The difficulty in life is the choice.
— George Moore, The Bending of the Bough

Health care professionals caring for patients with cardiovascular disease are faced with an increasingly complex array of difficult choices. In this issue (page 1233), Madhu Natarajan and colleagues\textsuperscript{1} describe how they evaluated over 8000 patients waiting for cardiac catheterization within the Hamilton, Ont., region during a 2-year period ending in March 2000. Only 37\% of the procedures were completed within the waiting times requested by the referring physicians. A disproportionately high fraction of the 50 deaths, 32 myocardial infarctions (MIs) and 156 episodes of unstable angina that occurred while patients were waiting for catheterization were in outpatients, who represented 59\% of all requests. Not surprisingly, multivariate analysis showed that those at greatest risk were older and had symptoms suggesting more advanced disease. Aortic stenosis and depressed left ventricular ejection fraction were also powerful predictors of unfavourable events in outpatients. Although central booking existed and procedure waiting times were prioritized according to urgency, this approach applied only to inpatients, whose names were then added to the central catheterization list. In contrast, outpatients were triaged to individual angiographers. This process was presumably modulated by the angiographers’ schedule, available laboratory time and their own prioritization.

Although some useful clues for practitioners are provided in this report, it is unclear whether the rather high proportion (33.5\%) of patients with normal coronary arteries or mild coronary disease underwent appropriate risk stratification with noninvasive testing, what this revealed and how it was applied to the waiting time for cardiac catheterization. In contrast, those deemed to be at high risk on stress testing and those with low left ventricular ejection fraction, advanced symptoms or recent MI were proportionately more likely to wait beyond their requested waiting time. These findings seem somewhat paradoxical. This disconnect between the knowledge of factors associated with increased risk and the length of procedural delay provides insight into a more opportune use of a limited resource. Hence, patients identified through appropriate risk stratification to be at low risk should have the longest delay. We are also uncertain what proportion of inpatients had unstable angina or acute MI. Notwithstanding the priority for such patients, there exists, even within this subset, a genuine hierarchy of urgency that should influence procedure timing.

Are these Hamilton waiting times reasonable, and is the frequency of adverse events while waiting for catheterization acceptable? Arguably, any death of a patient on a waiting list is 1 death too many. Yet a median waiting time of 60 days for outpatient catheterization is likely to be representative of many other Canadian institutions. Sources of delay are unfortunately many (Table 1). For patients presenting with acute coronary syndromes, the need for timely urgent intervention can be sensibly guided by intelligent and contemporaneous risk assessment that continually assesses response to medical therapy. Depending on these factors, we should be aiming at intervention either immediately or within 48 hours of admission.\textsuperscript{2,3}

Natarajan and colleagues\textsuperscript{1} have done us a service by recalling the need for appropriate audit at individual centres concerning waiting times and adverse cardiac events. This experience again signals the need for national surveillance and standards to define and assess severity and urgency and to gauge expected benefit.\textsuperscript{4,5} Recent demonstration of improved outcomes among appropriately selected patients undergoing revascularization, as well as with the use of cardioverter/defibrillator devices for those at risk for or surviving sudden cardiac death, will only augment the already substantial pressure on Canadian cardiovascular resources.\textsuperscript{2,4} The need for systematic planning and cooperation among regional medical centres, provincial and federal governments, and national organizations, such as the Heart and Stroke Foundation of Canada and the Canadian Cardiovascular Society, has never been greater.

To be sure, the road ahead is filled with difficult choices,

Table 1: Sources of delay in access to cardiac catheterization

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<tr>
<th>Source of Delay</th>
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<tr>
<td>Patient’s recognition of symptoms</td>
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<tr>
<td>Assessment by primary physician</td>
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<tr>
<td>Primary physician’s investigation and evaluation</td>
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<tr>
<td>Referral to internists/cardiologists</td>
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<tr>
<td>Further evaluation, investigation and noninvasive testing</td>
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<tr>
<td>Waiting for cardiac catheterization</td>
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<td>Waiting for revascularization</td>
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but an approach such as that outlined in Table 2 can assist in the evaluation of patients awaiting cardiac catheterization. Systematic monitoring that is open and disseminated, with audit and feedback driving timely and effective triage, is the preferred method for the engagement of high-quality professionals working among rationally distributed resources. Over 5 years ago, a Canadian Cardiovascular Society national consensus conference examined this issue and called urgently for a national registry to collect accurate prospective data on performance characteristics and waiting times. Both that time and some patients have since expired: the book is overdue.

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References


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Table 2: Approach to managing waiting lists for cardiac catheterization

- Comprehensive assessment of key baseline risk factors
- Thorough noninvasive provocative stress testing
- Optimization of medical therapy, including evaluation of clinical response
- Systematic monitoring of waiting list, with appropriate reclassification as required
- Audit, feedback and dissemination of waiting lists and times
- Seamless movement to percutaneous coronary revascularization at the time of angiography if appropriate or triage to surgical revascularization

Organ and tissue donation in the intensive care unit

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When patients will not survive (e.g., after severe head injuries), intensive care unit (ICU) teams face the challenge of conducting empathetic, honest and compassionate discussions about organ and tissue donation with grieving families. Sometimes we fail to approach all families and neglect to seek consent for organ donation in all eligible circumstances. In consequence, alternative ways to achieve improved rates of consent to organ and tissue donation within ICUs have been proposed. In the United States, for example, these initiatives include mandating representatives of transplant procurement organizations rather than ICU or hospital-based teams to interact with families at the time of death in the ICU. In Canada, other initiatives have resulted in provincial legislation that requires ICU physicians to provide outside agencies with details of patients nearing death — details that would traditionally remain confidential. This particular requirement has proved to be the most contentious for critical care specialists.

The irony is that ICU physicians are as committed as any to the concept of successful organ and tissue donation, but we act and must act in the interests of our patient — the potential donor in the ICU — and his or her family. Transplant surgeons and transplant organizations have