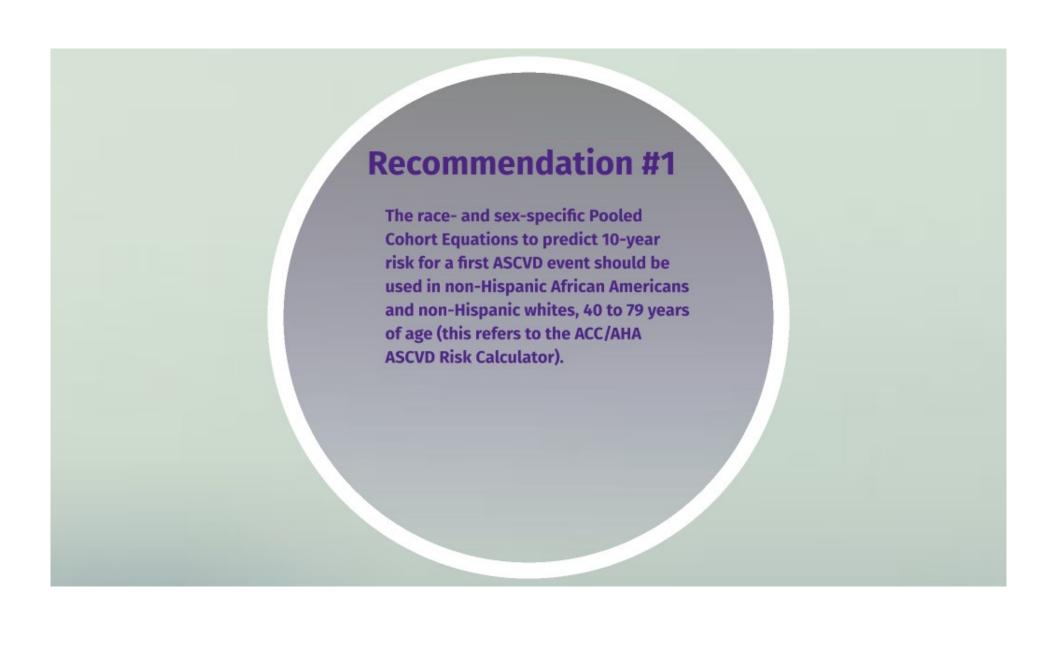


About the Guideline

- Developed by the American College of Cardiology/American Heart
 Association task force on practice guidelines in collaboration with
 the National Heart, Lung, and Blood Institute (NHLBI) in an effort
 to provide guidelines for the assessment of cardiovascular (CV)
 risk to promote optimal CV health and prevent CV disease.
- The objective of this publication was to develop an approach to quantify risk assessment in order to guide care and to address key clinical questions surrounding the assessment of CV risk targeting asymptomatic patients in the general population; i.e. those with no history of atherosclerotic cardiovascular disease (ASCVD).
- The three major areas of focus were developing a risk assessment tool and addressing 2 key clinical questions (CQ) for which 7 recommendations were made.
- The intent of the guideline is to target adults with no signs or symptoms of atherosclerotic cardiovascular disease (ASCVD) who would benefit from primary prevention.







CQ1: "What is the evidence regarding reclassification or contribution to risk assessment when high-sensitivity C-reactive protein (hs-CRP), apolipoprotein B (ApoB), glomerular filtration rate (GFR), microalbuminuria, family history, cardiorespiratory fitness, ankle-brachial index (ABI), carotid intima-media thickness (CIMT), or coronary artery calcium (CAC) score are considered in addition to the variables that are in the traditional risk scores?" (Goff et al., 2014).

This question was chosen to improve risk estimation beyond the traditional risk factors with markers for CV disease that could improve risk assessment and warrant routine measurement in clinical practice.

Recommendation #3

If, after quantitative risk assessment, a riskbased treatment decision is uncertain, 1 or more of the following may be considered to inform treatment decision making (Goff et al., 2014):

- Family history of premature CVD (1st degree relative): male < 55, female < 65
- · hs-CRP: ≥ 2mg/L
- CAC score ≥ 300 Agatston units or ≥ 75th percentile for age, sex, ethnicity
- · ABI < 0.9





CQ2: "Are models constructed to assess the long-term (≥ 15 years or lifetime) risk for a first cardiovascular disease (CVD) event in adults effective in assessing variation in long-term risk among adults at low and/or intermediate short-term risk, whether analyzed separately or combined?" (Goff et al., 2014).

The intent of this question was to evaluate models to use in addition to the 10-year risk assessment algorithm (ACC/AHA ASCVD Risk Calculator) to further assess lifetime risk and long-term outcomes, more particularly in those with low 10-year predicted risk.



Recommendation #7

Assessing 30-year or lifetime risk of ASCVD risk based on traditional risk factors may be considered in those aged 20 to 59 who are free from ASCVD and who are not at high short-term risk.

Note: Although there was not data supporting the use of lifetime risk to guide pharmacologic therapy decisions, it could potentially be used as information to communicate and motivate therapeutic lifestyle changes in younger adults starting at age 20 (despite the threshold of age 40 for the traditional 10-year ASCVD risk assessment) (Goff et al., 2014).

