Original article

Predictive Value of the CHA$_2$DS$_2$-VASc Score in Atrial Fibrillation Patients at High Risk for Stroke Despite Oral Anticoagulation

Eva Jover,[a] Vanessa Roldán,[b] Pilar Gallego,[b] Diana Hernández-Romero,[a] Mariano Valdés,[a] Vicente Vicente,[b] Gregory Y.H. Lip,[c] and Francisco Marín[a,*]

[a] Servicio de Cardiología, Hospital Universitario Virgen de la Arrixaca, Universidad de Murcia, El Palmar, Murcia, Spain
[b] Servicio de Hematología y Oncología Médica, Hospital Universitario Morales Meseguer, Universidad de Murcia, Murcia, Spain
[c] Haemostasis, Thrombosis and Vascular Biology Unit, University of Birmingham Centre for Cardiovascular Sciences, City Hospital, Birmingham, United Kingdom

A B S T R A C T

Introduction and objectives: The risk of stroke in atrial fibrillation is heterogeneous and depends upon underlying clinical conditions included in current risk stratification schemes. Recently, the CHA$_2$DS$_2$-VASc score has been included in guidelines to be more inclusive of common stroke risk factors seen in everyday clinical practice, and useful in defining "truly low risk" subjects. We aimed to assess the usefulness of CHA$_2$DS$_2$-VASc score to give us an additional prognostic perspective for adverse events and mortality among "real world" anticoagulated patients with atrial fibrillation who are often elderly with many comorbidities.
Methods: Consecutive outpatients with permanent/paroxysmal nonvalvular atrial fibrillation with CHA$_2$DS$_2$-VASc≥2 and stabilized oral anticoagulation (international normalized ratio 2.0-3.0) for at least the preceding 6 months were recruited. Patients with CHA$_2$DS$_2$-VASc≥2 were selected. Adverse cardiovascular events including stroke, acute coronary syndrome, or heart failure; major bleeds; and mortality were recorded during more than 2.5-year-follow-up.
Results: Of 933 patients (93.5%) assessed, 432 were males, median age 76 (71-81) years. After a follow-up of 946 (782-1068) days, 109 patients (11.7%) had adverse cardiovascular events, 80 patients (8.6%) had major bleeds, 101 patients (10.8%) died, and 230 (24.6%) major adverse events (composite end-point). Increasing CHA$_2$DS$_2$-VASc score by 1 point had a significant impact on the occurrence of cardiovascular events (hazard ratio=1.27; 95% confidence interval, 1.13-1.44; *P*<.001), mortality (hazard ratio=1.36; 95% confidence interval, 1.19-1.54, *P*<.001); and major adverse events (hazard ratio=1.23; 95% confidence interval, 1.13-1.34; *P*<.001). CHA$_2$DS$_2$-VASc score was not associated with major bleeding episodes.
Conclusions: Among high risk atrial fibrillation patients on oral anticoagulation, CHA$_2$DS$_2$-VASc successfully predicts cardiovascular events and mortality, but not major bleeds.

© 2012 Sociedad Española de Cardiología. Published by Elsevier España, S.L. All rights reserved.

Valor predictivo de la escala CHA$_2$DS$_2$-VASc en pacientes con fibrilación auricular de alto riesgo embólico en tratamiento anticoagulante

R E S U M E N

Introducción y objetivos: El riesgo de ictus en la fibrilación auricular es heterogéneo y depende de las características clínicas subyacentes que se incluyen en los actuales esquemas de estratificación del riesgo. La escala de riesgo CHA$_2$DS$_2$-VASc se ha incluido recientemente en las guías de práctica clínica para una valoración más completa de los factores de riesgo de ictus que son frecuentes en la práctica clínica diaria y para una definición útil de individuos con un riesgo «realmente bajo». Nuestro objetivo es evaluar la utilidad de la escala CHA$_2$DS$_2$-VASc para obtener una perspectiva pronóstica adicional respecto a los eventos adversos y la mortalidad en la práctica clínica real en pacientes con fibrilación auricular anticoagulados, que a menudo son ancianos con múltiples comorbilidades.
Métodos: Se reclutó a pacientes ambulatorios consecutivos con fibrilación auricular permanente/paroxística no valvular y puntuación CHA$_2$DS$_2$-VASc ≥ 2 y anticoagulación oral estabilizada (razón internacional normalizada: 2,0-3,0) durante al menos los 6 meses previos a su inclusión en el estudio. Se seleccionó a pacientes con CHA$_2$DS$_2$-VASc ≥ 2. Se registraron los eventos adversos cardiovasculares (incluidos ictus, síndrome coronario agudo e insuficiencia cardiaca), las hemorragias mayores y la mortalidad durante un seguimiento de más de 2,5 años.
Resultados: De los 933 pacientes (93,5%) evaluados, 432 eran varones, con una mediana de edad de 76 (71-81) años. Tras un seguimiento de 946 (782-1.068) días, 109 pacientes (11,7%) presentaron eventos adversos cardiovasculares, 80 (8,6%) sufrieron hemorragias mayores, 101 (10,8%) fallecieron y

* Corresponding author. Servicio de Cardiología, Hospital Universitario Virgen de la Arrixaca, Ctra. Madrid-Cartagena s/n, 30120 El Palmar, Murcia, Spain.
  E-mail address: fcomarino@hotmail.com (F. Marín).

230 (24,6%) sufrieron eventos adversos mayores (objetivo de valoración combinado). Un aumento de 1 punto en la escala CHA$_2$DS$_2$-VASc tuvo una repercusión significativa en la frecuencia de eventos cardiovasculares (*hazard ratio*=1,27; intervalo de confianza del 95%, 1,13-1,44; p<0,001), la mortalidad (*hazard ratio*=1,36; intervalo de confianza del 95%, 1,19-1,54; p<0,001) y los eventos adversos mayores (*hazard ratio*=1,23; intervalo de confianza del 95%, 1,13-1,34; p<0,001). La escala CHA$_2$DS$_2$-VASc no se asoció a los episodios de hemorragia mayor.

Conclusiones: En los pacientes con fibrilación auricular de alto riesgo tratados con anticoagulación oral, la escala de riesgo CHA$_2$DS$_2$-VASc predice satisfactoriamente los eventos cardiovasculares y la mortalidad, pero no las hemorragias mayores.

© 2012 Sociedad Española de Cardiología. Publicado por Elsevier España, S.L. Todos los derechos reservados.

Abbreviations

AF: atrial fibrillation
INR: international normalized ratio
MAE: major adverse events
OAC: oral anticoagulation

INTRODUCTION

Atrial fibrillation (AF) increases 5-fold the risk for stroke and thromboembolism.[1] Nonetheless, the stroke risk in AF patients is not homogeneous[2] but depends on the presence of other underlying clinical conditions.[3] These risk factors have been used to formulate stroke risk schemes that are used in clinical practice to stratify the embolic risk (low, moderate, or high) in AF and to choose proper antithrombotic agents, especially since until recently we only had an "inconvenient" anticoagulant, the vitamin K antagonist.[4,5] Oral anticoagulation (OAC) is highly effective in reducing stroke risk and mortality rates in patients with AF,[6] but also raises the risk for bleeds, at least in the historical trials.[7,8] More contemporary data show that the risk of major bleeding with acetylsalicylic acid may not be significantly different from OAC, especially in the elderly.[9–11]

Several risk stratification schemes have been derived from nonwarfarin arms of clinical cohort trials and/or expert consensus groups.[12] The most popular risk stratification scheme has been the CHADS$_2$ (congestive heart failure, hypertension, age, diabetes, stroke [doubled]) score[13] because it is easy to remember and calculate[4,5] and in some studies may have a better predictive value than other scores.[13] More recently, the value of the CHADS$_2$ scheme has been debated, given its noninclusion of many stroke risk factors and other limitations.[14,15] Thus, the CHADS$_2$ score has been refined with the CHA$_2$DS$_2$-VASc (congestive heart failure, hypertension, age≥75 [doubled], diabetes, stroke [doubled]-vascular disease and sex category [female]) emphasizing a risk factor-based approach.[12] Existing risk factors have been reclassified and new risk factors have been included (such as female sex and vascular disease).[12,16] The CHA$_2$DS$_2$-VASc consistently outperforms the CHADS$_2$ score in identifying low risk patients, and is as good as–and possibly better than–the CHADS$_2$ score in identifying those who develop stroke and thromboembolism.[12,17,18] Therefore, the European Society of Cardiology guidelines[4,5] encourage the use of CHADS$_2$ and CHA$_2$DS$_2$-VASc to refine stratification of patients and to aid decisions for thromboprophylaxis. A possible criticism has suggested that this risk score cannot give us more information after initiating OAC.[19] A recent Spanish study has even shown that the CHA$_2$DS$_2$-VASc risk stratification scheme better discriminated between patients at a low and intermediate risk of thromboembolic complications when compared to others.[20]

This study aims to assess the usefulness of CHA$_2$DS$_2$-VASc score to differentiate and predict adverse cardiovascular outcome and mortality among patients with AF on OAC. We aimed to assess the usefulness of CHA$_2$DS$_2$-VASc score to give us an additional prognostic perspective for adverse events and mortality among "real world" anticoagulated patients with AF who are often elderly patients with many comorbidities.

METHODS

Study Population

We recruited 998 consecutive outpatients diagnosed as having permanent or paroxysmal nonvalvular AF from our outpatient anticoagulation clinic. All patients received acenocoumarol OAC and had stabilized international normalized ratio values (INR 2.0-3.0) for at least the 6 months before study inclusion. The CHA$_2$DS$_2$-VASc score was calculated as previously described.[12] We selected those patients with a CHA$_2$DS$_2$-VASc score≥2 (high risk for stroke). For this reason, 65 (6.5%) patients were excluded. Finally 933 patients were included in the present study and followed for more than 2 years.

Inclusion criteria were an age older than 18 years, absence of any hematological disorder or contraindication for OAC in the last 6 months, absence of ischemic events (acute coronary syndrome, interventional procedures, stroke, or hemodynamic instability) requiring hospitalization at least for 6 months before a patient's enrollment, absence of rheumatic AF and prosthetic heart valves. Clinical and demographic characteristics as well as details from the antithrombotic therapies received/prescribed were recorded from their medical records (Table 1).

Follow-up was performed through visits in our outpatient anticoagulation clinic. During the study period, there were no changes in the anticoagulant drug class. Dental procedures were managed without retiring OAC. We detected 60 programmed surgeries with bridging therapy with low molecular heparins without adverse events associated to them. Adverse events were recorded, including thrombotic and cardiovascular events (such as stroke both ischaemic and embolic, acute coronary syndrome, acute heart failure), major bleeding events, and global mortality and cardiovascular death. Major bleeds were determined according to the 2005 International Society on Thrombosis and Haemostasis criteria.[21] Besides, "major adverse events" (MAE) were defined as a composite end-point of cardiovascular events, major bleeding, and mortality.

Statistical Analysis

Variables are presented as counts (percentages) or median [inter quartile range] as appropriate for categorical and continuous data, respectively. Kolmogorov-Smirnov test was used to check for normal distribution of continuous data. The clinical impact of

Table 1

Baseline Clinical Characteristics of Atrial Fibrillation Patients on Oral Anticoagulation

Patients	N=933
Male sex	432 (46)
Age, years	76 [71-81]
Age≥75 years	570 (61)
Hypertension	796 (85)
Diabetes mellitus	253 (27)
Hypercholesterolemia	298 (32)
Current tobacco smoking habit	127 (14)
Congestive heart failure	360 (39)
Prior stroke or TIA	190 (20)
Coronary artery disease	185 (20)
Peripheral vascular disease	87 (9)
CHA$_2$DS$_2$-VASc score	4 [3-5]
CHADS$_2$ score	2 [2-3]
Concomitant treatment	
Antiplatelet therapy	160 (17)
ACE inhibitors	246 (26)
Angiotensin-renin blockers	212 (23)
Calcium antagonist	209 (22)
Beta-blockers	285 (30)
Statins	199 (21)
Digoxin	177 (19)
Diuretics	402 (43)

ACE, angiotensin converting enzyme; TIA, transient ischemic attack.
Data are expressed as no. (%) or median [interquartile range].

Table 2

Total Event Rates per Year

End-points	no., %	Rate, %/year
Cardiovascular events	109 (11.7)	4.5
Stroke	38 (4.1)	1.6
ACS	41 (4.4)	1.7
Acute HF	31 (3.3)	1.3
Major bleeding	80 (8.6)	3.3
Intracranial	17 (1.8)	0.7
Global death	101 (10.8)	2.7
Cardiovascular death	30 (3.2)	1.2
Hemorrhagic cause	9 (0.9)	0.4
MAE	230 (24.6)	9.5

ACS, acute coronary syndrome; HF, heart failure; MAE, major adverse events.
Size of the whole sample assessed was of 933 atrial fibrillation patients on oral anticoagulation and at high risk for stroke (CHA$_2$DS$_2$-VASc≥2). Median [interquartile range] follow-up period was 946 [782-1068] days.

the calculated CHA$_2$DS$_2$-VASc was determined using Cox regression modeling with the score as the dependent variable. For all the investigated adverse events (cardiovascular events, major bleeding, mortality, and composite end-point) the percentage of event-rates per year after stratification of patients from 2 to 9 points (according to the CHA$_2$DS$_2$-VASc scoring system) were calculated, with hazard ratio (HR) obtained for 1 point of each increase in risk scoring from Cox regression modeling. The accuracy of prognostic value from CHA$_2$DS$_2$-VASc score was determined by calculating the area under the receiver-operator characteristic curve and the c-statistic value. The c-statistic quantifies and discriminates the ability (*P*-value≥.5), whereas HR quantifies the increased relative risk of adverse events across scores stratus. All *P*-values<.05 were accepted as statistically significant. Statistical analysis was performed using SPSS 15.0 for Windows (SPSS, Inc., Chicago, Illinois, United States).

RESULTS

Baseline clinical characteristics of the 933 (93.5%) patients included and assessed for CHA$_2$DS$_2$-VASc score≥2 and adverse events are shown in Table 1. The median age was 76 [71-81] years old, with 432 (46%) of them males. All patients assessed had CHA$_2$DS$_2$-VASc score≥2 and the median CHA$_2$DS$_2$-VASc score was 4 [3-5] and the median CHADS$_2$ score was 2 [2-3].

Median follow-up period was over 2.5 years (median 946 [782-1068] days). During this period, 109 patients (11.7%) presented with an adverse cardiovascular event, 80 patients (8.6%) had a major bleeding event, and 101 patients (10.8%) died; 30 (3.2%) of them died as a result of vascular death and 9 (0.9%) after a hemorrhagic event. As a composite end-point of cardiovascular

events, MAE major bleeding and mortality was observed in 230 patients (24.6%) (Table 2).

CHA$_2$DS$_2$-VASc Score and Adverse Events

In Table 3 and Figure we present the percentage of event rates per year according to the CHA$_2$DS$_2$-VASc score. We clearly show increasing event rates for 1 unit-increasing CHA$_2$DS$_2$-VASc score for cardiovascular events (Table 3A), major bleeding episodes (Table 3B), death rate (Table 3C), and MAE (Table 3D).

The CHA$_2$DS$_2$-VASc score had a c-statistic of 0.61 (95% confidence interval [95%CI], 0.59-0.66; *P*<.001) for cardiovascular events, while for mortality the c-statistic was 0.64 (95%CI, 0.58-0.70; *P*<.001), and for MAE, 0.61 (95%CI, 0.57-0.65; *P*<.001) (Table 4). The c-statistic for major bleeding episodes was not significant (0.54; 95%CI, 0.48-0.61; *P*=.179).

The increases in the CHA$_2$DS$_2$-VASc score showed a significant association with the development of clinical events, with the occurrence of cardiovascular events (HR=1.27; 95%CI, 1.13-1.44; *P*=.001), all-cause mortality (HR=1.36; 95%CI, 1.19-1.54; *P*<.001) and MAE (HR=1.23; 95%CI, 1.13-1.34; *P*<.001), Table 4. There was no significant association between CHA$_2$DS$_2$-VASc score and major bleeding episodes (HR=1.14; 95%CI, 0.98-1.32; *P*=.092).

DISCUSSION

The findings of the present study suggest that the CHA$_2$DS$_2$-VASc scoring system may be a useful tool to predict adverse events beyond thromboembolic risk in AF patients taking OAC. We found that one-unit-increasing CHA$_2$DS$_2$-VASc score[12] in high risk patients–which ranges from 2 to 9 points–was significantly associated with higher event rate, in particular cardiovascular events and mortality, despite all patients included taking OAC. There also was no statistically significant association between CHA$_2$DS$_2$-VASc score and major bleeding events.

We found that increasing scores across the CHA$_2$DS$_2$-VASc scoring strata–explored by 1-unit increments–consistently increased by 1.23 and 1.36-fold the risk (HR) to suffer any of the adverse events or mortality. Thus, subtype stratification into different high-risk categories derived from the calculation of CHA$_2$DS$_2$-VASc score may reflect the reality of risk for those AF patients at high risk on OAC. In a cohort study of 11 245 patients, Baruch et al.[22] concluded that high risk patients may be treated with more aggressive therapeutic strategies than those at

Table 3
Percentage of Event Rates per Year According the CHA$_2$DS$_2$-VASc Score

CHA$_2$DS$_2$-VASc score	% Event rate/year, %	No	Yes	Total
A. Cardiovascular events				
2	1.41	105	4	109
3	3.19	177	16	193
4	4.50	219	29	248
5	5.34	186	30	216
6	8.13	82	22	104
7	4.37	39	5	44
8	7.22	13	3	16
9	0	3	0	3
Total		824	109	933
B. Major bleeding rate				
2	2.11	103	6	109
3	2.79	179	14	193
4	3.72	224	24	248
5	3.38	197	19	216
6	3.69	94	10	104
7	4.37	39	5	44
8	4.80	14	2	16
9	0	3	0	3
Total		853	80	933
C. Death rate				
2	2.12	103	6	109
3	2.19	182	11	193
4	3.57	225	23	248
5	4.62	190	26	216
6	9.24	79	25	104
7	4.37	39	5	44
8	9.61	12	4	16
9	12.82	2	1	3
Total		832	101	933
D. Major adverse event rate				
2	4.23	97	12	109
3	6.97	158	35	193
4	12.55	185	63	248
5	10.68	156	60	216
6	14.05	66	38	104
7	13.11	29	15	44
8	14.42	10	6	16
9	12.82	2	1	3
Total		703	230	933
E. Stroke rate				
2	0	109	0	109
3	1.20	187	6	193
4	1.92	237	11	248
5	1.60	207	9	216
6	3.70	94	10	104
7	0.87	43	1	44
8	2.40	15	1	16
9	0	3	0	3
Total		895	38	985

accurate thromboprophylaxis decisions for the management of those "classical high risk patients" depending on their categorization into a "high risk subtype stratum."

AF patients are at high risk for both cardiovascular and bleeding events.[23] Notably, a great number of risk factors included in the CHADS$_2$ score are also bleeding risk factors,[24] ie, prior stroke, elderly, renal impairment or hypertension,[25] assessed by the popular HAS-BLED score.[26] It means that as the risk for stroke and thromboembolism increases—measured by, for example, the CHADS$_2$ score—the bleeding risk also increases.[27,28] With the novel OAC agents, the move has been to be more inclusive, rather than exclusive, of stroke risk factors.[29] Thus, the CHA$_2$DS$_2$-VASc includes newer risk factors and refines point assignment to others, and in several independent cohorts, the ability of the CHA$_2$DS$_2$-VASc score to predict or assess the impact in the occurrence of adverse events has been compared with other current risk stratification schemes, whereby CHA$_2$DS$_2$-VASc consistently better identifies patients truly at low to moderate risk for stroke and thromboembolism and is as good–and possibly better–at identifying "high" risk for thromboembolism.[12,17,30–32] We recently showed how HAS-BLED score may give important prognostic information regarding death and cardiovascular events, and not only bleeding risk.[33] However, we were not able to demonstrate a significant predictive role of CHA$_2$DS$_2$-VASc score regarding bleeding risk in the present cohort. The median HAS-BLED score in our population was 2 [2-3]. It may explain, at least in part, the lower bleeding risk in our population. We have recently demonstrated in a population on acenocoumarol OAC that bleeding rates only exceeded thrombotic events at HAS-BLED score≥3[33] as previously demonstrated.[27] Moreover, acenocoumarol, given its pharmacokinetic features[34] which may increase the risk of having INR>6, must be the better recommendation for patients at "low hemorrhage risk" to achieve OAC into during time in therapeutic range (TTR). Noteworthy, although a lesser number of studies have compared therapeutic effects of acenocoumarol vs warfarin, acenocoumarol appears to lead to less stable TTR,[34,35] a disadvantageous effect of acenocoumarol therapy which is not found in our selected population. Thereby, the relative low bleeding risk and acenocoumarol based-on anticoagulation, together with the higher TTR at entry of our cohort, may result in a more stable population with reduced thrombotic and hemorrhagic risk. It may explain the modest predictive value of CHA$_2$DS$_2$-VASc reported in our study. Accordingly, future investigational research should explore the clinical impact and predictive value of CHA$_2$DS$_2$-VASc score in those patients at 60% to 65% TTR on acenocoumarol-based anticoagulation (as warfarin-based populations finding consistent predictive value for thrombotic and hemorrhagic events after assessing the CHADS-VASc score) and/or HAS-BLED score≥3.

Most of the current risk stratification schemes are derived from nonwarfarin arms of historical clinical trial cohorts (which randomized <10% of subjects screened), in which the risk factors are often inadequately defined or recorded. Moreover, their predictive ability in patients receiving OAC is lesser validated.[36] A few validation studies—some recent meta-analyses—have not been based on clinical trial cohorts, have applied the published schemes to unselected patients encountered in general clinical practice to compare their predictive value[12,17,31,32] and in some cases their published results were performed in selected patients without indication for OAC,[30] unlike our study.

moderate risk. Other authors have previously evaluated the risk for stroke/TE in individual AF patients according to their underlying clinical conditions,[7] for example, to target their optimal INR and improve thromboprophylaxis decisions, but results were unsuccessful. This perhaps needs to be investigated in order to aid more
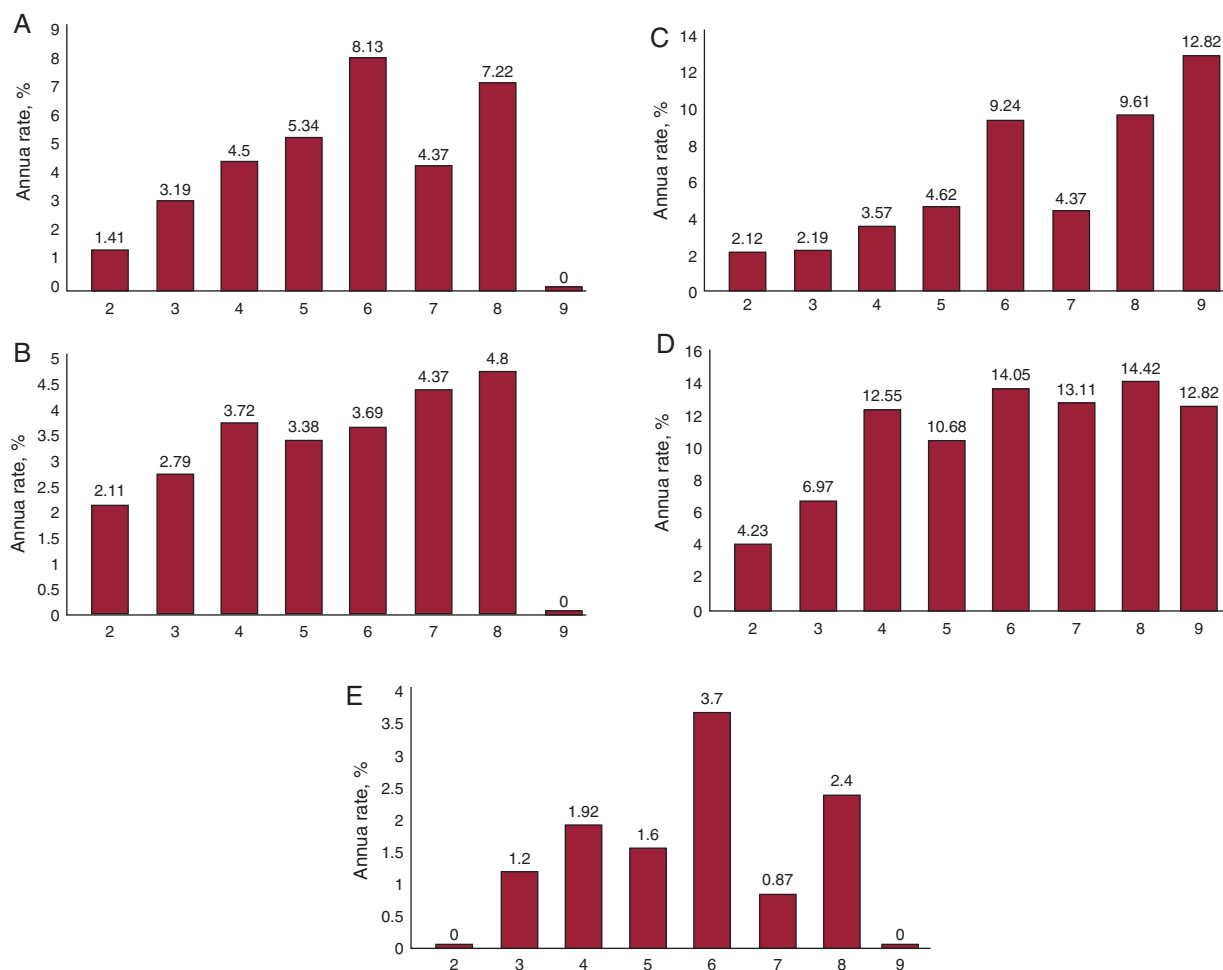
Limitations

We included only patients under steady oral OAC to homogenize the cohort, and so other potential variables were excluded.

Figure. A: cardiovascular events according to CHA$_2$DS$_2$-VASc (annual rate). B: haemorrhagic events according to CHA$_2$DS$_2$-VASc score (annual rate). C: mortality according to CHA$_2$DS$_2$-VASc score (annual rate). D: major adverse events according to CHA$_2$DS$_2$-VASc score (annual rate). E: stroke according to CHA$_2$DS$_2$-VASc score (annual rate).

We have recruited a population with good anticoagulation control at entry, while other clinical cohort studies recruited only patients with TTRs of 60% to 75%. Therefore, our results may not be applicable to unstable anticoagulation patients (with low TTRs) who are more prone to suffer adverse events or to patients under early OAC who are more likely to have thrombotic events[27,37]. Our patients were only anticoagulated with acenocoumarol (the vitamin K antagonist most widely used in Spain) which differs from warfarin in its shorter half-life; that seems to have some advantages in clinical practice. We have found a modest predictive value of the CHA$_2$DS$_2$-VASc score (<70%), unlike available data from previous reports. The good OAC at entry, the use of acenocoumarol, and thus the more stable population assessed in our study might explain the modest c-statistic reported. Moreover, the exponential increase in stroke rate, previously reported,[13] is blunted at higher scores probably due to the reduced number of patients in our study having high risk of stroke. It may be a limitation to achieving statistical differences.

Table 4

Predictive Value and Clinical Impact of Increasing CHA$_2$DS$_2$-VASc Score in End-Point Occurrence: C Statistic Indices and Hazard Ratios by Cox Regression Analysis

End-point	CHA$_2$DS$_2$-VASc			
	Predictive value c-statistic (95%CI)	P-value	Cox analysis HR (95CI%)	P-value
Cardiovascular events	0.61 (0.59-0.66)	<.001	1.27 (1.13-1.44)	.001
Major bleeding events	0.54 (0.48-0.61)	.179	1.14 (0.98-1.32)	.092
Mortality	0.64 (0.58-0.70)	<.001	1.36 (1.19-1.54)	<.001
MAE	0.61 (0.57-0.65)	<.001	1.23 (1.13-1.34)	<.001

95%CI, 95% confidence interval; HR, hazard ratio; MAE, major adverse events (composite end-point including cardiovascular events, major bleeding and mortality).
Increasing CHA$_2$DS$_2$-VASc and CHADS$_2$ scores mean an increase in one unit of the each risk stratification scores.
All P-values<.05 were considered significant.

CHA_2DS_2-VASc is a refinement of the $CHADS_2$ score and offers consistently better discrimination of patients at low and moderate risk,[18] and is as good—and possibly better—at identifying patients at high risk of developing thromboembolic events. Hence, the exponential increasing stroke risk by $CHADS_2$ may be labile when assessed by CHA_2DS_2-VASc due to the more intense stratification risk into a higher number of high risk categories. We have assessed Caucasian-based populations without any prevalence of other races, thus our results might be specific to our patient population and the way they were managed.

CONCLUSIONS

In conclusion, the CHA_2DS_2-VASc score successfully predicts cardiovascular events and mortality, but not major bleeds, among high risk AF patients on OAC.

ACKNOWLEDGEMENTS

CONFLICTS OF INTEREST

Gregory Y.H. Lip has served as a consultant for Bayer, Astellas, Merck, AstraZeneca, Sanofi, BMS/Pfizer, Biotronik, Portola, and Boehringer Ingelheim and has been on the speakers bureau for Bayer, BMS/Pfizer, Boehringer Ingelheim, and Sanofi-Aventis. Francisco Marín has served as a consultant for Bayer and Boehringer Ingelheim, and has been on the speakers bureau for Boehringer Ingelheim and Boston Scientific.

REFERENCES

1. Wolf PA, Abbott RD, Kannel WB. Atrial fibrillation as an independent risk factor for stroke: the Framingham Study. Stroke. 1991;22:983–8.
2. Lip GYH, Lim HS. Atrial fibrillation and stroke prevention. Lancet Neurol. 2007;6:981–93.
3. Watson T, Arya A, Sulke N, Lip GY. Relationship of indices of inflammation and thrombogenesis to arrhythmia burden in paroxysmal atrial fibrillation. Chest. 2010;137:869–76.
4. Camm AJ, Kirchhof P, Lip GYH, Schotten U, Savelieva I, Ernst S, et al. Guidelines for the management of atrial fibrillation: the Task Force for the Management of Atrial Fibrillation of the European Society of Cardiology (ESC). Eur Heart J. 2010;31:2369–3429.
5. Camm AJ, Kirchhof P, Lip GYH, Schotten U, Savelieva I, Ernst S, et al.; Grupo de Trabajo para el Manejo de la Fibrilación Auricular de la Sociedad Europea de Cardiología (ESC). Guías de práctica clínica para el manejo de la fibrilación auricular–2.ª edición corregida 8 de abril de 2011. Rev Esp Cardiol. 2010;63:1483. e1–83.
6. Hart RG, Pearce LA, Aguilar MI. Meta-analysis: antithrombotic therapy to prevent stroke in patients who have nonvalvular atrial fibrillation. Ann Intern Med. 2007;146:857–67.
7. Singer DE, Chang Y, Fang MC, Borowsky LH, Pomernacki NK, Udaltsova N, et al. Should patient characteristics influence target anticoagulation intensity for stroke prevention in nonvalvular atrial fibrillation?. The ATRIA study. Circ Cardiovasc Qual Outcomes. 2009;2:297–304.
8. Gorin L, Fauchier L, Nonin E, De Labriolle A, Haguenoer K, Cosnay P, et al. Antithrombotic treatment and the risk of death and stroke in patients with atrial fibrillation and a CHADS2 score=1. Thromb Haemost. 2010;103:833–40.
9. Mant J, Hobbs FD, Fletcher K, Roalfe A, Fitzmaurice D, Lip GY, et al.; BAFTA investigators; Midland Research Practices Network (MidReC). Warfarin versus aspirin for stroke prevention in an elderly community population with atrial fibrillation (the Birmingham Atrial Fibrillation Treatment of the Aged Study, BAFTA): a randomised controlled trial. Lancet. 2007;493–503. 370.
10. Van Walraven C, Hart RG, Connolly S, Austin PC, Mant J, Hobbs FD, et al. Effect of age on stroke prevention therapy in patients with atrial fibrillation: the atrial fibrillation investigators. Stroke. 2009;40:1410–6.
11. Connolly SJ, Eikelboom J, Joyner C, Diener HC, Hart R, Golitsyn S, et al.; AVERROES Steering Committee and Investigators. Apixaban in patients with atrial fibrillation. N Engl J Med. 2011;364:806–17.
12. Lip GY, Nieuwlaat R, Pisters R, Lane D, Crijns H. Refining clinical risk stratification for predicting stroke in atrial fibrillation using a novel risk factor based approach. The Euro Heart Survey on Atrial Fibrillation. Chest. 2010;137: 263–72.
13. Gage BF, Waterman AD, Shannon W, Boechler M, Rich MW, Radford MJ. Validation of clinical classification schemes for predicting stroke: results from the National Registry of Atrial Fibrillation. JAMA. 2001;285:2864–70.
14. Keogh C, Wallace E, Dillon C, Dimitrov BD, Fahey T. Validation of the CHADS2 clinical prediction rule to predict ischaemic stroke. A systematic review and meta-analysis. Thromb Haemost. 2011;106:528–38.
15. Karthikeyan G, Eikelboom JW. The CHADS2 score for stroke risk stratification in atrial fibrillation—friend or foe? Thromb Haemost. 2010;104:45–8.
16. Newman AB, Siscovick DS, Manolio TA, Polak J, Fried LP, Borhani NO, et al. Ankle-arm index as a marker of atherosclerosis in the Cardiovascular Health Study. Circulation. 1993;88:837–45.
17. Lip GY, Frison L, Halperin JL, Lane DA. Identifying patients at high risk for stroke despite anticoagulation: a comparison of contemporary stroke risk stratification schemes in an anticoagulated atrial fibrillation cohort. Stroke. 2010;41: 2731–8.
18. Azoulay L, Simon T, Dell'Aniello S, Renoux C, Suissa S. Comparison of the CHADS2 and CHA2DS2-VASc scores in predicting stroke events in patients with atrial fibrillation. Circulation. 2010;122:A18044.
19. Anguita M, Worner F, Domenech P, Marín F, Ortigosa J, Pérez-Villacastín J, et al. Nuevas evidencias, nuevas controversias: análisis crítico de la guía de práctica clínica sobre fibrilación auricular 2010 de la Sociedad Europea de Cardiología. Rev Esp Cardiol. 2012;65:7–13.
20. Abu-Assi E, Otero-Raviña F, Allut Vidal G, Coutado Méndez A, Vaamonde Mosquera L, Sánchez Loureiro M, et al.; on behalf of Grupo Barbanza researchers. Comparison of the reliability and validity of four contemporary risk stratification schemes to predict thromboembolism in non-anticoagulated patients with atrial fibrillation. Int J Cardiol. 2011. http://dx.doi.org/10.1016/j.ijcard.2011.10.096.
21. Schulman S, Kearon C; Subcommittee on Control of Anticoagulation of the Scientific and Standardization Committee of the International Society on Thrombosis and Haemostasis. Definition of major bleeding in clinical investigations of antihemostatic medicinal products in non-surgical patients. J Thromb Haemost. 2005;3:692–4.
22. Baruch L, Gage BF, Horrow J, Juul-Möller S, Labovitz A, Persson M, et al. Can patients at elevated risk of stroke treated with anticoagulants be further risk stratified? Stroke. 2007;38:2459–63.
23. Roldán V, Marín F, Muiña B, Torregrosa JM, Hernández-Romero D, Valdés M, et al. Plasma von Willebrand factor levels are an independent risk factor for adverse events including mortality and major bleeding in anticoagulated atrial fibrillation patients. J Am Coll Cardiol. 2011;57:2496–504.
24. Lip GY, Andreotti F, Fauchier L, Huber K, Hylek E, Knight E, et al. Bleeding risk assessment and management in atrial fibrillation patients Executive Summary of a Position Document from the European Heart Rhythm Association [EHRA], endorsed by the European Society of Cardiology [ESC] Working Group on Thrombosis. Thromb Haemost. 2011;106:997–1011.
25. Hughes M, Lip GY. Guideline Development Group for the NICE national clinical guideline for management of atrial fibrillation in primary and secondary care Risk factors for anticoagulation-related bleeding complications in patients with atrial fibrillation: a systematic review. QJM. 2007;100:599–607.
26. Pisters R, Lane DA, Nieuwlaat R, De Vos CB, Crijns HJ, Lip GY. A novel user-friendly score (HAS-BLED) to assess 1-year risk of major bleeding in patients with atrial fibrillation: The Euro Heart Survey. Chest. 2010;138:1093–100.
27. Hylek EM, Evans-Molina C, Shea C, Henault LE, Regan S. Major hemorrhage and tolerability of warfarin in the first year of therapy among elderly patients with atrial fibrillation. Circulation. 2007;115:2689–96.
28. Poli D, Antonucci E, Grifoni E, Abbate R, Gensini GF, Prisco D. Bleeding risk during oral anticoagulation in atrial fibrillation patients older than 80 years. J Am Coll Cardiol. 2009;54:999–1002.
29. Rasmussen LH, Larsen TB, Due KM, Tjønneland A, Overvad K, Lip GY. Impact of vascular disease in predicting stroke and death in patients with atrial fibrillation: the Danish Diet Cancer and Health cohort study. J Thromb Haemost. 2011;9:1301–7.
30. Olesen JB, Fauchier L, Lane DA, Taillandier S, Lip GY. Risk factors for stroke and thromboembolism in relation to age amongst patients with atrial fibrillation: The Loire Valley Atrial Fibrillation Project. Chest. 2012;141:147–53.
31. Olesen JB, Lip GY, Hansen ML, Hansen PR, Tolstrup JS, Lindhardsen J, et al. Validation of risk stratification schemes for predicting stroke and thromboembolism in patients with atrial fibrillation: nationwide cohort study. BMJ. 2011;342:d124.
32. Lin LY, Lee CH, Yu CC, Tsai CT, Lai LP, Hwang JJ, et al. Risk factors and incidence of ischemic stroke in Taiwanese with nonvalvular atrial fibrillation — a nation wide database analysis. Atherosclerosis. 2011;217:292–5.
33. Gallego P, Roldán V, Torregrosa JM, Gálvez J, Valdés M, Vicente V, et al. Relation of the HAS-BLED bleeding risk score to major bleeding, cardiovascular events and mortality in anticoagulated patients with atrial fibrillation. Circ Arrhythm Electrophysiol. 2012;5:312–8.

34. Oliva E, Galán P, Pacheco AM. Comparación de la calidad y el riesgo hemorrágico del tratamiento anticoagulante oral con acenocumarol frente a warfarina. Med Clin (Barc). 2008;131:96–7.

35. Pattacini C, Manotti C, Pini M, Quintavalla R, Dettori AG. A comparative study on the quality of oral anticoagulant therapy (warfarin versus acenocoumarol). Thromb Haemost. 1994;71:188–91.

36. Poli D, Antonucci E, Grifoni E, Abbate R, Gensini GF, Prisco D. Stroke risk in atrial fibrillation patients on warfarin Predictive ability of risk stratification schemes for primary and secondary prevention. Thromb Haemost. 2009;101:367–72.

37. Wan Y, Heneghan C, Perera R, Roberts N, Hollowell J, Glasziou P, et al. Anticoagulation control and prediction of adverse events in patients with atrial fibrillation: a systematic review. Circ Cardiovasc Qual Outcomes. 2008;1:84–91.