## **Letters** | Correspondance

well-managed patients from warfarin to an NOAC is a disservice to our patients and to our health care system. Improved TTRs translate into fewer strokes and hemorrhages and lower health care costs.

The need to improve warfarin management in Canada. Like our non-family physician specialist colleagues, family physicians are not happy with the "standard care model" of warfarin management. It is inefficient and provides suboptimal INR control. We need government-funded access to better tools to provide optimal warfarin management. What are these

- Computer software that is capable of dosing warfarin and measuring TTR must replace the manual warfarin-dosing system.10 In addition, we need a single warfarin database in Canada as part of a national AF registry similar to Sweden's.
- Point-of-care INR testing must replace laboratory INR testing in most instances. Testing options need to be widened to provide patients with improved access and convenience. In New Zealand, the Community Pharmacy Anti-coagulation Management Service study achieved a mean TTR of 78.6% overall and 80.3% after 6 months.11
- · Patient self-management systems should use pointof-care INR testing in a structured program taught by diabetes or anticoagulation educators and be supervised by family physicians. Such programs have existed in Germany for 25 years; their TTRs average greater than 80% (Dr Stephan Kress, oral communication, September 2014). Patients in these programs are tested weekly. There are 200 000 German patients who self-manage. Patients who are unable to self-manage warfarin dosing usually have caregivers who are trained to assist them. We need to train caregivers.
- Use of 1-mg warfarin tablets in most cases instead of our 9 different warfarin strengths might simplify warfarin dosing, avoid tablet confusion, and permit daily or weekly dose adjustments of 0.5 mg. In Germany, patients in the self-management programs use 1-mg tablets only (Dr Stephan Kress, oral communication, September 2014).

To implement the use of these tools in Canada, we need government funding for the following elements:

- computer software (\$24 per patient per annum)<sup>12</sup>;
- point-of-care INR strips (\$7 per strip)<sup>13</sup> and devices (\$375 per device)<sup>14</sup>; and
- · patient training by diabetes or anticoagulant educators (4 hours per patient) (Dr Stephan Kress, oral communication, September 2014).

The total cost, including monitoring and the warfarin drug, is half the cost of the NOACs and provides TTRs greater than 70%, further reducing strokes, hemorrhages, and their associated costs to a level unattainable by NOACs. Finally, in addition with warfarin, we can now cheaply monitor the degree of anticoagulation and compliance (INR) plus the quality of warfarin management (TTR), and we can affordably and promptly reverse warfarin in the event of major or minor bleeding.

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### **Competing interests**

Dr Trusler is Vice President of INR Online Canada Limited, a not-for-profit Canadian company dedicated to the improvement of warfarin management

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# Response

le thank Dr Trusler for his interest in our articles, 1,2 but we disagree with his claim that we "ignored some very significant facts" pertaining to a comparison of the efficacy and safety of new oral anticoagulants (NOACs) with warfarin for stroke prevention in patients with atrial fibrillation (AF).

The objective of our articles<sup>1,2</sup> was not to compare NOACs with warfarin, which has been comprehensively reviewed elsewhere.3,4 Instead, we specifically explained the following:

[T]his review focuses on treating patients who are currently taking NOACs and does not consider the process for choosing an appropriate anticoagulant for AF or VTE [venous thromboembolism], whether an NOAC or warfarin.1

We also stated:

[Warfarin] remains a treatment option for patients with AF or VTE [venous thromboembolism] in whom excellent anticoagulation control is attainable.1

Rather than engage in a potentially protracted debate on the relative merits and drawbacks of NOACs and warfarin as anticoagulants, we urge readers to reach their own conclusions by reviewing the evidence and by considering patient values and preferences, as well as the costs, of these treatment options. We also would refer readers to clinical practice guidelines developed by the Canadian Cardiovascular Society, the American College of Chest Physicians, and the European Society of Cardiology, which endorse the use of NOACs as a firstline anticoagulant option for stroke prevention for most patients with AF based on at least comparable efficacy and safety, and less intracranial hemorrhage compared with warfarin 5-7

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### Competing interests

None declared

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# We stand by our conclusion

e thank Dr Lam for his comment pertaining to our Tools for Practice article on the effects of nonsteroidal anti-inflammatory drugs (NSAIDs) on fracture healing.2 Animal studies can be very useful for hypothesis generation; however, in this case the human and the animal data differ. The references Lam provided pertained to animal studies and the review article primarily also described animal studies.1 The human studies referenced in the Boursinos et al study<sup>3</sup> did not show a deleterious effect of NSAIDs on fracture healing.

Recently, another randomized controlled study comparing an NSAID (ie, ibuprofen) with morphine for children with uncomplicated fractures found that ibuprofen provided equivalent short-term pain relief with fewer adverse events. We hope that the authors will also report nonunion rates.

Until evidence from randomized controlled studies demonstrate adverse effects of NSAIDs on human fracture healing, we stand by our original conclusion that NSAIDs can be used for short-term pain relief for children and adults with orthopedic injuries or fractures.

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## Competing interests

None declared

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