin calcium tablets)

This leaflet is Part III of a three-part "Product Monograph" published when CADUET was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about CADUET. Contact your doctor or pharmacist if you have any questions about this medication. Please read this information carefully.

ABOUT THIS MEDICATION

What is CADUET used for:

CADUET is a product that contains two active ingredients: amlodipine and atorvastatin and is to be used when your doctor deems it appropriate to use both medications.

Your doctor has prescribed CADUET to you to control your hypertension (high blood pressure) and/or to prevent angina attacks (chest pains) and to help lower your cholesterol or other fats in the blood (such as triglycerides). Even if you do not have high cholesterol, your doctor may still prescribe CADUET if you have high blood pressure and other risk factors in order to prevent your risk of cardiovascular disease such as heart attacks.

CADUET is just part of the treatment your doctor will plan with you to help keep you healthy. Depending on the condition of your health and your lifestyle, your doctor may recommend:
- a change in your diet to control your weight, reduce your cholesterol, reduce intake of saturated fats and increase fiber
- exercise that is right for you
- quitting smoking or avoiding smoky places
- giving up alcohol or drinking less

CADUET is not recommended in children.

What CADUET does:

The amlodipine portion of CADUET belongs to a class of medications called calcium channel blockers. Calcium channel blockers, like amlodipine, block the transfer of calcium into the cells of the heart and blood vessels. This helps the blood vessels to relax, thereby lowering blood pressure and resulting in less work for the heart.

The atorvastatin portion of CADUET belongs to the class of medicines known as "statins", more specifically called HMG-CoA reductase inhibitors. HMG-CoA reductase is an enzyme involved in regulating cholesterol levels in your body. Statins are used along with changes to exercise and diet to help control the amount of cholesterol produced by the body. High levels of cholesterol and other fats can cause heart disease by clogging the blood vessels that feed blood and oxygen to the heart.

Atorvastatin can help your body:
- Decreases LDL (bad) cholesterol, triglyceride levels and other lipids/fats in the blood.
- Increase HDL (good) cholesterol
- Decrease the Total Cholesterol HDL-Cholesterol Ratio (TC: HDL-C Ratio). The ratio represents the balance between bad and good cholesterol.

Follow your doctor's instructions carefully.

When CADUET should not be used:

Do not take CADUET if you:

- are allergic to either amlodipine besylate and/or atorvastatin calcium or other medications known as dihydropyridines (for example, felodipine, and nifedipine) or any of the nonmedicinal ingredients (see What the nonmedicinal ingredients are).

- have been diagnosed with low blood pressure (less than 90 mmHg systolic).

- have active liver disease or unexplained increases in liver enzymes.

- are pregnant or breast-feeding.

What the medicinal ingredients are:

amlodipine besylate and atorvastatin calcium.

What the nonmedicinal ingredients are:

calcium carbonate, colloidal silicon dioxide (anhydrous), croscarmellose sodium, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, opadry II white 85F28751 or opadry II blue 85F10919, polysorbate 80, pregelatinized starch, and purified water.

What dosage forms it comes in:

CADUET tablets are available in 8 combinations of amlodipine/ atorvastatin: 5mg/10mg, 5mg/20mg, 5mg/40mg, 5mg/80mg, 10mg/10mg, 10mg/20mg, 10mg/40mg, 10mg/80mg. CADUET tablets which contain 5 mg amlodipine will be film-coated white, and those with 10 mg amlodipine will be film-coated blue.

WARNINGS AND PRECAUTIONS

Serious Warnings and Precautions
Tell your doctor if you have any muscle pain, tenderness, soreness or weakness during treatment with CADUET.

Never change the dose of CADUET unless your doctor tells you to.

Before you use CADUET talk to your doctor or pharmacist if

- you have thyroid problems
- you have had a stroke or a mini stroke called a transient ischemic attack (TIA)
- you regularly drink *three or more* alcoholic drinks daily
- you are taking any other cholesterol lowering medication such as fibrates (gemfibrozil, fenofibrate), niacin or ezetimibe
- you are taking any other prescription, non-prescription or over-the-counter products especially those listed under the section "Interactions With This Medication"
- you have a family history of muscular disorders
- you had any past problems with the muscles (pain, tenderness), after using an HMG-CoA reductase inhibitor ("statin") such as atorvastatin LIPITOR®), fluvastatin (LESCOL®), lovastatin (MEVACOR®), pravastatin (PRAVACHOL®), rosuvastatin (CRESTOR®) or simvastatin (ZOCOR®)
- you have kidney or liver problems
- you have diabetes (as the dosage of CADUET may need to be adjusted)
- you have undergone surgery or other tissue injury
- you do excessive physical exercise
- you are 65 years old or older
- you have taken any of the following medicines before, and you had an allergic reaction:
- amlodipine or other derivatives
  - nifedipine (e.g. ADALAT XL®)
  - felodipine (e.g. PLENDIL®, RENEDIL®)
- atorvastatin (atorvastatin) or other derivatives
  - simvastatin (e.g. ZOCOR®)
  - lovastatin (e.g. MEVACOR®)
  - pravastatin (e.g. PRAVACHOL®)
  - fluvastatin (e.g. LESCOL®)
  - rosuvastatin (e.g. CRESTOR®)
- you are taking cyclosporine (e.g. SANDIMMUNE®, NEORAL®)

Slightly increased blood sugar can occur when you take CADUET. Discuss with the doctor your risk of developing diabetes.

CADUET may cause muscle pain, aching or weakness that does not go away even after stopping the drug.

Pregnancy:

- Before using this medication, discuss the following with your doctor:
  - If you are breast-feeding your baby, you should not take CADUET. This medicine may be present in your breast milk.
  - Cholesterol compounds are essential elements for the development of a fetus. Cholesterol-lowering drugs can harm the fetus. If you are of child-bearing age, discuss with your doctor the potential hazards to the fetus and the importance of birth control methods.
  - This medicine should not be used by pregnant women. If you become pregnant, discontinue use immediately and discuss with your doctor.

INTERACTIONS WITH THIS MEDICATION

As with most medicines, interactions with other drugs are possible. Tell your doctor, nurse, or pharmacist about all the medicines you take, including drugs prescribed by other doctors, vitamins, minerals, natural supplements, or alternative medicines. Also mention if you drink alcoholic beverages.

Drugs that may interact with CADUET include:

- corticosteroids (cortisone-like medicines)
- cyclosporine (e.g. SANDIMMUNE®, NEORAL®), tacrolimus
- gemfibrozil (LOPID®)
- fenofibrate (e.g. LIPIDIL MICRO®)
- bezafibrate (e.g. BEZALIP®)
- lipid-modifying doses of niacin (nicotinic acid)
- spironolactone
- cimetidine
- erythromycin, clarithromycin or azole antifungal agents (ketoconazole or itraconazole)
- nefazodone (e.g. SERZONE®)
- indinavir sulfate (e.g. CRIXIVAN®), nelfinavir mesylate (e.g. VIRACEPT®), ritonavir (e.g. NORVIR®), saquinavir mesylate

(INVIRASE™), lopinavir/ritonavir (e.g. KALETRA®), telaprevir (INCIVEK™), tipranavir (APTIVUS®), darunavir (PREZISTA®), fosamprenavir (TELZIR ®), boceprevir (VICTRELIS®)
- fusidic acid (e.g. FUCIDIN)
- digoxin
- diltiazem
- efavirenz, rifampin
- sildenafil (VIAGRA®) may lower your blood pressure
- beta-blockers (medicines used to lower your blood pressure)
- antacids (frequent use) and CADUET should be taken 2 hours apart
- St. John's Wort (hypericum perforatum)
- grapefruit juice
- colchicine

Certain non-prescription drugs may be harmful to you or may interfere with CADUET.

PROPER USE OF THIS MEDICATION

Take CADUET exactly as prescribed by your doctor, nurse or pharmacist. Swallow CADUET with water.

Usual Dose: The dose range for CADUET is 5/10 to 10/80 mg once daily. The maximum dose of CADUET is 10/80 mg once daily. CADUET is available in a broad dose-range. Your doctor will decide on the best dose for you. Your individual health and tolerance to medications will help your doctor decide how much of each ingredient will be best for you.

Follow your doctor's dosing instructions carefully. Take your CADUET as a single dose once a day. Never change the dose unless your doctor tells you to. It does not matter if you take CADUET with food or without food. Ideally, so you won't forget, you should get in the habit of taking your medicine at the same time every day. Follow the plan that you and your doctor make for diet, exercise, weight control and smoking cessation.

We often cannot see or feel the problems that high cholesterol and high blood pressure cause until a lot of time has passed or until a major event like a heart attack occurs. You and your doctor will be following your cholesterol and blood pressure levels to bring them down and keep them in a safe range. Here are some important tips to consider:
- Report to all follow-up visits scheduled by your doctor. This is important to follow through with all lab tests requested by your doctor. These tests will help to follow/track your health in general, for example, your kidneys, liver and blood sugar levels.
- It is important to take all of your medication and to refill your prescription on time so as not to miss any doses.
- Don't drink too much alcohol while you are taking CADUET. Talk to your doctor about how much is too much for you.
- If you get sick, have an operation, or need medical treatment, inform your doctor or pharmacist that you are taking CADUET.
- If you have to take any other medicine – prescription, non-prescription or over-the-counter - while you are taking CADUET, talk to your doctor or pharmacist first.
- If you have to see a different doctor for any reason, be sure to inform him/her that you are taking CADUET.

Missed Dose:
If you miss taking your medicine, take it as soon as you can. But if it is almost time for your next dose, skip the missed dose and just take the next dose. Don't take a double dose.

Overdose:

If you think you have taken too much of CADUET contact your doctor, nurse, pharmacist, hospital emergency department or regional Poison centre immediately, even if there are no symptoms.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Along with its intended action, any medication may cause unwanted effects. CADUET is generally well tolerated. However, check with your doctor or pharmacist promptly if any of the following persist or become troublesome:
- stomach pain or upset, vomiting or throwing up, loss of appetite and inability to eat, or malaise (general feeling of being unwell), burping
- gas, constipation, diarrhea
- headache, neck pain
- fever
- hair loss
- skin rash, hives, itchiness
- insomnia (difficulty sleeping), drowsiness, fatigue, nightmares
- impotence (inability of develop or maintain an erection of the penis)
- blurred vision, ringing in the ears

Possible side effects reported with some statins:
- breathing problems including persistent cough and/or shortness of breath or fever

- mood related disorders including depression
- poor memory, confusion and memory loss

CADUET can cause abnormal blood test results. Your doctor will decide when to perform blood tests and will interpret the results.

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

Symptom/Effect		Talk with your doctor or pharmacist		Stop taking drug and seek immediate emergency medical attention
		Only if severe	In all cases	
Common	Swelling of the ankles	✓		
Rare	Pancreatitis (severe upper abdominal pain that radiates to the back)		✓	
	Muscle pain that you cannot explain		✓	
	Muscle cramps, tenderness or weakness		✓	
	Generalized weakness, especially if you don't feel well		✓	
	Brownish or discoloured urine		✓	
	Increased frequency, severity, duration of angina (chest pains)		✓	
	Low blood pressure: (dizziness, or dizziness when rising from sitting or laying down)	✓		
	Change in rhythm or pace of heart beats		✓	
	Numbness or tingling in hands/fingers	✓		
	Shortness of breath		✓	
	Abnormal vision		✓	
	Jaundice (yellowing of the skin and eyes) from a liver disorder called hepatitis (inflammation of the liver)		✓	
Unknown	Increased blood sugar: frequent urination, thirst and hunger	✓		
	Extrapyramidal symptoms: muscle stiffness, body spasms, upward eye rolling, exaggeration of reflexes, drooling, difficulty moving how and when you want			✓

This is not a complete list of side effects. If you notice anything unusual or any unexpected effects while taking CADUET, contact your doctor or pharmacist.

HOW TO STORE IT

Keep your medicine at 25°C; excursions are permitted to 15-30°C (room temperature), away from warm and damp places, like the bathroom or kitchen. **Always keep medicine well out of the reach of children.**

REPORTING SUSPECTED SIDE EFFECTS

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at www.healthcanada.gc.ca/medeffect
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
  - Fax toll-free to 1-866-678-6789, or
  - Mail to: Canada Vigilance Program
    Health Canada
    Postal Locator 0701E
    Ottawa, Ontario
    K1A 0K9

  Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada Web site at www.healthcanada.gc.ca/medeffect.

NOTE: Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

Please note that the information contained in the enclosed pamphlet is general. Your doctor and pharmacist are your primary sources of information about your health and the medicine you take. Consult with your doctor or pharmacist if you have questions about your health, any medication you take or the information we are providing you.

This document plus the full Product Monograph, prepared for health professionals can be found at: http://www.pfizer.ca or by contacting the sponsor, Pfizer Canada Inc., at: 1-800-463-6001

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