

Therapeutic Class Overview

Statins (HMG-CoA Reductase Inhibitors)

INTRODUCTION

- The 3-hydroxy-3-methylglutaryl-coenzyme A (HMG-CoA) reductase inhibitors (also known as statins) include single entity agents (atorvastatin, fluvastatin, lovastatin, pitavastatin, pravastatin, rosuvastatin, and simvastatin), as well as fixed-dose combination products (amlodipine/atorvastatin, ezetimibe/atorvastatin, and ezetimibe/simvastatin). The statins work by inhibiting HMG-CoA reductase, which is the rate-limiting enzyme involved in hepatic cholesterol synthesis. This enzyme catalyzes the conversion of HMG-CoA to mevalonate, which is a cholesterol precursor. Inhibition of HMG-CoA reductase decreases hepatic cholesterol synthesis, causing up-regulation of low-density lipoprotein cholesterol (LDL-C) receptors. Statins also decrease the release of lipoproteins from the liver.
- The statins are the most effective class of oral drugs to lower LDL-C. Depending on the agent selected, moderate-intensity statins can decrease LDL-C by 30 to 49% and high-intensity statins can decrease LDL-C levels $\geq 50\%$. The effects on LDL-C are dose-dependent and log-linear. Statins also decrease triglycerides (TG) and increase high-density lipoprotein cholesterol (HDL-C) by varying levels (Stone et al, 2014).
- Ezetimibe inhibits the intestinal absorption of cholesterol, which decreases the delivery of cholesterol to the liver. This causes a reduction of hepatic cholesterol stores and an increase in clearance of cholesterol from the blood.
- Amlodipine is a calcium channel blocker that is approved for the treatment of hypertension (HTN), chronic stable angina and vasospastic angina, as well as to reduce the risks of hospitalization or revascularization in patients with angiographically confirmed coronary artery disease (CAD).
- Statins that are included in this review are listed in Table 1. All products are now available in a generic formulation except for ALTOPREV® (lovastatin extended-release tablet), FLOLIPID (simvastatin oral suspension), LIVALO (pitavastatin tablet), and ZYPITAMAG (pravastatin tablet) (Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations, 2018).
- The combinations niacin/lovastatin (ADVICOR®) and niacin/simvastatin (SIMCOR®) were removed from the market because the Food and Drug Administration (FDA) determined that a reduction in TG and increase in HDL-C do not contribute to decreased cardiovascular events according to the newest evidence (AbbVie, 2016).
- The agents included in this review are listed in Table 1 by brand name. Since there are some branded agents that contain the same generic component, the remaining tables in the review are organized by generic name.

Table 1. Medications Included Within Class Review

Drug	Manufacturer	FDA Approval Date	Generic Availability
ALTOPREV (lovastatin extended-release)	Covis Pharma	06/26/2002	-
CRESTOR (rosuvastatin)	AstraZeneca Pharmaceuticals	08/12/2003	✓
FLOLIPID (simvastatin oral suspension)	Salerno Pharmaceuticals LP	04/21/2016	-
LESCOL (fluvastatin)*	Novartis	12/31/1993	✓
LESCOL XL (fluvastatin extended-release)	Novartis	10/06/2000	✓
LIPITOR (atorvastatin)	Pfizer	12/17/1996	✓
LIVALO, ZYPITAMAG (pitavastatin)€	Kowa Company (LIVALO) Medicure (ZYPITAMAG)	08/03/2009	-
MEVACOR (lovastatin)*	Merck & Co., Inc	08/31/1987	✓

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Drug	Manufacturer	FDA Approval Date	Generic Availability
PRAVACHOL (pravastatin)	Bristol Myers Squibb Company	10/31/1991	✓
ZOCOR (simvastatin)	Merck & Co., Inc.	12/31/1991	✓
CADUET (amlodipine/atorvastatin)	Pfizer	01/30/2004	✓
LIPTRUZET† (ezetimibe/atorvastatin)	Watson Labs Teva	04/26/2017	✓
VYTORIN® (ezetimibe/simvastatin)	Merck & Co., Inc.	07/23/2004	✓

*The brands, LESCOL and MEVACOR, have been discontinued, but the generic formulations are available.

€The brand NIKITA was discontinued.

†The brand, LIPTRUZET, by Merck was discontinued in 2015. A generic formulation by Watson Labs Teva was recently approved by the FDA, however, current market availability is unknown.

(Drugs@FDA, 2018; Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations, 2018)

INDICATIONS
Table 2. FDA-approved indications

Indications	Single-Entity Agents							Combination Products		
	atorvastatin	fluvastatin	lovastatin	pitavastatin	pravastatin	rosuvastatin	simvastatin	amlodipine/ atorvastatin	ezetimibe/ atorvastatin	ezetimibe/ simvastatin
Hypertriglyceridemia										
Reduce elevated TG in patients with hypertriglyceridemia							✓			
Treatment of adult patients with hypertriglyceridemia in combination with diet	✓				✓	✓		✓ (atorvastatin)		
Primary Hypercholesterolemia and Mixed Dyslipidemia										
Reduce elevated total cholesterol (TC), LDL-C, apolipoprotein B (apo B), and TG and to increase HDL-C in patients with primary hyperlipidemia or hypercholesterolemia and mixed dyslipidemia	✓	✓	✓ § (ER)	✓	✓	✓	✓	✓ (atorvastatin)	✓	✓
Reduce TC, LDL-C, and apo B levels in children with heterozygous familial hypercholesterolemia (HeFH) if after an adequate trial of diet therapy the following findings are present: LDL-C remains ≥189 (lovastatin only) or 190 mg/dL OR LDL-C remains ≥160 mg/dL and there is a positive family history of premature cardiovascular disease (CVD) or two or more other cardiovascular risk factors are present in the pediatric patient	✓ ¶	✓ #	✓ ** (IR)		✓ ††	✓ ††	✓ ***	✓ (atorvastatin)		
Reduce elevated TG and very high LDL-C in patients with primary dysbetalipoproteinemia							✓			
Reduce TC and LDL-C in patients with homozygous familial hypercholesterolemia (HoFH) as an adjunct to other lipid-lowering treatments or if such treatments are unavailable	✓						✓	✓ (atorvastatin)	✓	✓
Reduce TC, LDL-C, and apo B in adults with HoFH						✓				
Reduce LDL-C, TC, non HDL-C and apo B in children and adolescents with HoFH, as monotherapy or with other lipid-lowering therapies						✓ Ⓐ				
Reduction of elevated TC and LDL-C levels in patients with primary hypercholesterolemia			✓ § (IR)							

Treatment of patients with primary dysbetalipoproteinemia who do not respond adequately to diet	✓				✓	✓		✓ (atorvastatin)		
Prevention of CVD										
Adjunctive therapy to diet to slow the progression of atherosclerosis in adult patients as part of a treatment strategy to lower TC and LDL-C to target levels						✓				
Reduce the risk of myocardial infarction (MI) and stroke in patients with type 2 diabetes, and without clinically evident coronary heart disease (CHD), but with multiple risk factors for CHD such as retinopathy, albuminuria, smoking, or HTN	✓							✓ (atorvastatin)		
Reduce the risk of MI, stroke, revascularization procedures, and angina in adult patients without clinically evident CHD, but with multiple risk factors for CHD such as age, smoking, HTN, low HDL-C, or a family history of early CHD	✓							✓ (atorvastatin)		
Reduce the risk of MI, undergoing myocardial revascularization procedures, and cardiovascular mortality with no increase in death from noncardiovascular causes in patients with hypercholesterolemia without clinically evident CHD					✓					
Reduce the risk of MI, unstable angina, and coronary revascularization procedures in patients without symptomatic CVD			✓ γ							
Reduce the risk of non-fatal MI, fatal and non-fatal stroke, revascularization procedures, hospitalization for congestive heart failure, and angina in patients with clinically evident CHD	✓							✓ (atorvastatin)		
Reduce the risk of stroke, MI, and arterial revascularization procedures in patients without clinically evident CHD but with an increased risk of CVD based on age ≥50 years old in men and ≥60 years old in women, high sensitivity C-reactive protein ≥2 mg/L, and the presence of at least one additional CVD risk factor such as HTN, low HDL-C, smoking, or a family history of premature CHD						✓				
Reduce the risk of total mortality by reducing coronary death, MI, undergoing myocardial revascularization procedures, stroke and stroke/transient ischemic attack, and to slow the progression of coronary atherosclerosis in patients with clinically evident CHD					✓					
Reduce the risk of total mortality by reducing CHD deaths, non-fatal MI and stroke, and need for coronary and non-coronary							✓			

revascularization procedures in patients at high risk of coronary events because of existing CHD, diabetes, peripheral vascular disease, history of stroke or other cerebrovascular disease										
Reduce the risk of undergoing coronary revascularization procedures and slow the progression of coronary atherosclerosis in patients with clinically evident CHD		✓								
Slow the progression of coronary atherosclerosis in patients with CHD as part of a treatment strategy to lower TC and LDL-C to target levels			✓							
Other										
Reduce the risk of hospitalization for angina and to reduce the risk of a coronary revascularization procedure in patients with recently documented CAD by angiography and without heart failure or an ejection fraction <40%								✓ (amlodipine)		
Symptomatic treatment of chronic stable angina								✓ (amlodipine)		
Treatment of confirmed or suspected vasospastic angina								✓ (amlodipine)		
Treatment of HTN, to lower blood pressure								✓ (amlodipine)		

Abbvr: CAD=coronary artery disease, CHD=coronary heart disease, ER=extended-release, IR=immediate-release, HTN=hypertension, MI=myocardial infarction.

§When the response to diet restricted in saturated fat and cholesterol and to other nonpharmacological measures alone has been inadequate.

¶In boys and postmenarchal girls 10 to 17 years of age.

#In adolescent boys and adolescents girls who are at least one year post-menarche, 10 to 16 years of age.

**In adolescent boys and girls who are at least one year post-menarche, 10 to 17 years of age.

††In children and adolescent patients eight to 17 years of age

‡In children and adolescents ages seven to 17 years of age

γFor ER lovastatin, for patients at high risk; for IR lovastatin, for patients with average to moderately elevated TC and LDL-C and below average HDL-C

(Prescribing information: ALTOPREV[®], 2018; CADUET[®], 2017; CRESTOR[®], 2017; FLOLIPID, 2017; Fluvastatin, 2017; LESCOL XL[®], 2017; LIPITOR[®], 2017; LIVALO[®], 2016 Lovastatin 2017; PRAVACHOL[®], 2017; VYTORIN[®], 2018; ZOCOR[®], 2018; ZYPITAMAG, 2018)
Clinical Pharmacology, 2018

Information on indications, mechanism of action, pharmacokinetics, and safety has been obtained from the prescribing information for the individual products, except where noted otherwise.

CLINICAL EFFICACY SUMMARY

- Numerous clinical trials have demonstrated that the statins (single-entity and combination products) can effectively lower LDL-C, non-HDL-C, total cholesterol (TC), and TG, as well as positively impact other lipid/lipoprotein parameters. Additionally, many studies have compared active treatment to placebo or compared combination therapy to monotherapy. In these studies, the more aggressive treatment regimens often improved lipid parameters to a greater extent than the less-intensive treatment regimens (Ai et al, 2008; Alvarez-Sala et al, 2008; Arca et al, 2007; Avis et al, 2007; Avis et al, 2010; Ballantyne et al, 2003; Ballantyne et al, 2004; Ballantyne et al, 2005; Ballantyne et al, 2006; Ballantyne et al, 2007; Ballantyne et al, 2008; Bardini et al, 2010; Bays et al, 2004; Bays et al, 2010; Bays et al, 2013; Bays et al, 2008a; Bays et al, 2008b; Becker et al, 2008; Betteridge et al, 2007a; Betteridge et al, 2007b; Braamskamp et al, 2015; Brown et al, 1990; Bullano et al, 2006; Bullano et al, 2007; Calza et al, 2008; Catapano et al, 2006; Charland et al, 2010; Chenot et al, 2007; Clearfield et al, 2006; Coll et al, 2006; Conard et al, 2008; Constance et al, 2007; Davidson et al, 2002; Deedwania et al, 2007a; Derosa et al, 2009; Erdine et al, 2009; Eriksson et al, 1998; Eriksson et al, 2011; Faergeman et al, 2008; Farnier et al, 2007; Farnier et al, 2008; Farnier et al, 2009; Feldman et al, 2004; Feldman et al, 2006; Ferdinand et al, 2006; Ferdinand et al, 2012; Flack et al, 2008; Florentin et al, 2011; Foody et al, 2010; Fox et al, 2007a; Fox et al, 2007b; Gagné et al, 2002; Gaudiani et al, 2005; Goldberg et al, 2004; Goldberg et al, 2006; Goldberg et al, 2009; Grimm et al, 2010; Gumprecht et al, 2011; Hall et al, 2009; Harley et al, 2007; Hing Ling et al, 2012; Hobbs et al, 2009; Hogue et al, 2008; Hunninghake et al, 2001; Illingworth et al, 1994; Insull et al, 2007; Jones et al, 2003; Jones et al, 2009a; Jones et al, 2009b; Kerzner et al, 2003; Kipnes et al, 2010; Knapp et al, 2001; Koshiyama et al, 2008; Kumar et al, 2009; Lee et al, 2007; Leiter et al, 2007; Leiter et al, 2008; Lewis et al, 2007; Lloret et al, 2006; Marais et al, 2008; May et al, 2008; Mazza et al, 2008; Melani et al, 2003; Meredith et al, 2007; Messerli et al, 2006; Milionis et al, 2006; Mohiuddin et al, 2009; Motomura et al, 2009; Neutel et al, 2009; Nicholls et al, 2010; Ose et al, 2007; Ose et al, 2009; Ose et al, 2010; Park et al, 2005; Park et al, 2010; Pearson et al, 2007; Piorkowski et al, 2007; Polis et al, 2009; Preston et al, 2007; Reckless et al, 2008; Robinson et al, 2009; Rodenburg et al, 2007; Roeters van Lennep et al, 2008; Rogers et al, 2007; Rosenson et al, 2009; Rotella et al, 2010; Roth et al, 2010; Saito et al, 2002; Sansanayudh et al, 2010; Sasaki et al, 2008; Shafiq et al, 2007; Stalenhoef et al, 2005; Stein et al, 2003; Stein et al, 2004; Stein et al, 2007; Stein et al, 2008; Viigimaa et al, 2010; Vuorio et al, 2014; Winkler et al, 2007; Winkler et al, 2009; Wlodarczyk et al, 2008; Wolffenbuttel et al, 2005; Yoshitomi et al, 2006; Zieve et al, 2010).
- All of the statins, with the exception of pitavastatin, have been shown to have beneficial effects on CHD outcomes, and the majority of them (atorvastatin, pravastatin, rosuvastatin, and simvastatin) have also been shown to decrease the risk of stroke (Afilalo et al, 2007; Afilalo et al, 2008; Ahmed et al, 2006; Amarenco et al, 2009a; Amarenco et al, 2009b; Asselbergs et al, 2004; Athyros et al, 2002; Athyros et al, 2007; Baigent et al, 2005; Barter et al, 2007; Briel et al, 2006; Bushnell et al, 2006; Byington et al, 1995; Cannon et al, 2004; Cannon et al, 2006; Cannon et al, 2015; Chan et al, 2010; Cholesterol Treatment Trialists' (CTT) Collaborators, 2008; Chonchol et al, 2007; Colhoun et al, 2004; Collins et al, 2003; Crouse et al, 2007; de Lemos et al, 2004; Deedwania et al, 2006; Deedwania et al, 2007b; Downs et al, 1998; Everett et al, 2010; Ford et al, 2007; Furberg et al, 1994; Hitman et al, 2007; Hulten et al, 2006; Khush et al, 2007; Knopp et al, 2006; Koenig et al, 2001; Koga et al, 2018; LaRosa et al, 2005; LaRosa et al, 2007; Liem et al, 2002; Meaney et al, 2009; Mood et al, 2007; Mora et al, 2010; Murphy et al, 2007; Nakamura et al, 2006; Neil et al, 2006; Nicholls et al, 2006; Nissen et al, 2004; Nissen et al, 2005; Nissen et al, 2006; No authors listed, 1994; No authors listed, 2002; No authors listed, 2007; Olsson et al, 2007; O'Regan et al, 2008; Pedersen et al, 2005; Pitt et al, 1999; Pitt et al, 2012; Ray et al, 2005; Ray et al, 2006; Ridker et al, 2008; Ridker et al, 2009; Ridker et al, 2010; Rossebø et al, 2008; Sacks et al, 1996; Sakamoto et al, 2007; Sato et al, 2008; Schmermund et al, 2006; Schoenhagen et al, 2006; Schouten et al, 2009; Schwartz et al, 2005; Scirica et al, 2006; Serruys et al, 2002; Sever et al, 2003; Sever et al, 2005; Shah et al, 2008; Shepherd et al, 1995; Shepherd et al, 2007; Shepherd et al, 2006; Shepherd J et al, 2002; Strandberg et al, 2009; Tavazzi L et al, 2008; Taylor et al, 2013; The ALLHAT Officers and Coordinators for the ALLHAT Collaborative Research Group, 2002; The Long-term Intervention with Pravastatin in Ischemic Disease (LIPID) Study Group, 1998; The Pravastatin Multinational Study Group for Cardiac Risk Patients (PMS-CRP), 1993; Thompson et al, 2004; Tikkanen et al, 2009; Waters et al, 2006; Wenger et al, 2007; Yu et al, 2007).
- Two early primary prevention trials (West of Scotland Coronary Prevention Study [WOSCOPS] and Air Force/Texas Coronary Atherosclerosis Prevention Study [AFCAPS/TexCAPS]) demonstrated that the use of statins significantly reduced the risk for major coronary events (Downs et al, 1998; Shepard et al, 1995).
- Specifically, the WOSCOPS trial (N=6959) demonstrated that compared to placebo, pravastatin (40 mg/day) was associated with a significant 31% reduction in the risk of the combined endpoint of CHD death and nonfatal MI (P<0.001). A reduction in the secondary endpoint of cardiovascular death was also significant in favor of pravastatin

(32%; $P=0.033$) (Shepard et al, 1995). Results of a 20-year observational follow-up of this trial continued to show beneficial effects of pravastatin on reduction of CHD. Among those with and without LDL-C ≥ 190 mg/dL ($N=5529$), pravastatin reduced the risk of CHD by 27% ($P=0.002$) and MACE by 25% ($P=0.004$). Among individuals with LDL-C ≥ 190 mg/dL ($N=2560$), pravastatin reduced the risk of CHD-related death, cardiovascular death, and all-cause mortality by 28% ($P=0.020$), 25% ($P=0.009$), and 18% ($P=0.004$), respectively (Vallejo-Vaz et al, 2017).

- The AFCAPS/TexCAPs trial ($N=6,605$) demonstrated similar benefits but with lovastatin (20 to 40 mg/day). In this trial, lovastatin was associated with a significant 37% reduction in the risk of the combined endpoint of fatal or nonfatal MI, unstable angina or sudden cardiac death ($P<0.001$). The AFCAPS/TexCAPs trial contained too few events to perform survival analysis on cardiovascular and CHD mortality (Downs et al, 1998).
- The Anglo-Scandinavian Cardiac Outcomes Trial (ASCOT, $N=10,305$) was terminated early (median duration, 3.3 years) due to the significant benefits observed with atorvastatin. In this trial, patients had average cholesterol concentrations but were at an increased risk for CHD due to the presence of HTN and three additional CHD risk factors. Compared to placebo, atorvastatin significantly reduced the risk of the combined endpoint of CHD death and nonfatal MI by 35% ($P=0.0005$) (Sever et al, 2003).
- Despite not demonstrating any benefit on all-cause mortality within the ASCOT trial ($P=0.1649$), atorvastatin has been associated with significant reductions in all-cause mortality in other primary prevention trials (Colhoun et al, 2004; Sever et al, 2003; Sever et al, 2005).
- A benefit in all-cause mortality, as well as other cardiovascular outcomes, with rosuvastatin in primary prevention was demonstrated in the Justification for the Use of Statins in Prevention: an Intervention Trial Evaluating Rosuvastatin (JUPITER) trial ($N=17,802$). This trial sought to evaluate the efficacy of rosuvastatin in reducing cardiac events in patients with elevated high sensitivity C-reactive protein levels, which they note as being a predictor for cardiac events. This trial was terminated early (median duration 1.9 years) due to the significant benefits observed with rosuvastatin. Compared to placebo, rosuvastatin significantly reduced the risk of a first major cardiovascular event (nonfatal MI, nonfatal stroke, hospitalization for unstable angina, revascularization procedure or cardiovascular death) by 44% ($P<0.0001$). When analyzed individually, rosuvastatin was associated with a significant benefit for all primary outcomes, as well as all-cause mortality ($P=0.02$) (Ridker et al, 2008).
- Meta-analyses support the findings observed in the individual primary prevention trials (Adams et al, 2018; Baigent et al, 2005; CTT Collaborators et al, 2008; Mora et al, 2010; O'Regan et al, 2008; Taylor et al, 2011; Nunes et al, 2017).
- The Incremental Decrease in Endpoints Through Aggressive Lipid Lowering (IDEAL) trial ($N=8,888$) compared intensive lipid lowering therapy with atorvastatin 80 mg/day to moderate therapy with simvastatin 20 mg/day (with the potential to increase to 40 mg/day based on improvements in lipid profile). In this trial, atorvastatin did not significantly reduce the risk of the primary composite endpoint of CHD death, nonfatal MI, or cardiac arrest with resuscitation (hazard ratio [HR], 0.89; 95% confidence interval [CI], 0.78 to 1.01; $P=0.07$). Atorvastatin was associated with a significant reduction in the risk of major cardiovascular events compared to simvastatin (12.0 vs 13.7%; HR, 0.87; $P=0.02$). Atorvastatin was associated with a significant reduction in the risk of any CHD event compared to simvastatin (20.2 vs 23.8%; HR, 0.84; $P<0.001$) and for the risk of any cardiovascular events compared to simvastatin (26.5 vs 30.8%; HR, 0.84; $P<0.001$). For the individual events, atorvastatin had a lower rate of nonfatal acute MI than simvastatin (7.2% vs. 6.0%; HR, 0.83; 95% CI, 0.71 to 0.98; $P=0.02$), but the treatments were no different in terms of all-cause ($P=0.81$) or noncardiovascular ($P=0.47$) mortality. In addition, intensive therapy with atorvastatin 80 mg/day was associated with a significantly higher incidence of discontinuations due to adverse events ($P<0.001$) (Pedersen et al, 2005). A total of 94 patients (2.2%) receiving atorvastatin and 135 patients (3.2%) receiving simvastatin developed peripheral arterial disease (HR, 0.7; 95% CI, 0.53 to 0.91; $P=0.007$) (Stoekenbroek et al, 2015).
- Several trials have demonstrated that statins are effective in delaying the progression of atherosclerotic disease in patients with CHD. Included in these is the head-to-head REVERSAL trial that demonstrated that intensive lipid lowering with atorvastatin 80 mg/day was associated with a significantly lower median percentage change in atheroma volume compared to moderate lipid lowering with pravastatin 40 mg/day after 18 months ($P=0.02$) (Byington et al, 1995; Chan et al, 2010; Crouse et al, 2007; Furberg et al, 1994; Karlson et al, 2018; Nicholls et al, 2006; Nissen et al, 2004; Nissen et al, 2005; Nissen et al, 2006; Schmermund et al, 2006; Schoenhagen et al, 2006). A meta-analysis comparing the efficacy and safety of atorvastatin and pitavastatin on the regression of atherosclerosis did not find a statistically significant difference between these agents when evaluating changes in plaque volume, lumen volume, and external elastic membrane. However, atorvastatin was potentially more effective than pitavastatin at reducing LDL-C and improving HDL-C (Liu et al, 2018).

- The majority of secondary prevention trials have evaluated the use of statins initiated three to six months after an acute cardiac event; however, evidence supports the use of these agents initiated right after an acute event (Briel et al, 2006; Cannon et al, 2004; de Lemos et al, 2004; Liem et al, 2002).
- The Myocardial Ischemia Reduction with Aggressive Cholesterol Lowering (MIRACL) trial (N=3,086), a placebo-controlled trial with atorvastatin, is noteworthy as it demonstrated that when initiated in the hospital following an acute coronary syndrome (ACS), atorvastatin was safe and associated with a 16% reduction in the composite of death, nonfatal acute MI, resuscitated cardiac arrest, or recurrent symptomatic myocardial ischemia after 16 weeks (P=0.048) (Schwartz et al, 2005). However, a 2018 RCT that included 4191 patients with ACS and planned PCI found that 2 loading doses of atorvastatin 80 mg before and 24-hours after surgery did not reduce the rate of MACE at 30 days when compared to placebo (absolute difference, 0.85%; 95% CI, -0.70% to 2.41%; hazard ratio, 0.88; 95% CI, 0.69-1.11; P=0.27) (Berwanger et al, 2018).
- The Improved Reduction of Outcomes: Vytorin Efficacy International Trial (IMPROVE-IT) investigated the efficacy of the addition of ezetimibe to simvastatin for the prevention of stroke and other adverse cardiovascular events in 18,144 patients. After 7 years, the combination of ezetimibe and simvastatin significantly reduced the risk of stroke of any etiology (HR, 0.83; 95% CI, 0.70-0.98; P=0.029) and ischemic stroke (HR, 0.76; 95% CI, 0.63-0.91; P=0.003) when compared to simvastatin monotherapy. Significant benefits were also observed in the subgroup of patients with prior stroke (Bohula et al, 2017).
- Of the head-to-head trials, the Pravastatin or Atorvastatin Evaluation and Infection Therapy–Thrombolysis in Myocardial Infarction 22 (PROVE IT–TIMI 22) trial (N=4,162) again compared intensive lipid therapy with atorvastatin 80 mg/day to standard therapy with pravastatin 40 mg/day (with a potential to increase to 80 mg/day based on improvements in lipid profile). Patients who were hospitalized with an ACS within the preceding 10 days were enrolled. After two years, atorvastatin significantly reduced the combined endpoint of all-cause mortality, MI, unstable angina requiring hospitalization, coronary revascularization performed >30 days after randomization, and stroke by 16% compared to pravastatin (P=0.005). Among the individual endpoints, atorvastatin was significant for reducing the risk of revascularization (P=0.04) and unstable angina (P=0.02). In this trial, discontinuations due to adverse events were similar between the two treatments (P=0.11) (Cannon et al, 2004).
- A meta-analysis which assessed the efficacy of high dose atorvastatin in patients who underwent percutaneous coronary intervention (PCI) (N=2,850) found that atorvastatin significantly reduced the risk of MI in patients with PCI compared to placebo (RR, 0.62; 95% CI, 0.49 to 0.78) (Lu et al, 2017).
- A meta-analysis evaluated the efficacy and safety of dosing statins on alternative days (N=505) compared to daily dosing (N=518). Although there was no differences on TG, the reduction in TC (P<0.00001) and LDL-C (P=0.003) was significantly greater in the daily dosing group (Awad et al, 2017).
- A Cochrane review assessed the effectiveness of statins in children aged 4 to 18 years with HeFH and found that statin treatment is effective. Statin therapy was found to be safe with no significant safety issues in the short-term (Vuorio et al, 2017).

SAFETY SUMMARY

- Statins are contraindicated in documented hypersensitivity to the agent, unexplained elevations in serum transaminases, active liver disease, and patients who are pregnant or nursing.
- The statins are generally well-tolerated, and the most common side effects are gastrointestinal disturbances, headache, insomnia, myalgia, and rash. Muscle aches and weakness are reported by 1 to 2% of patients taking statins. The symptoms are usually mild and generally do not lead to discontinuation, however, myopathy can sometimes take the form of rhabdomyolysis, with or without acute renal failure secondary to myoglobinuria. Rare fatalities have occurred. The risk of myopathy is increased by high levels of HMG-CoA reductase inhibitory activity in plasma. All statins can increase hepatic transaminase levels and creatinine kinase.
- Increases in hemoglobin A1c (HbA1c) and fasting serum glucose have been reported with statins. New-onset diabetes is increased in patients treated with statins; however, it is dose-related, occurs primarily in patients on metformin and a sulfonylurea, appears to be less common with pravastatin and possibly pitavastatin, and occurs overall to a lesser extent than the associated decrease in atherosclerotic cardiovascular disease (ASCVD) (Jellinger et al, 2017).
- Pravastatin is the only statin that does not undergo cytochrome (CYP) 450 metabolism, and is therefore associated with a lower risk for drug interactions. Atorvastatin (to a lesser extent), lovastatin, and simvastatin are primarily metabolized by the CYP3A4 isoenzyme, while fluvastatin, pitavastatin, and rosuvastatin are metabolized by the CYP2C9 isoenzyme, which may result in differences in their drug interaction profiles (Wiggins et al, 2016).

- The 2016 scientific statement written by the American Heart Association (AHA) stated that the risk for interactions between statins and other cardiovascular drugs may be unavoidable for heart patients, but it can be reduced with proper clinical management. A review of all of the medications that statin-treated patients are taking should be done at each patient visit, so that potential drug interactions can be identified early. Some key recommendations include:
 - Concomitant use of lovastatin, pravastatin, or simvastatin with gemfibrozil should be avoided. When gemfibrozil is used with other statins, a lower statin dose should be utilized.
 - A non-CYP3A4-metabolized statin should be used in combination with verapamil and diltiazem (calcium channel blockers). The dose of lovastatin or simvastatin should be limited to 20 mg daily or less when given with the calcium channel blocker amlodipine.
 - The concomitant use of cyclosporine, everolimus, sirolimus, or tacrolimus should be avoided with lovastatin, simvastatin, and pitavastatin, as the combination could be potentially harmful.
 - Numerous other drug interactions are listed, many of which require dose adjustment of statin therapy or drug level monitoring (e.g. digoxin) (Wiggins et al, 2016).

DOSING AND ADMINISTRATION

Table 3. Dosing and Administration

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
Single-Entity Agents				
atorvastatin	Tablet: 10 mg 20 mg 40 mg 80 mg	<p><u>Hyperlipidemia:</u> Tablet: initial, 10 to 40 mg once daily; maintenance, 10 to 80 mg/day</p> <p><u>Adjunct to diet for the treatment of patients with elevated serum TG levels, reduce TC and LDL-C in patients with HoFH as an adjunct to other lipid lowering treatments or if such treatments are unavailable, treatment of patients with primary dysbetalipoproteinemia:</u> Tablet: 10 to 80 mg/day</p> <p><u>HeFH in pediatric patients 10 to 17 years old:</u> Tablet: initial dose 10 mg/day, maximum dose 20 mg/day</p>	After initiation and/or upon titration, lipid levels should be analyzed within two to four weeks and dosage adjusted accordingly.	<p>May be administered with or without food.</p> <p>Tablets may be taken at any time during the day.</p>
fluvastatin	Capsule: 20 mg 40 mg Extended-release tablet: 80 mg	<p><u>Hypercholesterolemia (including HeFH and nonfamilial) and mixed dyslipidemia in adults:</u> Capsule: 40 mg once daily or 40 mg twice daily</p> <p>Patients requiring LDL-C reductions ≥25% should initiate fluvastatin therapy at 40 mg once daily or 80 mg in divided doses of the 40 mg capsule given twice daily.</p> <p>Patients requiring LDL-C reductions < 25% should initiate a starting dose of 20 mg.</p>	<p>After initiation and/or upon titration, lipid levels should be analyzed after four weeks and dosage adjusted accordingly.</p> <p>Max dose is 20 mg twice daily when used with cyclosporine or fluconazole.</p>	<p>Capsules should be taken in the evening if dosed once daily. If 80 mg/day is used, it should be administered in two divided doses (immediate-release capsule).</p> <p>May be administered with or without food.</p> <p>Tablets may be taken at any time</p>

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
		Extended-release tablet: 80 mg once daily <u>HeFH in pediatric patients:</u> Capsule: 20 mg daily, maximum dose 40 mg twice daily Extended-release tablet: 80 mg once daily		during the day (extended-release tablet). Tablets should be swallowed whole. (extended-release tablet).
lovastatin	Extended-release tablet: 20 mg 40 mg 60 mg Tablet: 10 mg 20 mg 40 mg	<u>Hyperlipidemia:</u> Extended-release tablet: initial, 20 to 60 mg once daily; maintenance, 20 to 60 mg/day Tablet: initial, 20 mg once daily; maintenance, 10 to 80 mg/day in single or two divided doses; maximum, 80 mg/day <u>Prevention of CVD:</u> Extended-release tablet: initial, 20 to 60 mg once daily; maintenance, 20 to 60 mg/day Tablet: initial, 20 mg once daily; maintenance, 10 to 80 mg/day in single or two divided doses; maximum, 80 mg/day	Prior to initiation and periodically during therapy, lipid levels should be analyzed and dosage adjusted accordingly.	Extended-release tablet should be taken at bedtime. Extended-release tablets should be swallowed whole. Immediate-release tablet should be taken with an evening meal.
pitavastatin	Tablet: 1 mg 2 mg 4 mg	<u>Hyperlipidemia:</u> Tablet: initial, 2 mg once daily; maintenance, 1 to 4 mg/day; maximum, 4 mg/day	After initiation and/or upon titration, lipid levels should be analyzed after four weeks and dosage adjusted accordingly. Do not exceed 4 mg once daily dosing due to increased risk of severe myopathy Max dose is 1 mg mg/day when used with erythromycin.	May be administered with or without food. Tablets may be taken at any time during the day.

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
			<p>Max dose is 2 mg mg/day when used with rifampin.</p> <p>Use caution in patients receiving ≥ 1 gram daily of niacin-containing products.</p>	
pravastatin	Tablet: 10 mg* 20 mg 40 mg 80 mg	<p><u>Hyperlipidemia:</u> Tablet: initial, 40 mg once daily; maintenance, 40 to 80 mg once daily</p> <p><u>Prevention of CVD:</u> Tablet: initial, 40 mg once daily; maintenance, 40 to 80 mg once daily</p> <p><u>Pediatric patients:</u> Ages eight to 13 years old: 20 mg once daily Ages 14 to 18 years old: 40 mg once daily</p>	<p>After initiation and/or upon titration, lipid levels should be analyzed after four weeks and dosage adjusted accordingly.</p> <p>Max dose in patients taking cyclosporine is 20 mg/day. Max dose in patients taking clarithromycin is 40 mg/day.</p>	<p>May be administered with or without food.</p> <p>Tablets may be taken at any time during the day.</p>
rosuvastatin	Tablet: 5 mg 10 mg 20 mg 40 mg	<p><u>Hyperlipidemia:</u> Tablet: initial, 10 to 20 mg once daily; maintenance, 5 to 40 mg/day</p> <p><u>Reduce TC, LDL-C and apo B in patients with HoFH:</u> Tablet: initial, 20 mg once daily;</p> <p>Ages seven to 17 years: Tablet: 20 mg once daily</p> <p><u>Reduce TC, LDL-C and apo B in pediatric patients with HeFH:</u> Aged eight to less than 10 years: Tablet: maintenance, 5 to 10 mg/day</p> <p>Aged 10 to 17 years: Tablet: maintenance, 5 to 20 mg/day</p>	<p>After initiation and/or upon titration, lipid levels should be analyzed within two to four weeks and dosage adjusted accordingly.</p> <p>Dosing in Asian patients: initial, 5 mg once daily</p> <p>Max dose is 5 mg once daily when used with cyclosporine and 10 mg once daily when used with gemfibrozil, atazanavir/</p>	<p>May be administered with or without food.</p> <p>Tablets may be taken at any time during the day.</p>

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
simvastatin	<p>Tablet: 5 mg 10 mg 20 mg 40 mg 80 mg</p> <p>Oral suspension: 20 mg/5 mL 40 mg/5 mL</p>	<p><u>Hyperlipidemia:</u> Tablet: initial, 10 or 20 mg once daily; maintenance, 5 to 40 mg/day</p> <p><u>Reduce TC and LDL-C in patients with HoFH as an adjunct to other lipid lowering treatments or if such treatments are unavailable:</u> Tablet: 40 mg once daily</p> <p><u>Prevention of CVD:</u> Tablet: initial, 10 or 20 mg once daily; maintenance, 5 to 40 mg/day</p> <p><u>Reduce TC, LDL-C and apo B in pediatric patients with HeFH:</u> Aged 10 to 17 years: Tablet: initial, 10 mg/day; maintenance, 10 to 40 mg/day; maximum dose is 40 mg/day</p>	<p>ritonavir, lopinavir/ritonavir, or simeprevir.</p> <p>After initiation and/or upon titration, lipid levels should be analyzed after four weeks and dosage adjusted accordingly.</p> <p>Dose should be decreased by 50% if initiating lomitapide.</p> <p>Simvastatin dosage should not exceed 20 mg/day (or 40 mg/day for patients who have previously taken simvastatin 80 mg/day chronically (e.g. for 12 months or more) without evidence of muscle toxicity) while taking lomitapide.</p> <p>Use caution in Chinese patients receiving doses >20 mg with niacin-containing products.</p> <p>Max dose is 10 mg/day when used with verapamil, diltiazem, or dronedarone.</p> <p>Max dose is 20 mg/day when</p>	<p>Tablets should be taken in the evening. The oral suspension should be taken on an empty stomach.</p> <p>Shake oral suspension bottle for at least 20 seconds. Use accurate measuring device.</p> <p>Due to the increased risk of myopathy, including rhabdomyolysis, particularly during the first year of treatment, use of the 80 mg dose should be restricted to patients who have been taking the 80 mg dose chronically without evidence of muscle toxicity.</p>

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
			<p>used with amiodarone, amlodipine, or ranolazine.</p> <p>Simvastatin is contraindicated for use with strong CYP3A4 inhibitors.</p> <p>For patients at high risk for a CHD event due to existing CHD, diabetes, peripheral vessel disease, history of stroke or other cerebrovascular disease, the recommended starting dose is 40 mg/day.</p> <p>Use caution in patients receiving ≥ 1 gram daily of niacin-containing products.</p>	

Combination Products

amlodipine/atorvastatin	<p>Tablet: 2.5/10 mg 2.5/20 mg 2.5/40 mg 5/10 mg 5/20 mg 5/40 mg 5/80 mg 10/10 mg 10/20 mg 10/40 mg 10/80 mg</p>	<p>Dosage of amlodipine/atorvastatin must be individualized on the basis of both effectiveness and tolerance for each individual component in the treatment of hypertension/angina and hyperlipidemia.</p> <p>Select doses of amlodipine and atorvastatin independently.</p> <p>The usual starting dose for amlodipine is 5 mg daily and for atorvastatin 10 to 20 mg daily. The maximum dose is amlodipine 10 mg daily and atorvastatin 80 mg daily.</p> <p>Patients requiring large LDL-C reductions (>45%) should initiate</p>	<p>After initiation and/or upon titration, lipid levels should be analyzed within two to four weeks and dosage adjusted accordingly.</p> <p>Dosage should be adjusted to achieve blood pressure goals. In general, wait seven to 14 days between titration steps.</p>	<p>May be administered with or without food.</p> <p>Tablets may be taken at any time during the day.</p>
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Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
		<p>atorvastatin therapy at 40 mg once daily.</p> <p><u>HeFH in pediatric patients 10 to 17 years old:</u> <i>Atorvastatin</i> Tablet: initial dose 10 mg/day, maximum dose 20 mg/day <i>Amlodipine [age 6 to 17 years old]</i> Tablet: initial dose 2.5 to 5 mg maximum dose 5 mg</p>	<p>Titration may proceed more rapidly if clinically warranted, provided the patient is assessed frequently.</p>	
ezetimibe/atorvastatin	Tablet: 10/10 mg 10/20 mg 10/40 mg 10/80 mg	<p><u>Usual starting dose:</u> 10/10 mg or 10/20 mg once daily. Usual dose range is 10/10 mg to 10/80 mg once daily.</p> <p>May initiate at 10/40 mg once daily for patients requiring a larger LDL-C reduction (> 55%).</p> <p><u>HoFH:</u> 10/40 mg or 10/80 mg once daily.</p>	<p>After initiation or titration of doses, lipid levels may be analyzed after two or more weeks.</p> <p>For patients taking clarithromycin, itraconazole, saquinavir + ritonavir, darunavir + ritonavir, or fosamprenair alone or with ritonavir: Do not exceed 10/20 mg once daily.</p> <p>For patients taking nelfinavir: Do not exceed 10/40 mg once daily.</p>	<p>Tablets may be taken at any time of the day.</p> <p>May be administered with or without food.</p>
ezetimibe/simvastatin	Tablet: 10/10 mg 10/20 mg 10/40 mg 10/80 mg	<p><u>Hyperlipidemia:</u> <u>Adjunct to diet to reduce elevated TC, LDL-C, apo B and TG levels and to increase HDL-C in patients with primary hypercholesterolemia and mixed dyslipidemia, reduce TC and LDL-C in patients with HoFH as an adjunct to other lipid lowering treatments or if such treatments are unavailable:</u> Tablet: initial, 10/10 or 10/20 mg once daily; maintenance, 10/10 to 10/40 mg/day</p>	<p>After initiation and/or upon titration, lipid levels should be analyzed within two or more weeks and dosage adjusted accordingly.</p> <p>Decrease dose of VYTORIN by 50% if initiating lomitapide.</p>	<p>May be administered with or without food.</p> <p>Tablets should be taken in the evening.</p> <p>Due to the increased risk of myopathy, particularly during the first year of treatment, use of</p>

Drug	Dosage Form: Strength	Usual Recommended Dose	Other Dosing Considerations	Administration Considerations
			<p>VYTORIN dosage should not exceed 10/20 mg once day (or 10/40 mg once daily for patients who have previously taken simvastatin 80 mg once day chronically, e.g., for 12 months or more, without evidence of muscle toxicity) while taking lomitapide.</p> <p>Max dose is 10/10 mg/day when used with verapamil, diltiazem, or dronedarone.</p> <p>Max dose is 10/20 mg/day when used with amiodarone, amlodipine, or ranolazine.</p> <p>VYTORIN is contraindicated for use with strong CYP3A4 inhibitors.</p> <p>Use caution in patients receiving ≥ 1 gram daily of niacin-containing products.</p>	<p>the 10/80 mg dose should be restricted to patients who have been taking the 10/80 mg dose chronically.</p>

*Pravachol 10 mg is no longer available, however, generic pravastatin 10 mg remains available.

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SPECIAL POPULATIONS

Table 4. Special Populations

Drug	Population and Precaution				
	Elderly	Pediatrics	Renal Dysfunction	Hepatic Dysfunction	Pregnancy* and Nursing
atorvastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Approved for use in children 10 to 17 years of age for the treatment of HeFH. Doses of >20 mg have not been studied in this population. Safety and efficacy in children <10 years of age have not been established.	No dosage adjustment required.	Contraindicated in active liver disease or in patients with unexplained persistent elevations or serum transaminases.	Unclassified† Contraindicated in pregnant women. Contraindicated during breastfeeding.
fluvastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Approved for use in children 9 to 16 years of age for the treatment of HeFH. Safety and efficacy in children for other approved indications have not been established.	No dosage adjustment required in mild to moderate renal dysfunction. Use with caution in severe renal dysfunction; doses above 40 mg per day have not been studied.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Pregnancy Category X Potential excretion into breast milk; contraindicated during breastfeeding
lovastatin	No dosage adjustment required in the elderly. The initial starting dose of lovastatin extended-release should not exceed 20 mg/day (ALTOPREV).	Approved for use in children 10 to 17 years of age for the treatment of HeFH (MEVACOR); maximum dose of 40 mg/day. Safety and efficacy in children <10 years of age have not been established (MEVACOR). Safety and efficacy in children have not been	Renal dosage adjustment is required; for creatinine clearances <30 mL/minute, use with caution and carefully consider doses >20 mg/day.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Pregnancy Category X (MEVACOR) No data on excretion in breast milk; not recommended (MEVACOR) Unclassified† (ALTOPREV) Contraindicated in pregnant women (ALTOPREV). Contraindicated during

Drug	Population and Precaution				
	Elderly	Pediatrics	Renal Dysfunction	Hepatic Dysfunction	Pregnancy* and Nursing
		established (ALTOPREV).			breastfeeding (ALTOPREV).
pitavastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Safety and efficacy in children have not been established.	Renal dosage adjustment is required; for creatinine clearances 15 to 59 mL/minute or end-stage renal disease receiving hemodialysis, an initial dose of 1 mg once daily and a maximum dose of 2 mg/day is recommended.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Unclassified† Contraindicated in pregnant women. Contraindicated during breastfeeding.
pravastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Approved for use in children eight to 18 years of age for the treatment of HeFH. Safety and efficacy in children <8 years of age have not been established.	Renal dosage adjustment is required in severe renal impairment; an initial dose of 10 mg/day is recommended.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Unclassified† Contraindicated in pregnant women. Pravastatin is present in breast milk; contraindicated during breastfeeding.
rosuvastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Approved for use in children 8 to 17 years of age for the treatment of HeFH and 7 to 17 years of age for the treatment of HoFH. Safety and efficacy in children <7 years of age have not been established.	No dosage adjustment required in mild to moderate renal dysfunction. Renal dosage adjustment required; for creatinine clearances <30 mL/minute, an initial dose of 5 mg/day and a maximum dose of 10 mg/day are recommended.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Unclassified† Contraindicated in pregnant women. Limited data indicate that the drug is in breast milk; contraindicated during breastfeeding.

Drug	Population and Precaution				
	Elderly	Pediatrics	Renal Dysfunction	Hepatic Dysfunction	Pregnancy* and Nursing
simvastatin	No evidence of overall differences in safety or efficacy observed between elderly and younger adult patients.	Approved for use in children 10 to 17 years of age for the treatment of HeFH. Doses greater than 40 mg have not been studied in this population. Safety and efficacy in children <10 years of age have not been established.	No dosage adjustment required in mild to moderate renal dysfunction. Renal dosage adjustment required for severe renal impairment: an initial dose of 5 mg/day with close monitoring is recommended.	Contraindicated in active liver disease or unexplained persistent elevations in serum transaminases.	Pregnancy Category X Unknown whether excreted in breast milk; contraindicated during breastfeeding.
Combination Products					
amlodipine/atorvastatin	Safety and efficacy in elderly patients have not been established. Elderly patients have decreased clearance of amlodipine; lower initial doses of amlodipine may be required.	Safety and efficacy in children have not been established. Safety and efficacy of atorvastatin in children <10 years and amlodipine in children <6 years of age have not been established	No dosage adjustment required.	Contraindicated in active liver disease.	Unclassified[†] Contraindicated for use during pregnancy and in women who may become pregnant. Contraindicated for use during breastfeeding.
ezetimibe/atorvastatin	The maximum dosage limit is 10/80 mg once daily for most patients.	Safety and efficacy have not been established.	No dosage adjustment is needed.	Contraindicated in patients with active hepatic disease or unexplained transaminase elevations.	Unclassified [†] Contraindicated for use during pregnancy and in women who may become pregnant. Contraindicated for use during breastfeeding.
ezetimibe/simvastatin	No evidence of overall differences in safety or efficacy observed between elderly	Safety and efficacy in children < 10 years old have not been established.	Use with caution doses exceeding 10/20 mg in patients with moderate to	Contraindicated in active liver disease or unexplained persistent elevations in	Unclassified[†] Contraindicated for use during pregnancy and in women who may

Drug	Population and Precaution				
	Elderly	Pediatrics	Renal Dysfunction	Hepatic Dysfunction	Pregnancy* and Nursing
	and younger adult patients; prescribe with caution.		severe renal dysfunction.	serum transaminases.	become pregnant. Contraindicated for use during breastfeeding.

* Pregnancy Category X = Contraindicated in pregnant women due to evidence of fetal abnormalities from adverse effects data from investigational or marketing experience. Risks of use of the drug in pregnant women clearly outweigh potential benefits.

† In accordance with the FDA's Pregnancy and Lactation Labeling Rule (PLLR), this product is not currently assigned a Pregnancy Category. Consult product prescribing information for details.

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CONCLUSION

- Statins are approved for the treatment of a variety of lipid disorders, including primary hypercholesterolemia, mixed dyslipidemia, and hypertriglyceridemia.
- The fixed-dose combination products (CADUET [amlodipine/atorvastatin], ezetimibe/atorvastatin, and VYTORIN [ezetimibe/simvastatin]) are indicated for use when dual therapy is appropriate.
- Statins decrease LDL-C according to the intensity of statin used and TG by 7% to 30%, as well as increase HDL-C by 5% to 15% when administered as monotherapy. The effects on LDL-C are dose-dependent and log-linear. Statins also decrease TG and increase HDL-C by varying levels.
- All products in this review are now available in a generic formulation except for ALTOPREV® (lovastatin extended-release), FLOLIPID (simvastatin oral suspension), LIVALO (pitavastatin), and Zypitamag (pitavastatin) (Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations, 2018).
- In general, therapeutic lifestyle changes, including diet, exercise and smoking cessation, remain an essential modality in the management of patients with hypercholesterolemia. When LDL-C lowering is required, initial treatment with a statin is recommended.
- In 2004, the National Cholesterol Education Program (NCEP) published guidelines on the Implications of Recent Clinical Trials for the NCEP Adult Treatment Panel III, which stated the following:
 - When LDL-C lowering drug therapy is employed in high-risk or moderately-high risk patients, it is advised that intensity of therapy be sufficient to achieve ≥30 to 40% reduction in LDL-C levels.
 - Standard statin doses are defined as those that lower LDL-C levels by 30 to 40%. The same effect may be achieved by combining lower doses of statins with other drugs or products such as bile acid sequestrants, ezetimibe, nicotinic acid, or plant stanols/sterols.
 - When LDL-C level is well above 130 mg/dL (e.g., ≥160 mg/dL), the statin dose may need to be increased or a second agent (e.g., a bile acid sequestrant, ezetimibe, nicotinic acid) may be required. Alternatively, maximizing dietary therapy (including use of plant stanols/sterols) combined with standard statin doses may be sufficient to attain goals.
 - Fibrates may have an adjunctive role in the treatment of patients with high TG and low HDL-C, especially in combination with statins.
 - In high risk patients with high TG or low HDL-C levels, consideration can be given to combination therapy with fibrates or nicotinic acid and a LDL lowering agent.
 - For the treatment of HeFH, LDL-C lowering drugs should be initiated in young adulthood. Statins are considered first-line therapy. Two-drug and sometimes three-drug therapy may be needed (Grundy et al, 2004).
- The 2013 American College of Cardiology/American Heart Association (ACC/AHA) Guidelines on Treatment of Blood Cholesterol to Reduce Atherosclerotic Cardiovascular Risk in Adults focus on primary and secondary atherosclerotic cardiovascular disease (ASCVD) risk reduction in adults (Stone et al, 2014).
 - These guidelines established four statin benefit groups: (1) individuals with clinical ASCVD (2) individuals with primary elevations of LDL-C >190 mg/dL (3) individuals with diabetes aged 40 to 75 years with LDL-C 70 to 189 mg/dL and without clinical ASCVD, and (4) individuals aged 40 to 75 years without clinical ASCVD or diabetes with LDL-C 70 to 189 mg/dL and estimated 10-year ASCVD risk >7.5%

- Intensity of statin therapy (high, moderate, and low) is the new goal of treatment in the benefit groups for use in primary and secondary prevention of ASCVD.
- A new cardiovascular risk tool, based on pooled cohort equations, has been created to estimate absolute 10-year ASCVD risk (defined as first occurrence nonfatal and fatal MI, and nonfatal and fatal stroke). The Pooled Cohort Equations should be used to estimate 10-year ASCVD risk for individuals without clinical ASCVD or diabetes and LDL-C 70 to 189 mg/dL to guide the initiation of statin therapy. For the primary prevention of ASCVD in individuals with diabetes (diabetes mellitus type-1 and type-2), estimated 10-year ASCVD risk can also be used to guide the intensity of statin therapy. For those with clinical ASCVD or with LDL-C \geq 190 mg/dL who are already in a statin benefit group, it is not necessary to estimate 10-year ASCVD risk (Stone et al, 2014).
- Statins are the primary medications to utilize for ASCVD risk reduction according to the 2013 guidelines, which focus on treatments proven to reduce ASCVD and not comprehensive lipid management.
- The 2015 AHA Scientific Statement on Familial Hypercholesterolemia (FH) recommends aggressive pharmacological treatment for patients with HeFH beginning at age eight to 10 years. Pharmacological treatment may also be considered in younger patients (less than eight years of age) with extreme elevation of LDL-C or those with other major risk factors suggesting very premature CVD. In HeFH pediatric patients, LDL-C goals are not well defined; however, treatment is recommended based on LDL-C levels and not based on genetic abnormalities or other clinical features. In adult patients with HeFH, the initial goal is to reduce LDL-C by 50% and treatment with a high-intensity statin (rosuvastatin or atorvastatin) is recommended. If LDL-C levels remain above goal after three months, then ezetimibe may be added. If LDL-C continues to be above goal after three months of two-drug therapy, then the addition of a PCSK9 inhibitor, bile acid sequestrant, or niacin can be considered. In patients with HoFH, lipid-lowering therapy should be initiated as soon as possible, with statins providing a 10 to 25% reduction in LDL-C (Gidding et al, 2015).
- The 2016 United States Preventive Services Task Force (USPSTF) recommendations for statin use for the primary prevention of cardiovascular disease in adults note the following:
 - Adults without a history of CVD should use a low- to moderate-dose statin for the prevention of CVD events and mortality when the following criteria are met: (1) they are aged 40 to 75 (2) they have one or more CVD risk factor such as dyslipidemia, diabetes, hypertension, or smoking (3) they have a calculated 10-year risk of a cardiovascular risk of 10% or more.
 - Although statin use may be beneficial for the primary prevention of CVD in some adults with a 10-year cardiovascular risk of <10%, the benefits are likely smaller. A low- to moderate-dose statin may be offered to certain adults without a history of CVD when all of the following criteria are met: (1) they are aged 40 to 75 years (2) they have one or more CVD risk factor (3) they have a calculated 10-year risk of a cardiovascular event of 7.5 to 10%.
 - There is insufficient evidence to assess the balance of benefits to risks of initiating a statin for the primary prevention of CVD and mortality in patients \geq 76 years without a history of MI or stroke (US Preventative Task Force, 2016).
- In 2017, the American Association of Clinical Endocrinologists/American College of Endocrinology (AACE/ACE) recommended the addition of another agent when statin therapy alone does not achieve therapeutic goals; their guidance offers cholesterol absorption inhibitors, bile acid sequestrants, and proprotein convertase subtilisin/kexin type 9 (PCSK-9) inhibitors as options (Jellinger et al, 2017). The recommendations for statin therapy for managing dyslipidemia and prevention of cardiovascular disease are stated as the following:
 - Statin therapy is recommended as the primary pharmacologic agent to achieve target LDL-C goals on the bases of morbidity and mortality outcome trials.
 - For clinical decision making, mild elevations in blood glucose levels and/or an increased risk of new-onset type 2 diabetes mellitus associated with intensive statin therapy do not outweigh the benefits of statin therapy for ASCVD risk reduction.
 - In individuals within high-risk and very high-risk categories, further lowering of LDL-C beyond established targets with statins results in additional ASCVD event reduction and may be considered.
 - Very high-risk individuals with established coronary, carotid, and peripheral vascular disease, or diabetes who also have at least 1 additional risk factor should be treated with statins to target a reduced LDL-C treatment goal of <70 mg/dL.
 - Extreme-risk individuals should be treated with statins to target an even lower LDL-C treatment goal <55 mg/dL.

- Numerous clinical trials have demonstrated that the statins (single entity and combination products) can effectively lower LDL-C, non-HDL-C, TC, and TG, as well as positively impact other lipid/lipoprotein parameters. Many studies have compared active treatment to placebo or compared combination therapy to monotherapy. In these studies, the more aggressive treatment regimens often improved lipid parameters to a greater extent than the less-intensive treatment regimens.
- All of the statins, with the exception of pitavastatin, have been shown to have beneficial effects on CHD outcomes, while the majority of them (atorvastatin, pravastatin, rosuvastatin, and simvastatin) have also been shown to decrease the risk of stroke.
- Atorvastatin, fluvastatin, pravastatin, rosuvastatin, and simvastatin have been shown to reduce cardiovascular events in patients with clinically evident CHD (secondary prevention). In addition, fluvastatin, lovastatin, pravastatin, and rosuvastatin have been shown to slow progression of coronary atherosclerosis in patients with CHD.
- No incremental benefit of the combination statin products on cardiovascular morbidity and mortality has been established over and above that demonstrated for the single entity statin products.
- The statins are generally well-tolerated, and the most common side effects are gastrointestinal disturbances, headache, insomnia, myalgia, and rash. Muscle aches and weakness are reported by one to two percent of patients taking statins. The symptoms are usually mild and generally do not lead to discontinuation. All statins can increase hepatic transaminase levels and creatinine kinase.
- Pravastatin is the only statin that does not undergo cytochrome (CYP) 450 metabolism, and is therefore associated with a lower risk for drug interactions. Atorvastatin (to a lesser extent), lovastatin, and simvastatin are primarily metabolized by the CYP3A4 isoenzyme, while fluvastatin, pitavastatin, and rosuvastatin are metabolized by the CYP2C9 isoenzyme, which may result in differences in their drug interaction profiles.
- There is insufficient evidence to support that one statin is safer or more efficacious than another statin.

Table 5. Advantages and Disadvantages of Statins

Drug	Advantages	Disadvantages
Atorvastatin	<ul style="list-style-type: none"> • Available generically both alone and in combination with ezetimibe • Has been documented to have more potency in cholesterol-lowering than certain other statins • Cardiovascular outcomes studies support the use of the 80 mg strength in certain populations (e.g., as secondary prophylaxis following ST elevation MI) 	<ul style="list-style-type: none"> • Associated with drug-drug interactions through the CYP3A4 isoenzyme system
Fluvastatin	<ul style="list-style-type: none"> • Available generically • Available in an extended-release formulation • Not associated with drug-drug interactions through the CYP3A4 isoenzyme system 	<ul style="list-style-type: none"> • Associated with drug-drug interactions through the CYP2C9 isoenzyme system
Lovastatin	<ul style="list-style-type: none"> • Available generically (immediate release formulation) • Available in an extended-release formulation 	<ul style="list-style-type: none"> • Associated with drug-drug interactions through the CYP3A4 isoenzyme system
Pitavastatin	<ul style="list-style-type: none"> • Not associated with drug-drug interactions through the CYP isoenzyme system 	<ul style="list-style-type: none"> • Effect on cardiovascular morbidity and mortality has not been determined
Pravastatin	<ul style="list-style-type: none"> • Available generically • Not associated with drug-drug interactions through the CYP isoenzyme system 	
Rosuvastatin	<ul style="list-style-type: none"> • Available generically 	

Drug	Advantages	Disadvantages
	<ul style="list-style-type: none"> Has been documented to have more potency in cholesterol-lowering than certain other statins 	
Simvastatin	<ul style="list-style-type: none"> Available as an oral suspension Tablet form is available generically Available both alone and in combination with ezetimibe 	<ul style="list-style-type: none"> Associated with drug-drug interactions through the CYP3A4 isoenzyme system

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