The association between heart failure and stroke risk has been described for >3 decades. Hypercoagulability and left ventricular blood stasis are thought to contribute to stroke risk in patients with heart failure, in addition to common risk factors such as hypertension and atrial fibrillation. Regardless of the mechanism, however, for the majority of patients with heart failure who do not have atrial fibrillation, this elevated stroke risk is appreciated but not managed.

The current 2013 American College of Cardiology Foundation/American Heart Association Guideline on Management of Heart Failure states that “anticoagulation is not recommended in patients with chronic heart failure with reduced ejection fraction without atrial fibrillation, a prior thromboembolic event, or a cardioembolic source.” The 2012 European Society of Cardiology Guidelines for the Diagnosis and Treatment of Acute and Chronic Heart Failure similarly acknowledge the elevated stroke risk, but recommend against treatment with anticoagulation.

The recommendations are based on the results of previous randomized, controlled trials that assessed anticoagulation with warfarin in patients with heart failure but without atrial fibrillation. These include the Warfarin/Aspirin Study in Heart Failure (WASH), Heart Failure Long-Term Antithrombotic Study (HELAS), and Warfarin and Antiplaquette Therapy in Chronic Heart Failure (WATCH) trials, as well as the Warfarin versus Aspirin in Reduced Cardiac Ejection Fraction (WARCEF) trial that was the largest and was conducted with patients receiving contemporary heart failure therapy. Consistently, these trials failed to show the benefit of warfarin for primary outcomes that included death and stroke. In addition, warfarin therapy in these trials was associated with increased bleeding, although the risk for the most devastating bleeding outcome, intracranial hemorrhage, remained very low and was not significantly different than with aspirin.

It is important to note, however, that when the stroke outcome is considered by itself, both WARCEF and subsequent meta-analyses including all 4 trials show that warfarin can effectively prevent stroke in patients with heart failure in sinus rhythm. In the WARCEF trial, the risk of stroke was only 0.72% per year in the warfarin arm, in comparison with 1.36% per year in the aspirin arm. Nonetheless, these numbers are a fraction of other adverse events such as death and heart failure hospitalizations. It is thus likely that the stroke prevention benefit of warfarin is hidden by competing risks from other clinical events.

In this issue of Circulation, Abdul-Rahim et al shed further light on the frequency of stroke in patients with heart failure without atrial fibrillation, while highlighting the potential for using predicted stroke risk to refine patient selection. In a rigorously performed analysis, the authors pooled 9585 patients from the Controlled Rosuvastatin Multinational Trial in Heart Failure (CORONA) and Gruppo Italiano per lo Studio della Sopravvivenza nell'Infarto Miocardico Heart Failure (GISSI-HF) trials. In the subset of 6054 patients without atrial fibrillation, a stroke rate of 1.11% per year was observed. A risk model for stroke prediction was derived from 5 clinical predictors, including age, New York Heart Association Class, diabetes mellitus treated with insulin, body mass index, and a history of previous stroke. The model was validated by using an independent cohort from the Candesartan in Heart Failure Assessment of Reduction in Mortality and Morbidity–Added and Alternative Heart Failure and Reduced Ejection Fraction (CHARM HF-REF) trials. For patients classified in the highest tertile of stroke risk by the model, stroke rate was 1.98% per year in the derivation cohort and 1.79% per year in the validation cohort. In patients in whom N-terminal pro-B-type natriuretic peptide was measured, N-terminal pro-B-type natriuretic peptide further simplified and improved stroke risk prediction when added to the model.

However, in interpreting these results, some caution must be taken given this is a retrospective analysis that combined 2 different clinical trials. Another concern is that only 16% of the 6054 patients without atrial fibrillation received anticoagulant therapy at baseline, in comparison with 62% of patients with atrial fibrillation. The stroke rate in nonanticoagulated patients with atrial fibrillation was 2.17% per year, but, because anticoagulation was not randomized, this risk may be over- or underestimated. These factors may limit comparison between this group and the subgroup of patients without atrial fibrillation in the highest tertile of predicted risk.

Additionally, bleeding risk was not a part of this analysis, thereby precluding considerations of net clinical benefits that often serve as the basis for decisions regarding anticoagulation. However, the stroke rate for the highest-risk tertile was...
similar to the 1.9% annual rate of stroke seen in patients who have atrial fibrillation with a CHA2DS2-VASc score of 2, for whom anticoagulation is thought to have net clinical benefit and is recommended by current guidelines. A recent decision analysis by Eckman and colleagues suggests that, for nonwarfarin oral anticoagulants, the stroke risk threshold for net clinical benefit could even be as low as 0.9% per year. Nonwarfarin anticoagulants are also thought to have more consistent anticoagulation effect, and further analysis of the WARCEF patients has shown that, for patients with heart failure in sinus rhythm, higher-quality anticoagulation as reflected by the time-in-therapeutic range is associated with better clinical outcomes. It is thus reasonable to propose, as the authors do, that a randomized, controlled trial should test whether newer oral anticoagulants can prevent stroke in patients with heart failure in sinus rhythm.

A substantial number of patients are at risk. Of the estimated 5.7 million Americans with heart failure, 2.6 million will have reduced ejection fraction. With the use of conservative assumptions, three-quarters of those patients, or ≈1.9 million, will be in sinus rhythm. A stroke risk of 1.8% per year for the 630,000 patients in the highest tertile of risk would translate into ≈11,000 strokes each year.

The question of whether newer oral anticoagulants can prevent stroke in patients with heart failure in sinus rhythm can only be answered through an adequately powered randomized, controlled trial. The risk score for stroke prediction proposed by Abdul-Rahim et al, along with other selection approaches, could guide the identification of potential trial participants. Alternatively, given the improved risk-benefit profile of newer oral anticoagulants, a more inclusive trial design similar to WARCEF may be appropriate. A better characterization of bleeding risk in this patient population may also improve the risk–benefit balance and guide patient selection.

The choice of end points will also need to be carefully considered in such a trial, given the competing risks of death and heart failure hospitalizations observed in the WATCH and WARCEF trials. Because patients have consistently expressed strong preferences for avoiding severe stroke and often consider severe stroke worse than death, it will be reasonable to emphasize stroke outcomes in trial design. Ultimately, an innovative trial testing the effect of newer oral anticoagulants on stroke in patients with heart failure in sinus rhythm will represent an opportunity to align risk, treatment, and patient-centered outcomes.

Sources of Funding
Dr Ye is supported by a National Institutes of Health K23 career development award (K23 HL121144).

Disclosures
Dr Homma reports receiving consulting fees from Boehringer Ingelheim, Daiichi Sankyo, Bristol- Myers Squibb, Pfizer, and St. Jude Medical. Dr Ye reports no conflicts.

References
14. Eckman MH, Singer DE, Rosand J, Greenberg SM. Moving the tipping point: the decision to anticoagulate patients with atrial fibrillation with a CHA2DS2-VASc score of 2, for whom anticoagulation is thought to have net clinical benefit and is recommended by current guidelines.


Key Words: Editorials ■ anticoagulants ■ heart failure
Stroke and Anticoagulation in Heart Failure Without Atrial Fibrillation: From Risk to Opportunity
Shunichi Homma and Siqin Ye

Circulation. 2015;131:1465-1467; originally published online March 25, 2015; doi: 10.1161/CIRCULATIONAHA.115.016329
Circulation is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2015 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circ.ahajournals.org/content/131/17/1465

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in Circulation can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Circulation is online at:
http://circ.ahajournals.org/subscriptions/