Statin dose in primary prevention: aim for the target!

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Benefits from statin therapy for the primary prevention of atherosclerotic cardiovascular disease (ASCVD) are supported by high-quality evidence from randomised controlled trials (RCT). However, when and how to prescribe statins to individuals without ASCVD is a matter of debate. Strategies for initial management of low-density lipoprotein cholesterol (LDL-c) and goals of treatment differ among different guidelines (table 1).1–3 The National Institute for Health and Care Excellence (NICE) guidelines, for instance, recommend daily use of atorvastatin 20 mg in the primary care scenario when the estimated 10-year risk of ASCVD, as assessed by QRISK2, is at least 10%, whereas other guidelines propose different risk equations and different cut-offs. Also, the NICE document recommends clinicians to aim at lowering non-high-density lipoprotein cholesterol (non-HDL-c) by at least 40%, whereas other societies propose different eligibility criteria and different targets.

An important step after a guideline publication is the assessment of its uptake among health practitioners and patients in the real world, as well as of the impact of its adherence on clinical outcomes. These analyses may not only verify its appropriateness, providing feedback for continuous improvement of recommendations but also identify targets to optimise delivery of health to the society.

In this context, a study by Akyea et al investigating whether a suboptimal LDL-c response to statins, based on the NICE guidelines, increases the risk of future ASCVD events would prevent significantly more outcomes as compared with a 40% LDL-c reduction. On the other hand, there may be subgroups (higher risk and/or lower LDL-c) where a higher intensive statin (eg, >40% LDL-c reduction) would provide an incremental absolute benefit of small or negligible magnitude, as compared with a less intensive statin (eg, 30% LDL-c reduction). On the other hand, there may be subgroups (higher risk and/or higher LDL-c) where an even higher intensive statin therapy (eg, >50% LDL-c reduction) would prevent significantly more events as compared with a 40% LDL-c

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This study allows important insights to those who deal with cholesterol management in primary care. First, the high rate of suboptimal LDL-c response is clearly alarming, although it is probably overestimated for the present day, once the studied cohort was not contemporary (over 40% of the subjects were recruited before 2006 and managed under older guidelines). Even so, a substantial proportion of individuals who do not attain LDL-c targets recommended by guidelines, especially in high-risk conditions, has already been reported by other investigators.

In the Akyea et al study, not surprisingly, optimal responders, as compared with non-optimal responders, were initially prescribed medium potent statins more frequently and low potent statins less often. This probably explains why baseline LDL-c was higher in optimal responders, indicating that higher LDL-c motivates the physician to be more aggressive upfront.

There are also different patient-related causes for a suboptimal LDL-c response to statins. Although an interindividual response to statins may occur according to the genetic background, most cases where LDL-c response is less than expected are probably due to lack of adherence or persistence to the treatment. Even though statin-associated side effects do prevent statin use in many patients, the nocebo effect, that is, the report of a side effect by the patient actually due to negative expectation, is a frequent cause for interrupting statins among the population. Of note, poor adherence to lipid-lowering therapy, together with low-intensity therapy (as opposed to high-intensity treatment), is associated with higher cardiovascular risk. In addition, continued statin use after an adverse reaction has been shown to be related to a lower incidence of death and ASCVD events.

Important issues on blood cholesterol management in primary prevention could not be addressed or answered by the Akyea et al study. NICE recommendation (>40% non-HDL-c reduction for everyone in primary prevention eligible for statins) is practical and facilitates guideline uptake by clinicians. However, the absolute benefit from statins depends on the baseline LDL-c and ASCVD risk.4 There may be subgroups (lower risk and/or lower LDL-c) where a higher intensive statin (eg, >40% LDL-c reduction) would provide an incremental absolute benefit of small or negligible magnitude, as compared with a less intensive statin (eg, 30% LDL-c reduction). On the other hand, there may be subgroups (higher risk and/or higher LDL-c) where an even higher intensive statin therapy (eg, >50% LDL-c reduction) would prevent significantly more events as compared with a 40% LDL-c
lowering. For example, a recent meta-analysis demonstrated that more intensive LDL-c-lowering therapy, compared with less potent regimens, is associated with greater reduction in total and cardiovascular mortality when baseline LDL-c is >2.6 mmol/L, but not when LDL-c is lower. Therefore, clinicians should use judgement to refine statin therapy beyond general guideline recommendations on a case-by-case basis in their practice, though one should always consider the incremental benefit of more aggressive statin use.

Another point that should deserve further investigation is the comparative analyses of the effectiveness of different guidelines on statin use for the primary prevention of ASCVD. As shown in Table 1, criteria for initiate statin therapy and the goals of treatment vary according to the guideline used. These differences are not subtle and may have relevant clinical implications in ASCVD prevention. In this regard, a modelling study from The Copenhagen General Population Study concluded that more events could be prevented by recommending more widespread use of statins compared with more restrictive guidelines.

In conclusion, the study by Akyea et al brings a clear message for both clinicians and patients, highlighting the importance of LDL-c lowering for primary prevention of ASCVD, specifically supporting the concept that not achieving NICE guideline recommendation is a risk factor for ASCVD.

Effective implementation of guidelines among healthcare practitioners and the general population has been a challenge for a long time. Both physicians and patients should be targets for approaches aiming at improving adherence to guidelines. For the clinicians, these strategies should include serious programmes of continuing medical education and the development of in-house protocols. Simplified recommendations and algorithms may also facilitate guideline implementation among physicians. Patients and society should be educated on the scientific evidence documenting the benefits of lipid-lowering therapy and antistatin propaganda based on pseudoscience should be strongly disavowed and demystified by health authorities. Reassurance of statin safety should be emphasised to both doctors and patients in order to diminish excessive, unrealistic concerns. Global efforts and multidirectional strategies are necessary to optimise the management of blood cholesterol in primary care and reduce the burden of ASCVD.

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REFERENCES


