Applying the 2005 Canadian Hypertension Education Program recommendations: 2. Assessing and reducing global atherosclerotic risk among hypertensive patients

E vidence from large population-based studies has confirmed that hypertension increases a patient's risk of cardiovascular disease about 2- to 3-fold. Even after adjusting for other risk factors, the presence of borderline systolic hypertension (≥ 160 mm Hg) alone increases the relative risk of a cardiovascular event by about 1.5.

Conversely, lowering blood pressure is known to reduce the risk of major cardiovascular events. Antihypertensive treatment, for example, reduces the risk of coronary artery disease by about 15% and cerebrovas-

cular events by about 35%.1

However, hypertension is only 1 risk factor for cardiovascular disease, and it has become evident that a patient's prognosis depends on the number and severity of atherosclerotic risk factors present.

The 2005 Canadian Hypertension Education Program recommendations emphasize risk factor assessment in managing hypertension.² Using 2 hypothetical clinical cases, we illustrate how global atherosclerotic risk is assessed and how those assessments are applied in hypertension management.

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Case 1: James is a 45-year-old man in whom hypertension has recently been diagnosed. A smoker, he has a history of dyslipidemia. His blood pressure is currently 160/100 mm Hg, and laboratory results show a total cholesterol level of 6.8 mmol/L, a low-density lipoprotein level of 5.2 mmol/L, a triglyceride level of 3.2 mmol/L, a high-density lipoprotein cholesterol (HDL-C) level of 1.0 mmol/L, and a cholesterol:HDL-C ratio of 6.8.

CASE 2: Sarah is a 45-year-old female marathon runner with hypertension. Her blood pressure has been consistently elevated at 160/100 mm Hg. She has no other atherosclerotic risk factors, and her blood lipid levels are excellent. She has a total cholesterol level of 5.8 mmol/L, a low-density lipoprotein level of 3.7 mmol/L, a triglyceride level of 1.3 mmol/L, a HDL-C level of 1.8 mmol/L, and a cholesterol:HDL-C ratio of 3.2.

Question: How do you determine the overall cardiovascular risk for patients with hypertension?

Comment: A number of methods to calculate a patient's absolute cardiovascular risk are now available, including the Framingham risk equation, the Cardiovascular Life Expectancy Model and the Dundee coronary risk disk. The Framingham risk equation is one of the most frequently used, and it takes into account age, sex, systolic and diastolic blood pressure, cholesterol levels, presence of diabetes mellitus and smoking status.

It estimates the 10-year risk of coronary artery disease (CAD) (defined as angina pectoris, recognized and unrecognized myocardial infarction, coronary insufficiency and CAD-related death).

Question: What are the 10-year risks of CAD for James and Sarah?

Comment: Using the Framing-ham risk equations (http://hin .nhlbi.nih.gov/atpiii/calculator.asp?usertype=prof or www.chiprehab.com/cvd for a bilingual version calibrated for Canada), James has

a 24.1% risk of CAD over the next 10 years. Although Sarah has the same blood pressure as James, her risk is much lower (4.1%) since she has no other atherosclerotic risk factors.

Question: Would the benefit of treatment be greatest for Sarah, who has hypertension only, or for James, who has the same blood pressure elevation as Sarah but more atherosclerotic risk factors?

Comment: Although both randomized trials and observational studies have consistently demonstrated a continuous increase in the relative risk of cardiovascular disease with increasing levels of blood pressure, the absolute risk can in fact be higher at lower levels of blood pressure if other risk factors are present. Similarly, the absolute benefit of antihypertensive therapy depends not only on the blood pressure level but also on the absolute risk of disease based on the number of cardiovascular risk factors present.3 As shown in Table 1, the number of pa-

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tients needed to treat to prevent a CAD-related event at 10 years is much lower for James, in whom the absolute risk is higher, than for Sarah. Table 1 also illustrates that the effectiveness of treatment depends more on the absolute risk of disease than on the blood pressure level.

Question: If the greatest treatment benefit is experienced by patients with the most risk factors, should lower blood pressure targets be used for these high-risk patients?

Comment: The optimal target blood pressure for safely reducing the risk of cardiovascular events has been validated for patients with diabetes and hypertension but remains uncertain for patients with specific atherosclerotic risk factors. Among patients with diabetes, one of the groups at highest risk, the results of the Hypertension Optimal Treatment (HOT)⁴ trial

showed an approximate 50% reduction in major cardiovascular events for patients in the lowest category of target diastolic blood pressure ($\leq 80 \text{ mm Hg}$) compared with the highest category ($\leq 90 \text{ mm Hg}$). Optimal target blood pressures in other populations at high risk of cardiovascular disease remain to be determined.

Question: James has a number of risk factors that require treatment. Should any 1 risk factor in particular be targeted to achieve the greatest reduction in his risk of future cardiovascular events?

Comment: Given that the greatest impact in blood pressure lowering has been demonstrated among patients with the greatest number of risk factors, emphasis should be placed on targeting these high-risk populations. However, rather than targeting 1 risk factor alone, efforts

should be made to treat all risk factors. Table 2 shows the impact of treating James's risk factors (returning lipid levels to normal and stopping smoking), as well as lowering his blood pressure. Treatment of each additional risk factor reduces his CAD risk and reduces the number needed to treat to prevent a CAD-related event at 10 years, compared with his original high-risk profile.

Question: Can these prediction models for estimating cardiovascular risk be applied to all populations?

Comment: Unfortunately, even well-validated models, when developed in one patient population, may be less accurate when applied to other populations. For instance, although the Framingham equations have been shown to reliably discriminate between low- and high-risk patients, they may be inaccurate

Table 1: Absolute coronary risk and number of patients needed to treat (NNT)* to prevent a coronary event over 10 years for 2 patients with different risk profiles, using Framingham estimates of coronary risk and change in blood pressure

Risk profile	Absolute 10-year coronary risk			
	Baseline blood pressure 160/100	Blood pressure treated to 150/95	Blood pressure treated to 139/89	
James • male • 45 yrs • dyslipidemia • smoker	24.1%	22.2% (NNT = 53)	17.9% (NNT = 16)	
Sarah • female • 45 yrs • no other risk factors	4.1%	3.3% (NNT = 125)	2.4% (NNT = 59)	

^{*}The NNT is calculated as NNT= 1/ absolute risk reduction as a proportion.

Table 2: Absolute coronary risk and number of patients needed to treat (NNT) to prevent a coronary event over 10 years for 1 patient with a changing risk profile, based on Framingham estimates of coronary risk and change in blood pressure, lipid levels and smoking cessation

Risk profile for 45-year-old male	Absolute 10-year coronary risk		
	Baseline blood pressure 160/100	Blood pressure treated to 150/95	Blood pressure treated to 139/89
 dyslipidemia smoker	24.1%	22.2% (NNT = 53)	17.9% (NNT = 16)
lipid levels back to normal has stopped smoking	9.4% (NNT = 7)*	8.6% (NNT = 6)*	6.8% (NNT = 6)*

^{*}The NNT is calculated on the assumption that returning lipid levels to normal and stopping results in an absolute risk reduction comparable to those who have never smoked and never had dyslipidemia.

in estimating absolute cardiovascular risks. The estimate of absolute risk may require additional calibration for a specific population. If potential limitations are recognized, these prediction models still have an important role to play in assessing risk and guiding treatment.

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dustry-sponsored, investigator-initiated, clinical trials assessing the usefulness of global risk assessment in clinical practice.

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Articles to date in this series

McAlister FA, Wooltorton E, Campbell NRC. The Canadian Hypertension Education Program (CHEP) recommendations: launching a new series. *CMAJ* 2005;173(5):508-9.

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