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Commentary: NOAC? No problem

Edward Y. Sako, MD, PhD

Since approval by the Food and Drug Administration in 2010, the use of non-vitamin K oral anticoagulants (NOACs) as an alternative to vitamin K antagonists (VKAs) is increasing. In the accompanying article, Beller and colleagues¹ point out in their introduction that fixed dosing, lack of food–drug interactions, and absence of frequent laboratory monitoring are all factors behind this increased use. Because of this trend, they sought to examine the use of NOACs following cardiac surgery.¹ They compared 2 cohorts of patients spanning the time from the initial Food and Drug Administration approval of these agents with the 2 groups divided to give roughly equal numbers. They found, not surprisingly, that NOAC use has increased significantly over this time frame.

It is of interest that this increase is only relative to the use of VKAs. The overall percentage of anticoagulation with either a VKA or NOAC following bioprosthetic valve implantation was essentially unchanged over this time period. In other words, the availability of NOACs did not appear to influence the decision whether to prescribe an anticoagulant postoperatively, only which agent to use.

Efficacy has now been shown to the extent that updated guidelines for the management of atrial fibrillation allow for NOACs as first-line therapy in appropriate patients.² Therefore, the increase seen in the coronary artery bypass group, mostly for the treatment of atrial fibrillation, now at least has some support. One limitation of the study was that it was not possible to correlate preoperative use of anticoagulation with postoperative use. Therefore, it is not known in those patients who were on anticoagulation preoperatively whether the choice of preoperative agents was related to the choice postoperatively.

Perhaps a larger issue concerns patients receiving bioprosthetic valve replacements. Current guidelines provide for reasonable anticoagulation with a VKA in the first 3 to 6 months



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CENTRAL MESSAGE

The use of NOACs as the preferred method of anticoagulation is increasing in part due to their relative ease of use. This should not preclude the requisite studies of efficacy and safety.

after bioprosthetic mitral valve replacement or bioprosthetic aortic valve replacement.³ This is primarily to decrease the risk of thromboembolism during this period. The role of NOACs in these situations has yet to be determined.

As summarized by the authors, this study demonstrates that trends in the increased substitution of NOACs as the anticoagulant of choice are also seen in the post-cardiac surgical population. As the drivers may include increased ease of use, they are right to call for prospective clinical trials to assess efficacy. Furthermore, these trials should also examine relative effectiveness of the individual NOACs as well as variabilities in dosing.⁴

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From the Department of Cardiothoracic Surgery, University of Texas Health Science Center at San Antonio, San Antonio, Tex.

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Address for reprints: Edward Y. Sako, MD, PhD, Department of Cardiothoracic Surgery, University of Texas Health Science Center at San Antonio, Mail Code 7841, 7703 Floyd Curl Dr, San Antonio, TX 78229-3900 (E-mail: sako@uthscsa.edu).

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