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CHA₂DS₂-VASc Score and Adverse Outcomes in Patients with Heart Failure with Reduced Ejection Fraction and Sinus Rhythm

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Abstract

Aims—To determine whether the CHA₂DS₂-VASc score can predict adverse outcomes such as death, ischemic stroke, and major hemorrhage, in patients with systolic heart failure in sinus rhythm.

Methods and Results—CHA₂DS₂-VASc scores were calculated for 1,101 patients randomized to warfarin and 1,123 patients randomized to aspirin. Adverse outcomes were defined as death or

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**Please see attached supplementary file for full list of WARCEF investigators

Disclosures:

No other potential conflict of interest relevant to this article was reported.

ischemic stroke, death alone, ischemic stroke alone, and major hemorrhage. Using proportional hazards models, we found that each 1-point increase in the CHA₂DS₂-VASc score was associated with increased hazard of death or ischemic stroke events (hazard ratio [HR] for the warfarin arm = 1.21 [1.13–1.30], p<0.001; for aspirin, HR = 1.20 [1.11–1.29], p<0.001). Similar increased hazards for higher CHA₂DS₂-VASc scores were observed for death alone, ischemic stroke alone, and major hemorrhage. Overall performance of the CHA₂DS₂-VASc score was assessed using c-statistics for full models containing the risk score, treatment assignment, and score-treatment interaction, with the c-statistics for the full models ranging from 0.57 for death to 0.68 for major hemorrhage.

Conclusions—The CHA₂DS₂-VASc score predicted adverse outcomes in patients with systolic HF in sinus rhythm, with modest prediction accuracy.

Keywords

heart failure; warfarin; sinus rhythm; stroke; bleeding

Background

Patients with heart failure with reduced ejection fraction (HFrEF) are at increased risk for adverse outcomes, including death and stroke.(1, 2) For patients with HFrEF in sinus rhythm, there has been recent interest in assessing whether the risk of these adverse outcomes can be accurately predicted by existing risk scores that incorporate established risk factors.(3–5) In particular, although the CHA₂DS₂-VASc risk score (incorporating risk factors including congestive heart failure, hypertension, age ≥ 75 years [2 points], age 65–75 years, diabetes, stroke/transient ischemic attack/thromboembolism [2 points], vascular disease [prior myocardial infarction, peripheral artery disease, or aortic plaque], and female sex) was developed for patients with atrial fibrillation,(6) Melgaard and colleagues recently described its use for predicting the risk of death and stroke in HFrEF patients in sinus rhythm.(3) We therefore undertook this analysis of patients enrolled in the Warfarin versus Aspirin in Reduced Cardiac Ejection Fraction Trial (WARCEF),(7) to evaluate whether the CHA₂DS₂-VASc risk score can predict death, ischemic stroke, and major hemorrhage in patients with HFrEF in sinus rhythm who received warfarin or aspirin.

Methods

The protocol for the randomized, double blinded WARCEF trial (<http://www.ClinicalTrials.gov> No. NCT00041938) has been described previously.(7, 8) Briefly, patients with left ventricular ejection fraction (LVEF) ≥ 35% and who were in SR were randomized to receive warfarin or aspirin. Additional eligibility criteria included age ≥ 18 years old, having no contraindications to warfarin, having a modified Rankin score of 4 or less, and on evidence-based heart failure medications (beta-blocker, angiotensin-converting-enzyme inhibitor or angiotensin-receptor blocker, or hydralazine and nitrates). Patients were excluded if they had a clear indication for warfarin or aspirin, or if they had a condition that conferred a high risk of cardiac embolism. A total of 2,305 participants were recruited from 168 centers in 11 countries from October 2002 to January 2010. The investigation conforms

with the principles outlined in the Declaration of Helsinki.(9) All subjects provided informed consent.

For the WARCEF trial, ischemic stroke was defined as a clinically relevant new lesion detected on computed tomography or magnetic resonance imaging or, in the absence of a new lesion, clinical findings that were consistent with the occurrence of clinical stroke and that lasted for longer than 24 hours. Major bleeding was defined as intracerebral, epidural, subdural, subarachnoid, spinal intramedullary, or retinal hemorrhage; any other bleeding causing a decline in the hemoglobin level of more than 2 g per deciliter in 48 hours; or bleeding requiring transfusion of 2 or more units of whole blood, hospitalization, or surgical intervention. A blinded independent end-point adjudication committee adjudicated all stroke and major bleeding events.

For each participant, the CHA₂DS₂-VASc risk score was calculated based on the approach described by Lip and colleagues.(6) For this analysis, 81 participants were excluded because incomplete data precluded the calculation of the CHA₂DS₂-VASc score. Because stroke and bleeding risk are expected to differ for patients receiving warfarin and aspirin, we performed all analyses separately for the warfarin and aspirin arms of the WARCEF trial on an intent-to-treat (ITT) basis. Baseline characteristics of the study participants were compared by treatment group using two-sample t-test for continuous variables and Chi-squared test for categorical variables. For each CHA₂DS₂-VASc score, we calculated incidence rate per 100 patient-years for the composite outcome of death and ischemic stroke, as well as for the individual outcomes of death, ischemic stroke, and major bleeding. Exact Poisson 95% confidence intervals of the incidence rates were provided. To determine the association between CHA₂DS₂-VASc score and each outcome above, as well as to investigate whether the CHA₂DS₂-VASc score influenced the treatment effect of warfarin versus aspirin, we constructed Cox proportional hazard models that incorporated treatment assignment, the CHA₂DS₂-VASc score, and their interaction term. Model discrimination was assessed using Harell's *c*-index. Model calibration was assessed using the survival-adapted Hosmer-Lemeshow χ^2 goodness of fit test.(10) The test statistic measures the difference between the predicted and the observed probabilities of risk. A small *p*-value suggests that the model is not well calibrated. *P*-values of <0.05 were considered statistically significant for all testing. All statistical analyses were conducted using SAS version 9.4 (SAS Institute, Cary, NC).

Results

For this analysis, a total of 2,204 participants were included, of whom 1,101 were enrolled in the warfarin arm and 1,123 were enrolled in the aspirin arm. The mean (SD) age of participants were 60.8 (11.3) years old, and 435 (19.6%) were female. Baseline characteristics of WARCEF participants by treatment arm are described in Table 1.

Because all WARCEF participants had a diagnosis of heart failure, the minimum CHA₂DS₂-VASc score was 1; the maximum was 9. The composite outcome of death and ischemic stroke occurred in 588 (27%) of the 2,204 participants; the risk significantly increased with each 1-point increase in CHA₂DS₂-VASc score (Figure) for participants enrolled in both warfarin (hazard ratio [HR] 1.21, 95% confidence interval [CI] 1.13–1.30, *p*<0.001) and

aspirin arms (HR 1.20, 95% CI 1.11–1.29, $p < 0.001$). Similar trends were observed for the individual outcomes of ischemic stroke (Table 2), death alone and major hemorrhage (Table 3). For ischemic stroke, the incidence rate per 100 patient-years was largely above 1% for CHA₂DS₂-VASc scores of 5 and above for patients receiving warfarin, and for CHA₂DS₂-VASc scores of 2 and above for patients receiving aspirin.

The treatment effect of warfarin versus aspirin did not significantly differ according to the CHA₂DS₂-VASc score for all outcomes examined. Model discrimination was modest for the composite outcome of death and ischemic stroke (c -index 0.58, 95% CI 0.56–0.60) and the individual outcome of death (c -index 0.57, 95% CI 0.55–0.60), but was higher for ischemic stroke (c -index 0.64, 95% CI 0.57–0.70) and for major hemorrhage (c -index 0.68, 95% CI 0.62–0.73) (Tables 2 and 3). The survival-adapted Hosmer-Lemeshow goodness-of-fit test p -values were larger than 0.05 for all outcomes, suggesting that none of the models had a significant lack of fit.

Discussion

In this retrospective analysis of the WARCEF trial, we found that in patients with heart failure in sinus rhythm, the CHA₂DS₂-VASc score was moderately predictive of adverse outcomes, including death or ischemic stroke, and the individual outcomes of death, ischemic stroke, and major hemorrhage. Increased CHA₂DS₂-VASc risk score was consistently associated with increased stroke risk for both the aspirin and warfarin arms of the WARCEF cohort. There was no significant interaction between treatment assignment and the CHA₂DS₂-VASc score for these outcomes.

Our results are largely consistent with those reported by Melgaard and colleagues, who found that the CHA₂DS₂-VASc score predicted death, ischemic stroke, and thromboembolism, in a nationwide cohort of 33,592 heart failure patients with reduced or preserved ejection fraction without concurrent atrial fibrillation.⁽³⁾ In that study, the c -statistics for the CHA₂DS₂-VASc ranged from 0.63 to 0.69 for the individual outcomes. These values are slightly higher than those in our analysis. However, at each level of the CHA₂DS₂-VASc score, the absolute risk for ischemic strokes was higher in the Danish cohort, ranging from 1.5% per year for a score of 1 to 7% for a score of 6, where as across the same range of CHA₂DS₂-VASc scores in the WARCEF cohort the stroke risk was 0.3% to 1.1% in the aspirin arm and 0.4% to 1.1% in the warfarin arm. Potential reasons for this include the protective effects of aspirin and warfarin, inclusion of younger patients in WARCEF, and statistical imprecision due to the sample size. Regardless of the reason, these differences suggest additional analyses of independent cohorts are needed to better calibrate how the CHA₂DS₂-VASc score can predict stroke risk in heart failure patients in sinus rhythm, especially as the threshold for benefit from initiating anticoagulation is commonly thought to be a stroke risk of between 1–2%.^(11, 12)

In addition to the CHA₂DS₂-VASc score, other approaches to risk stratification for heart failure patients in sinus rhythm have been examined recently. Abdul-Rahim and colleagues have found that in patients with heart failure with reduced ejection fraction who are in sinus rhythm, risk scores that included clinical risk factors (age, New York Heart

Association class, diabetes treated with insulin, body mass index, previous stroke) and N-terminal pro B-type natriuretic peptide accurately predicted strokes.(4) Similarly, we have previously demonstrated that major bleeding in WARCEF participants could be predicted by existing bleeding risk scores, such as the Hypertension, Abnormal Renal/Liver Function, Stroke, Bleeding History or Predisposition, Labile INR, Elderly, Drugs/Alcohol Concomitantly (HAS-BLED) score and the Outpatient Bleeding Risk Index (OBRI).(5) Taken together, these findings all suggest that risk prediction in heart failure patients in sinus rhythm can be further refined. Although current guidelines do not recommend routine anticoagulation in this patient population,(13, 14) ongoing studies, such as the COMMANDER-HF randomized, controlled trial, are assessing the potential benefit of newer oral anticoagulants such as rivaroxaban.(15) Given the superior safety profile of these agents compared with warfarin, improved risk stratification of heart failure patients in sinus rhythm, especially with regards to stroke and bleeding risk,(16, 17) will greatly aid the identification of patients who may benefit from such therapy.(11)

The limitations of our analysis include its retrospective design and the modest number of stroke events. As mentioned previously, since WARCEF was a randomized, controlled trial, it is possible that healthier patients were selected, which could lead to underestimation of stroke risk and affect the generalizability of our findings. Similarly, since all patients in WARCEF received either warfarin or aspirin, the true stroke risk of heart failure patients in sinus rhythm who are not on anticoagulant therapy may be higher than described here. We did not construct a model including hemodynamic or laboratory parameters using WARCEF participants, and it is possible such a model may be superior to the CHA₂DS₂-VASc risk score. However, any such model would also need to be validated externally, while the CHA₂DS₂-VASc risk score is easy to calculate and already widely accepted, making it more likely to be clinically useful should future studies continue to confirm our findings. Finally, we did not formally monitor for the occurrence of asymptomatic atrial fibrillation during the WARCEF study, although a small number of participants (~10%) did develop clinical atrial fibrillation during the follow-up period.(18) Although this may have contributed to the relationship between the CHA₂DS₂-VASc score and clinical outcomes observed in our analysis, this does not change our finding that the CHA₂DS₂-VASc score represents a potentially useful tool for risk stratification in this patient population.

Despite these limitations, our findings are timely additions to the literature on understanding stroke risk in heart failure patients in sinus rhythm. Our analysis confirms that the CHA₂DS₂-VASc score can predict stroke risk in this population, and can potentially inform future studies on new therapeutic approaches for stroke prevention for heart failure patients in sinus rhythm.

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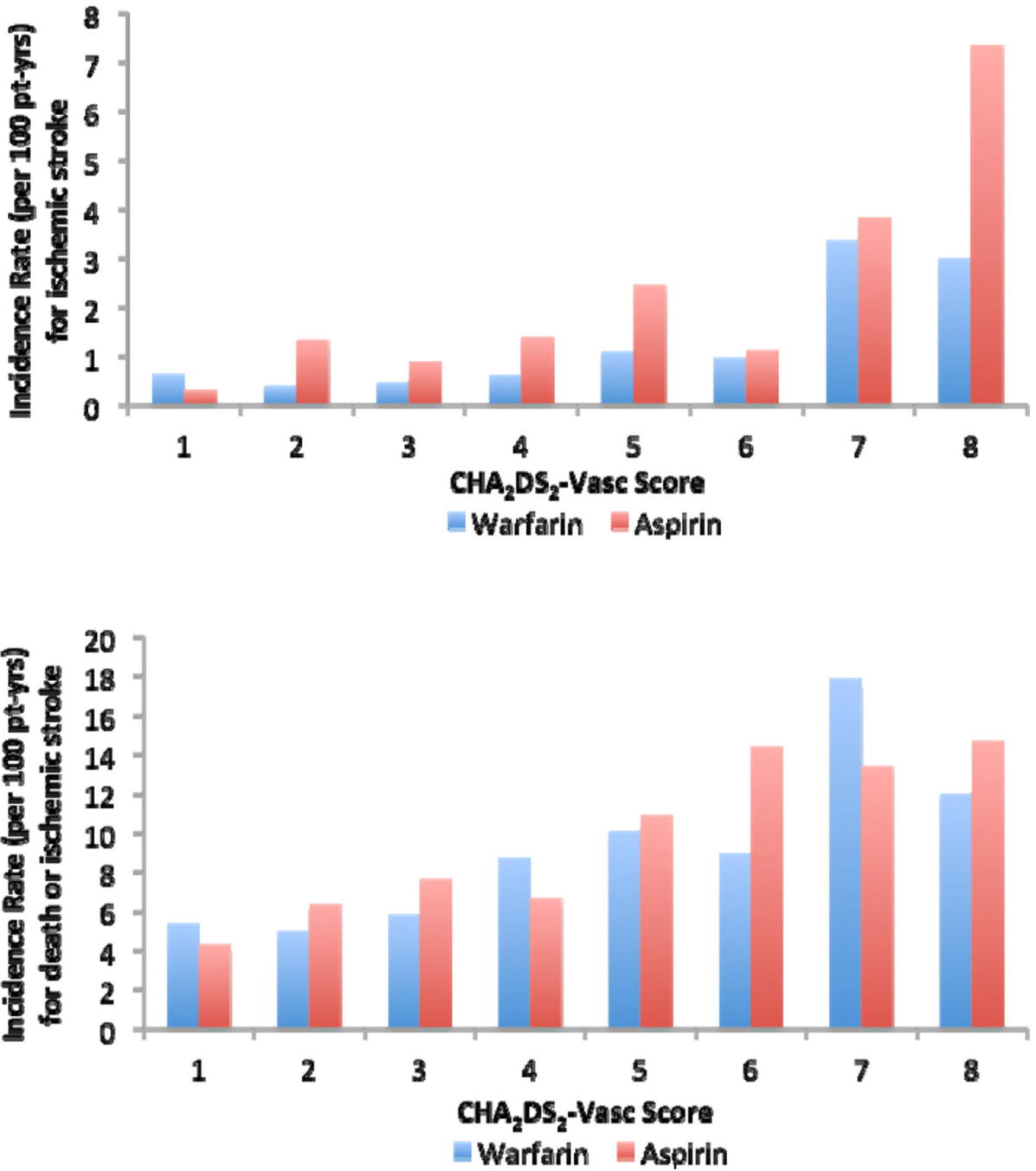


Figure 1. Adverse outcomes by CHA₂DS₂-VASc score. Incidence rate of ischemic stroke (top panel), and death or ischemic stroke (bottom panel), by categories of CHA₂DS₂-VASc score. Because only 1 patient had a CHA₂DS₂-VASc score of 9, data are not shown for that category.

Table 1

Baseline characteristics of the study participants, according to treatment group (warfarin versus aspirin). For continuous variables, mean \pm SD were reported, and *p*-values were calculated using two-sample *t*-test. For categorical variables, number /total number (%) were reported, and *p*-values were calculated using Chi-squared test.

Characteristic	Warfarin (n=1101)	Aspirin (n=1123)	p-value
Age - years	60.9 \pm 11.5	60.7 \pm 11.1	0.647
Male sex	879/1101 (79.8)	910/1123 (81.0)	0.477
Race or ethnic group			0.862
Non-Hispanic white	830/1101 (75.4)	856/1123 (76.2)	.
Non-Hispanic black	158/1101 (14.4)	153/1123 (13.6)	.
Hispanic	83/1101 (7.5)	79/1123 (7.0)	.
Other	30/1101 (2.7)	35/1123 (3.1)	.
Body-mass index - kg / m ²	28.9 \pm 5.9	29.2 \pm 6.0	0.197
Systolic blood pressure - mmHg	124.0 \pm 19.4	124.2 \pm 18.4	0.832
Diastolic blood pressure - mmHg	74.1 \pm 11.6	74.5 \pm 11.3	0.402
Pulse - beats/min	71.8 \pm 11.3	72.1 \pm 12.5	0.479
Hypertension	669/1101 (60.8)	692/1123 (61.6)	0.678
Diabetes Mellitus	359/1101 (32.6)	337/1123 (30.0)	0.187
Atrial Fibrillation	42/1101 (3.8)	38/1123 (3.4)	0.585
Peripheral Vascular Disease	130/1101 (11.8)	122/1123 (10.9)	0.483
Prior stroke or TIA	148/1101 (13.4)	137/1123 (12.2)	0.381
Ischemic Cardiomyopathy	475/1101 (43.1)	488/1123 (43.5)	0.882
Smoking status			0.424
Current smoker	211/1099 (19.2)	193/1122 (17.2)	.
Former smoker	563/1099 (51.2)	579/1122 (51.6)	.
Never smoked	325/1099 (29.6)	350/1122 (31.2)	.
NYHA classification			0.502
1	145/1098 (13.2)	160/1118 (14.3)	.
2	595/1098 (54.2)	626/1118 (56.0)	.
3	343/1098 (31.2)	319/1118 (28.5)	.
4	15/1098 (1.4)	13/1118 (1.2)	.
Ejection fraction - %	24.6 \pm 7.5	24.9 \pm 7.5	0.319

Table 2

Incidence rate per 100 patient-years for death or ischemic stroke and ischemic stroke only, by categories of CHA₂DS₂-VASc score. Hazard ratios are for each 1-point increase in CHA₂DS₂-VASc score, using models that contained treatment assignment, CHA₂DS₂-VASc score and their interaction.

		Death or Ischemic Stroke		Ischemic Stroke	
	CHA ₂ DS ₂ -VASc Score	Warfarin (285/1101)	Aspirin (303/1123)	Warfarin (27/1101)	Aspirin (54/1123)
no. of events/total no. of patients [Incidence rate per 100 patient-years] (95% CI)	1	17/85 [5.39] (3.14, 8.63)	14/82 [4.30] (2.35, 7.21)	2/85 [0.63] (0.08, 2.29)	1/82 [0.31] (0.01, 1.71)
	2	38/196 [4.97] (3.52, 6.82)	58/247 [6.36] (4.83, 8.22)	3/196 [0.39] (0.08, 1.15)	12/247 [1.32] (0.68, 2.30)
	3	63/296 [5.82] (4.47, 7.45)	69/267 [7.64] (5.95, 9.67)	5/296 [0.46] (0.15, 1.08)	8/267 [0.89] (0.38, 1.75)
	4	72/249 [8.74] (6.84, 11.0)	58/249 [6.66] (5.06, 8.61)	5/249 [0.61] (0.20, 1.42)	12/249 [1.38] (0.71, 2.41)
	5	46/142 [10.1] (7.36, 13.4)	62/179 [10.9] (8.33, 13.9)	5/142 [1.09] (0.35, 2.55)	14/179 [2.45] (1.34, 4.12)
	6	28/89 [8.96] (5.95, 13.0)	26/61 [14.4] (9.39, 21.1)	3/89 [0.96] (0.20, 2.81)	2/61 [1.11] (0.13, 3.99)
	7	16/32 [17.9] (10.2, 29.1)	14/33 [13.4] (7.33, 22.5)	3/32 [3.36] (0.69, 9.82)	4/33 [3.83] (1.04, 9.81)
	8	4/11 [12.0] (3.26, 30.6)	2/4 [14.7] (1.78, 53.1)	1/11 [2.99] (0.08, 16.6)	1/4 [7.35] (0.19, 41.0)
	9	1/1 [103] (2.61, 575)	0/1 [0.00] (0, 221)	0/1 [0.00] (0, 381)	0/1 [0.00] (0, 221)
HR (95% CI)		1.21 (1.13, 1.30) <i>p</i> <0.001	1.20 (1.11, 1.29) <i>p</i> <0.001	1.35 (1.08, 1.70) <i>p</i> =0.009	1.28 (1.08, 1.51) <i>p</i> =0.004
<i>p</i> -value for interaction		0.819		0.693	
C-index (95% CI)		0.58 (0.56-0.60)		0.64 (0.57-0.70)	
Goodness of fit test <i>p</i> -value		0.773		0.224	

Table 3

Incidence rate per 100 patient-years for death and major hemorrhage, by categories of CHA₂DS₂-VASc score. Hazard ratios are for each 1-point increase in CHA₂DS₂-VASc score, using models that contained treatment assignment, CHA₂DS₂-VASc score and their interaction.

		Death		Major Hemorrhage	
	CHA ₂ DS ₂ -Vasc Score	Warfarin (258/1101)	Aspirin (249/1123)	Warfarin (63/1101)	Aspirin (29/1123)
no. of events/total no. of patients [Incidence rate per 100 patient-years] (95% CI)	1	15/85 [4.76] (2.66, 7.85)	13/82 [3.99] (2.12, 6.82)	1/85 [0.32] (0.01, 1.77)	1/82 [0.31] (0.01, 1.71)
	2	35/196 [4.58] (3.19, 6.36)	46/247 [5.04] (3.69, 6.73)	10/196 [1.31] (0.63, 2.40)	5/247 [0.55] (0.18, 1.28)
	3	58/296 [5.36] (4.07, 6.93)	61/267 [6.76] (5.17, 8.68)	13/296 [1.20] (0.64, 2.05)	5/267 [0.55] (0.18, 1.29)
	4	67/249 [8.13] (6.30, 10.3)	46/249 [5.28] (3.87, 7.04)	12/249 [1.46] (0.75, 2.54)	8/249 [0.92] (0.40, 1.81)
	5	41/142 [8.95] (6.43, 12.2)	48/179 [8.41] (6.20, 11.2)	13/142 [2.84] (1.51, 4.86)	5/179 [0.88] (0.28, 2.04)
	6	25/89 [8.00] (5.18, 11.8)	24/61 [13.3] (8.50, 19.7)	6/89 [1.92] (0.70, 4.18)	4/61 [2.21] (0.60, 5.66)
	7	13/32 [14.6] (7.75, 24.9)	10/33 [9.57] (4.59, 17.6)	6/32 [6.72] (2.47, 14.6)	1/33 [0.96] (0.02, 5.33)
	8	3/11 [8.96] (1.85, 26.2)	1/4 [7.35] (0.19, 41.0)	2/11 [5.97] (0.72, 21.6)	0/4 [0.00] (0, 27.1)
	9	1/1 [103] (2.61, 575)	0/1 [0.00] (0, 221)	0/1 [0.00] (0, 381)	0/1 [0.00] (0, 221)
HR (95% CI)		1.20 (1.11, 1.30)	1.18 (1.09, 1.28)	1.36 (1.17, 1.58)	1.26 (1.00, 1.59)
		p<0.001	p<0.001	p<0.001	p=0.047
p-value for interaction		0.777		0.608	
C-index (95% CI)		0.57 (0.55-0.60)		0.68 (0.62-0.73)	
Goodness of fit test p-value		0.428		0.751	