Statins: The Backbone of Treatment of Dyslipidemia

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Abstract: Statins are a panacea for secondary prevention of atherosclerotic cardiovascular disease and primary prevention in high-risk individuals. They are very well tolerated and side effects like muscle toxicity and increased risk of new onset of diabetes are seen in a minority of cases. They are also recommended in diabetic patient because the benefit is many times more than the risk of diabetes. Statins reduce total cholesterol, LDL cholesterol, Apo B, non-HDL cholesterol, and triglycerides, and also increase high-density lipoprotein (HDL) cholesterol levels in most patients with hypercholesterolemia and combined hyperlipidemia. Statins are not indicated in individuals with Frederickson Class I and V hyperlipidemias. Extensive literature supports use of statins in coronary heart disease (CHD) patients for treatment of dyslipidemia and secondary prevention. It has also been recognized that in secondary prevention and ACS populations lower LDL may be better. Trials have compared moderate with more robust LDL-C reduction, using maximum doses of atorvastatin or simvastatin. Available statins differ in their ability to reduce atherogenic lipoproteins and raise the level of high-density lipoprotein (HDL) cholesterol. Depending on dose used and specific statin, LDL cholesterol reduction of 18% to 55% can be expected. Atorvastatin and rosuvastatin are the most potent statins for lowering LDL-C cholesterol levels, yielding average reductions that approach 50% for atorvastatin and exceed 50% for rosuvastatin at the highest dose. Reduction in triglycerides with statins ranges from 7% to 30%, and is higher in hypertriglyceridemic populations and at higher statin doses. HDL levels usually rise by 5% to 10%. No consistent dose response relationship between statin dose and degree of HDL increase is seen.

Keywords: Dyslipidemia, Statin, Atherosclerosis

1. Background

Cardiovascular diseases cause 18 million deaths and a similar number of nonfatal cardiovascular events all over the world. [1] Dyslipidemia (LDL-C) account for approximately half the population-attributable risk of myocardial infarction [2] and approximately one quarter of the risk of ischemic stroke. [3]

Multitudinal studies have produced sufficient evidence for a causal relationship between blood cholesterol concentrations and cardiovascular and cerebrovascular disease. [4]

Although there has been noteworthy improvement in atherosclerotic cardiovascular disease (ASCVD) outcomes in recent decades, ASCVD remains the leading cause of morbidity and mortality globally. [4]

The indication for lipid-lowering therapy to prevent cardiovascular disease events in established cardiovascular disease is unambiguous. The benefit of lipid-lowering interventions for cardiovascular disease risk reduction is proven by mendelian randomization studies, prospective epidemiological cohort studies, and randomized trials. [5]

However, the decision on implementing a lipid-lowering intervention in the primary prevention setting is a major challenge in clinical practice for several reasons. Data on the association between the concentrations of the entire range of bloodstream lipids and very long-term cardiovascular outcomes in the general population are rather sparse.

Although HOPE-3 evaluated cholesterol lowering with the use of a low dose of rosuvastatin in a diverse population of persons who did not have cardiovascular disease and who were at intermediate risk. There was a significant reduction
in the risk of cardiovascular events with the use of rosuvastatin.

New evidence has confirmed that the key initiating event in atherogenesis is the retention of low-density lipoprotein (LDL) cholesterol (LDL-C) and other cholesterol-rich apolipoprotein (Apo) B containing lipoproteins within the arterial wall. Both European and American guidelines recommend that before starting medication in dyslipidemia patients total cardiovascular risk burden needs to be calculated. Prevention of ASCVD in a given person should relate to his or her total CV risk, the higher the risk, the more intense the action should be. Ideally, risk charts should be based on country-specific cohort data. But unfortunately, these are not available for most countries.

2. High Blood Cholesterol and ASCVD

Cholesterol, Lipoproteins, and Apolipoprotein B

Serum cholesterol and its lipoprotein carriers (LDL, very low-density lipoprotein [VLDL], and HDL) are known to be related to ASCVD. LDL-C is the dominant form of atherogenic cholesterol. VLDL is the chief carrier of triglycerides, and HDL cholesterol (VLDL-C) is also atherogenic. HDL-C is seemingly not atherogenic. Chylomicrons transport dietary fat; chylomicron atherogenicity is uncertain. The combination of LDL-C and VLDL-C is called non–HDL-C and is more atherogenic than either lipoprotein alone. The main protein embedded in LDL and VLDL is apolipoprotein B (apoB), and like non–HDL-C, apoB is a stronger indicator of atherogenicity than LDL-C alone. Multiple studies suggest that optimal total cholesterol levels are about 150 mg/dL (3.8 mmol/L), which corresponds to an LDL-C level of about 100 mg/dL (2.6 mmol/L). Adult populations with cholesterol concentrations in this range manifest low rates of ASCVD. [5, 6].

Measurement of LDL-C

The standard calculation method for LDL-C is the Friedewald formula: LDL-C=(TC)–(triglycerides/5)–(HDL-C)

When triglyceride levels are not elevated, this equation is sufficiently accurate. In hypertriglyceridemia, however, Friedewald-calculated LDL-C can be erroneous. In these cases measuring Apolipoprotein B is helpful.

Effect of food intake on cholesterol levels:

After normal food intake, LDL-C differs minimally with time. Fasting and nonfasting TC and HDL-C levels appear to have fairly similar prognostic value and associations with CVD outcomes. Thus, nonfasting samples can be used for risk assessment in primary prevention and for assessment of baseline LDL-C levels before the initiation of a statin in primary and secondary prevention. [7] In adults with a family history of premature ASCVD or genetic hyperlipidemia, a fasting lipid profile is reasonable for initial evaluation. The unreliability of the Friedewald-calculated LDL-C levels rise at lower levels of LDL-C.

Measurements of Apolipoprotein B and Lipoprotein (a)

Two lipoprotein entities related to LDL-C are apoB and lipoprotein (a) [Lp(a)]. Because apoB is the major apolipoprotein embedded in LDL and VLDL, several investigators identify strength of association between apoB and ASCVD. Others report a high correlation between apoB and non–HDL-C. Under certain circumstances, particularly in patients with hypertriglyceridemia, the measurement of apoB may have advantages. Nevertheless, apoB measurement carries extra expense, and its measurement in some laboratories may not be reliable. A relative indication for its measurement would be triglyceride ≥200 mg/dL. A level >130 mg/dL corresponds to an LDL-C level ≥160 mg/dL and constitutes a risk-enhancing factor. [8]

Lp(a) measurement should be considered at least once in each adult person’s lifetime to identify those with very high inherited Lp(a) levels >180 mg/dL (>430 nmol/L) who may have a lifetime risk of ASCVD equivalent to the risk associated with heterozygous familial hypercholesterolemia. Risk stratification for management of dyslipidemia

The general Framingham CVD risk score is being used by American Guidelines for management of blood cholesterol, adults are categorized as having low (<5%), borderline (5% to <7.5%), intermediate (7.5% to <20%), or high (>20%) 10-year risk. [9]

The Europeans uses The SCORE (Systematic Coronary Risk Estimation) system. The 2016 European Society of Cardiology/European Atherosclerosis Society Guidelines for the management of dyslipidemias and the 2016 European Guidelines on cardiovascular disease prevention in clinical practice, in that: (i) age has been extended from age 65 to 70; (ii) the interaction between age and each of the other risk factors has been incorporated, thus reducing the overestimation of risk in older persons in the original Systematic Coronary Risk Estimation charts; and (iii) the cholesterol band of 8 mmol/L has been removed since such persons will qualify for further evaluation in any event.

Familial dyslipidemias

Plasma lipid levels are, to a very large extent, determined by genetic factors. In its more extreme forms this is manifested as familial dyslipidemias. Among these, FH is the most common and is strongly related to CVD (Table 1).

2.1. Familial Combined Hyperlipidemia

The combination of ApoB >120 mg/dL and TGs >1.5 mmol/L >133 mg/dL) with a family history of premature CVD can be used to identify people who most probably have FCH. [10]

2.2. Familial Hypercholesterolaemia

2.2.1. Heterozygous Familial Hypercholesterolaemia

It causes premature CVD due to lifelong elevation of plasma levels of LDL-C. If left untreated, men and women with HeFH typically develop early CAD before the ages of 55 and 60 years respectively. The risk of CHD among individuals with definite or probable HeFH is estimated to be
increased at least 10-fold.

Cholesterol-lowering treatment should be initiated as soon as possible after a diagnosis has been made.

2.2.2. Homozygous Familial Hypercholesterolaemia

HoFH is a rare and life-threatening disease. The clinical picture is characterized by extensive xanthomas, marked premature and progressive CVD, and TC >13 mmol/L (>500 mg/dL). Most patients develop CAD and aortic stenosis before the age of 20 years and die before 30 years of age. The patients should be treated with intensive LDL-lowering drug therapy and, when available, with lipoprotein apheresis. This treatment (every 1-2 weeks) can decrease plasma LDL-C levels by 55-70%. [11]

2.2.3. Familial Hypercholesterolaemia in Children

FH is diagnosed in children based on phenotypic criteria including elevated LDL-C plus a family history of elevated LDL-C, premature CAD, and/or positive genetic testing. [12]

Treatment of children with FH includes a healthy lifestyle and statin treatment. A heart-healthy diet should be adopted early in life and statin treatment should be considered at 6-10 years of age. Statin treatment should be started with low doses and the dose should be increased to reach goals.

2.3. Familial Dysbetalipoproteinaemia

It produces a characteristic clinical syndrome in which both TC and TGs are elevated before treatment, usually both in the range of 7-10 mmol/L. In severe cases, patients develop tuberoeruptive xanthomas, particularly over the elbows and knees, and palmar xanthomata in the skin creases of the hands and wrists. The risk of CAD is very high, and accelerated atherosclerosis of the femoral and tibial arteries is also prevalent.

Most cases respond well to treatment with a statin or, if dominated by high TGs, a fibrate; often a combination of a statin and a fibrate may be needed.

Action to prevent acute pancreatitis is severe hypertriglyceridaemia

The risk of pancreatitis is clinically significant if TGs are >10mmol/L (880 mg/dL), particularly when occurring in association with familial chylomiconaemia, and actions to prevent acute pancreatitis are mandatory. [13, 14]

Restriction of calories and fat content (10-15% recommended) in the diet, and alcohol abstinence are obligatory. Fibrate therapy (fenofibrate) should be initiated, with n-3 fatty acids (2-4 g/day) as adjunct therapy. Lomitapide may also be considered in severe cases. In patients with DM, insulin therapy should be initiated to achieve good glycaemic control. In general, a sharp decrease of TG values is seen within 2-5 days. In the acute setting, plasmapheresis is able to rapidly lower TG levels. Volanesorsen has been recently approved by the EMA as an adjunct to diet in adult patients with genetically confirmed Familial Chylomiconaemia Syndrome (FCS) who are at high-risk for pancreatitis.

3. Therapeutic Modalities

3.1. Diet Composition, Weight Control, and Physical Activity

Patients should consume a dietary pattern that emphasizes intake of vegetables, fruits, whole grains, legumes, healthy protein sources (low-fat dairy products, low-fat poultry (without the skin), fish/seafood, and nuts), and non-tropical vegetable oils; and limit intake of sweets, sugar-sweetened beverages and red meats. This dietary pattern should be adjusted to appropriate calorie requirements, personal and cultural food preferences, and nutritional therapy for other medical conditions including diabetes. In general, adults should be advised to engage in aerobic physical activity 3-4 sessions per week, lasting on average 40 minutes per session and involving moderate-to vigorous-intensity physical activity.

3.2. Lipid-Lowering Drugs

Among lipid-lowering drugs, statins are the cornerstone of therapy, in addition to healthy lifestyle interventions. Other LDL-lowering drugs include ezetimibe, bile acid sequestrants, and PCSK9 inhibitors. Triglyceride-lowering drugs are fibrates and niacin; they have a mild LDL lowering action, but RCTs do not support their use as add-on drugs to statin therapy. [15]

3.2.1. Statin Therapy

The intensity of statin therapy is divided into 3 categories: high-intensity, moderate-intensity, and low intensity. [16]

High-intensity statin therapy typically lowers LDL-C levels by ≥50%, moderate-intensity statin therapy by 30% to 49%, and low-intensity statin therapy by <30% (Table 4).

Monitoring response of LDL-C to statin therapy:

In large RCTs of cholesterol-lowering therapy, LDL-C lowering has been consistently shown to reduce the risk of ASCVD. One large meta-analysis of statin clinical trials showed a progressive reduction in risk of major ASCVD events with lower on-treatment LDL-C levels. In another larger meta-analysis of 14 statin trials, it was observed that a 38.7-mg/dL (1-mmol/L) reduction of LDL-C levels is accompanied by a 21% reduction in ASCVD risk.

As a rough guide, a lowering of LDL-C levels of 1% gives an approximate 1% reduction in the risk of ASCVD—somewhat more at higher baseline LDL-C levels and somewhat less at lower baseline levels. [17]

3.2.2. Nonstatin Therapies

Ezetimibe is the most commonly used nonstatin agent. It lowers LDL-C levels by 13% to 20% and has a low incidence of side effects. Bile acid sequestrants reduce LDL-C levels by 15% to 30% depending on the dose. Bile acid sequestrants are not absorbed and do not cause systemic side effects, but they are associated with gastrointestinal complaints (eg, constipation) and can cause severe hypertriglyceridaemia when fasting triglycerides are ≥300 mg/dL (≥3.4 mmol/L). PCSK9 inhibitors are powerful LDL-lowering drugs. They
been shown to further reduce LDL-C levels by 43% to 64%

but if LDL-C levels remains ≥70 mg/dL (≥1.8 mmol/L) on

prefers statin-associated side effects because the large

possible adverse effects of a small increase in the  incidence

patients. SAMS often result in nonadherence and can

been observed. The number needed to cause one case of

presence of other risk factors for diabetes such as  overweight

or insulin resistance. Overall, the absolute reduction in the

risk of CVD in high-risk patients clearly outweighs the possible adverse effects of a small increase in the incidence of diabetes. [21, 22] This effect is probably related to the

mechanism of action of statins, as Mendelian randomization studies have confirmed the increased risk of DM in individuals with HMG-CoA reductase polymorphisms that reduce cholesterol synthesis.

3.3. Management of Dyslipidemia

Treatment goals according to European guidelines for low-density lipoprotein cholesterol across categories of total cardiovascular disease risk are summarized in Table 2.

Adults 40 to 75 years of age in primary prevention can be
classified as borderline risk (10-year risk of ASCVD (5% to <7.5%), intermediate-risk (7.5% to <20%), and high-risk (20%). For intermediate-risk patients, moderate- to high-intensity statin therapy should be considered during risk discussion of treatment options. Additional considerations favoring use of statins in intermediate-risk patients include other independent risk conditions and, in selected individuals, risk-enhancing factors associated with greater ASCVD risk. The statin eligibility criteria for primary prevention according to 5 major ASCVD guidelines are outlined in Table 3.

3.3.1. Treatment of Dyslipidemias in Older People

Treatment with statins is recommended for primary prevention, according to the level of risk, in older people aged <75. Initiation of statin treatment for primary prevention in older people aged >75 may be considered, if at high risk or above.

3.3.2. Treatment of Dyslipidemias in DM

In patients with T2DM at very-high risk, an LDL-C reduction of >50% from baseline and an LDL-C goal of <1.4 mmol/L (<55mg/dL) is recommended. In patients with T2DM at high risk, an LDL-C reduction of >50% from baseline and an LDL-C goal of <1.8 mmol/L (<70 mg/dL) is recommended. Statins are recommended in patients with T1DM who are at high or very-high risk. Intensification of statin therapy should be considered before the introduction of combination therapy. If the goal is not reached, statin combination with ezetimibe should be considered. Statin therapy is not recommended in pre-menopausal patients with DM who are considering pregnancy or not using adequate contraception.

The most important way to prevent ASCVD is to promote a healthy lifestyle throughout life. Prevention strategies must include a strong focus on lifestyle optimization (improvements in diet, physical activity, and avoidance of tobacco use and exposure to second hand smoke) to minimize the risk of future ASCVD events.

In adults at intermediate risk (>7.5% to <20% 10-year ASCVD risk) or selected adults at borderline risk (5% to <7.5% 10-year ASCVD risk), risk-based decisions for preventive interventions (e.g., statin therapy) remain uncertain, it is reasonable to measure a coronary artery calcium score to guide clinician–patient risk discussion.

Statin therapy is first-line treatment for primary prevention of ASCVD in patients with elevated low density lipoprotein
cholesterol levels (>190 mg/dL), those with diabetes mellitus, who are 40 to 75 years of age, and those determined to be at sufficient ASCVD risk after a clinician–patient risk discussion.

3.3.3. Management of Dyslipidemia in Women

Statin treatment is recommended for primary prevention of ASCVD in high-risk women. Statins are recommended for secondary prevention in women with the same indications and goals as in men.

Lipid-lowering drugs should not be given when pregnancy is planned, during pregnancy, or during the breastfeeding period. However, for severe FH patients, bile acid sequestrants (which are not absorbed) and/or LDL apheresis may be considered.

3.3.4. Severe Hypercholesterolemia

(LDL-C ≥190 mg/dL [≥4.9 mmol/L])

Patients with severe hypercholesterolemia have a high lifetime risk, and decisions about statins in these patients do not require ASCVD risk scoring. These patients derive net ASCVD risk reduction benefit from interventions that increase expression of LDL receptors. In selected patients with severe hypercholesterolemia whose LDL-C is inadequately controlled with drug therapy, LDL apheresis is an option.

4. Conclusion

Statins are proven agents for prevention of atherosclerotic cardiovascular disease and are recommended by all Guidelines throughout the globe. Statins are powered to reduce LDL-C by 1 mmol/L and this translates into reduction of cardiovascular events by 20 to 24%. If the LDL goals are not achieved with statins, non statins drugs like ezetimibe, bempedoic acid and PCSK-9 inhibitors are utilized.

| Table 1. Genetic disorders of lipoprotein metabolism. |
| --- |
| **Disorder** | **Prevalence** | **Gene (s)** | **Effect on lipoproteins** |
| HeFH | 1 in 200-250 | LDLR | Increased LDL-C |
| HoFH | 1 in 160 000-320 000 | LDLR, APO B, PCSK9 | Increased LDL-C |
| FCH | 1 in 100-200 | USF1, modifying genes | Increased LDL-C, VLDL-C and ApoB |
| Familial dysbetalipoproteinaemia | 1 in 5000 | APO B, PCSK9, ApoE | Increased IDL and chylomicron remnants |
| Familial lipoprotein lipase deficiency (familial chylomicron syndrome) | 2 in 106 | LPL, APO C2, ApoAV, GPIHBP1, LMF1 | Increased chylomicrons and VLDL-C |
| Tangier disease (analphalipoproteinaemia) | 1 in 106 | ABCA1 | Decreased HDL-C |
| Familial LCAT deficiency | 1 in 106 | LCAT | Decreased HDL-C |

Apo=apolipoprotein; FCH=familial combined hyperlipidemia; HDL-C=high-density lipoprotein cholesterol; HeFH=heterozygous familial hypercholesterolaemia; HoFH=homozygous familial hypercholesterolaemia; LDL-C=low-density lipoprotein cholesterol; LCAT=lecithin cholesterol acyltransferase; LDL-C=low-density lipoprotein cholesterol; VLDL=very low-density lipoprotein cholesterol

| Table 2. Treatment goals for low-density lipoprotein cholesterol across categories of total cardiovascular disease risk. |
| --- |
| **Category** | **Treatment Goals (LDL-C mg/dl)** |
| Low Risk | SCORE<1% |
| Moderate | SCORE ≥1% and <5% |
| •Young patients (T1DM <35 years; T2DM <50 years) with DM duration <10 years without other risk factors |
| •Markedly elevated single risk factor (310 mg/dL) or LDL-C >4.9 mmol/L (190 mg/dL) or BP ≥180/110 mmHg |
| •FH without other major risk factors |
| •Moderate CKD (eGFR 30–59 mL/min) |
| •DM w/o target organ damage, with DM duration ≥10 years or other additional risk factor |
| Very High|ASCVD (clinical/imaging) |
| •SCORE ≥10% |
| •FH with ASCVD or with another major risk factor |
| •Severe CKD (eGFR <30 mL/min) |
| •DM & target organ damage: ≥3 major risk factors; or early onset of T1DM of long duration ≥20 years | 116 |
| Treatment Goals | 100 |
| Moderate | SCORE ≥5% and <10% |
| •Markedly elevated single risk factor (310 mg/dL) or LDL-C >4.9 mmol/L (190 mg/dL) or BP ≥180/110 mmHg |
| •DM w/o target organ damage, with DM duration ≥10 years or other additional risk factor |
| Very High|ASCVD (clinical/imaging) |
| •SCORE ≥10% |
| •FH with ASCVD or with another major risk factor |
| •Severe CKD (eGFR <30 mL/min) |
| •DM & target organ damage: ≥3 major risk factors; or early onset of T1DM of long duration ≥20 years | 70 |
| Treatment Goals | 55 |
Table 3. Statin Eligibility Criteria According to 5 Major ASCVD Primary Prevention Guidelines.

| Criteria          | NICE                                      | USPSTF                                | CCS                  | ESC/EAS                          | ACC/AHA                           |
|-------------------|-------------------------------------------|---------------------------------------|----------------------|----------------------------------|-----------------------------------|
| Lipid-based       | Cholesterol level                         | LDL-C >190mg/dL or TC >290 mg/dL      | LDL-C ≥193 mg/dL     | LDL-C >232 mg/dL or TC >309 mg/dL| LDL-C ≥190 mg/dL or [TG >2.3 mmol/L (200 mg/dL)]. |
| Risk-based        |                                           |                                       |                      |                                 |                                   |
| Age range, yrs    | 40-75                                     | 40-75                                 | 40-75                | 40-65                            | >40 PCE ≥7.5% predicted 10-y risk of any ASCVD; LDL-C 70-189 mg/dL or diabetes (plus LDL-C level ≥70 mg/dL); risk-enhancers: family history of ASCVD, persistently elevated LDL-C levels ≥160 mg/dL, CKD, metabolic syndrome, persistently elevated triglycerides ≥175 mg/dL, hs-CRP levels ≥2.0 mg/dL; Lp(a) levels ≥50mg/dL, apoB levels ≥130mg/dL, and ankle-brachial index <0.9 |
| Eligibility       |QRISK2 ≥ 10% predicted 10- y risk of any ASCVD or nondialysis-dependent CKD| PCE ≥10% predicted 10-y risk of any ASCVD plus ≥1 ASCVD risk factor | FRS ≥20% predicted 10-y risk of any ASCVD or age 40-75 y; FRS ≥10% to <20% predicted 10-y risk of any ASCVD; LDL-C ≥135 mg/dL or diabetes or CKD (age ≥50 y) and eGFR level <60 mL/min/1.73m2 | LDL-C level ≥155 mg/dL; SCORE 5% to <10% predicted 10-y risk of fatal ASCVD or age 40-65 y; LDL-C >97 mg/dL; SCORE ≥10% predicted 10-y risk of fatal ASCVD or diabetes; or nondialysis-dependent CKD and eGFR level <60 mL/min/1.73m2 |

Table 4. High, Moderate and Low-Intensity Statin Therapy Percent LDL-C reductions with the primary statin medications use.

| Criteria          | High Intensity | Moderate Intensity | Low Intensity |
|-------------------|----------------|-------------------|---------------|
| LDL-C lowering    | ≥50%           | 30%–49%           | <30%          |
| Statins           | Atorvastatin (40 mg) 80 mg Rosuvastatin 20 mg (40 mg) | Atorvastatin 10 mg (20 mg) Rosuvastatin (5 mg) 10 mg Simvastatin 20–40 mg | Pravastatin 40 mg (80 mg) Lovastatin 40 mg (80 mg) Fluvastatin XL 80 mg Fluvastatin 40 mg BID Pitavastatin 1–4 mg |

Table 5. Drugs potentially interacting with statins metabolized by cytochrome P450 3A4 leading to increased risk of myopathy and rhabdomyolysis.

| Anti-infective agents | Calcium antagonists | Other              |
|-----------------------|---------------------|--------------------|
| Itraconazole          | Verapamil           | Amiodarone         |
| Ketoconazole          | Diltiazem           | Cyclosporine       |
| Posaconazole          | Amlodipine          | Danazol            |
| Erythromycin          |                     | Ranolazine         |
| Clarithromycin        |                     | Gemfibrozil        |
| Telithromycin         |                     | Grapefruit juice   |

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