Joint Mexican position document on the treatment of atrial fibrillation

Posicionamiento conjunto acerca del tratamiento para fibrilación auricular

Gerardo Rodríguez-Diez1*, Manlio F. Márquez2, Pedro Iturralde-Torres2, Luis G. Molina-Fernández de L.3, Gerardo Pozas-Garza4, Alejandro Cordero-Cabra5, and Ulises Rojel-Martínez6

1Arrhythmia and Cardiac Pacing Unit, Centro Médico Nacional Siglo XXI, Instituto de Seguridad Social y Servicios Sociales de los Trabajadores del Estado, Mexico City, 2Electrocardiology Service, Instituto Nacional de Cardiología Ignacio Chávez, Mexico City, 3Electrophysiology Unit, Hospital General de México, Mexico City, 4Instituto de Cardiología y Medicina Vascular, Hospital Zambrano Hellion TEC-Salud, Nuevo Léon; 5Servicio de Electrophisiología, Centro Médico Nacional de Occidente-Hospital de Especialidad, Instituto de Seguridad Social y Servicios Sociales de los Trabajadores del Estado (ISSSTE), Guadalajara; 6Arrhythmia and Cardiac Pacing Unit, Centro Médico Sur de los Servicios Médicos de Salud de Puebla, Puebla, Mexico

Abstract

Atrial fibrillation (AF) is a frequent arrhythmia; its prevalence is near 2% in the general population; in Mexico, more than one-half million people are affected. AF needs to be considered as a public health problem. Because AF is an independent risk factor associated with mortality, due to embolic events, heart failure, or sudden death; early diagnosis is of utmost importance. In unstable patients with a recent onset of AF, electrical cardioversion should be practiced. In stable patients, once thromboembolic measures have been taken, it is necessary to assess whether it is reasonable to administer an antiarrhythmic drug to restore sinus rhythm or performed electrical cardioversion. For recidivating cases of paroxysmal and persistent presentation, the most effective strategy is performed pulmonary vein isolation with either radiofrequency or cryoballoon energy. Permanent AF is that in which recovery of sinus rhythm is not possible, the distinguishing feature of this phase is the uncontrollable variability of the ventricular frequency and could be treated pharmacologically with atrioventricular (AV) nodal blockers or with a VVIR pacemaker plus AV nodal ablation. The presence of AF has long been associated with the development of cerebral and systemic (pulmonary, limb, coronary, renal, and visceral) embolism. The prevention of embolisms in "valvular" AF should perform with Vitamin K antagonists (VKA). For patients with AF not associated with mitral stenosis or a mechanical valve prosthesis, a choice can be made between anticoagulant drugs, VKA, or direct oral anticoagulants. Antiplatelet agents have the weakest effect in preventing embolism.

Key words: Atrial fibrillation. Drug treatment. Tromboprofilaxis. Cryoballoon ablation. Radiofrequency ablation.

Resumen

La fibrilación auricular (FA) es una arritmia frecuente; su prevalencia es cercana al 2% en la población general, en México se ven afectados más de medio millón de personas por eso debe considerarse como un problema de salud pública. Debido a que la FA es un factor de riesgo independiente asociado a mortalidad, por eventos embólicos, insuficiencia cardíaca o
muerte súbita, la identificación y diagnóstico temprano es de suma importancia. En el inicio reciente de FA en pacientes inestables, se debe practicar la cardioversión eléctrica. En pacientes estables, una vez que se han tomado medidas tromboembólicas, es necesario evaluar si es razonable administrar un medicamento antiarrítmico para restaurar el ritmo sinusal o realizar una cardioversión eléctrica. Para los casos que recibivan, ya sea paroxística o persistente, la estrategia más efectiva es realizar el aislamiento de la venas pulmonares con radiofrecuencia o crioablación con balón. La FA permanente es aquella en la que no es posible la recuperación del ritmo sinusal, la característica distintiva de esta fase de la FA es la variabilidad incontrolable de la frecuencia ventricular. Puede tratarse farmacológicamente con bloqueadores nódales AV o con un marcapasos VVIR mas ablación del nodo AV. La presencia de FA se ha asociado durante mucho tiempo con el desarrollo de embolia cerebral y sistémica (pulmonar, de extremidades, coronaria, renal y visceral). La prevención de embolias en la FA “valvular” debe realizarse con antagonistas de la vitamina K (AVK). Para los pacientes con FA no asociados con estenosis mitral o una prótesis valvular mecánica, se puede elegir entre medicamentos anticoagulantes, AVK o anticoagu- gulantes orales directos (DOAC). Los agentes antiplaquetarios tienen el efecto más débil para prevenir la embolia.

**Palabras clave:** Fibrilación atrial. Tratamiento farmacológico. Tromboprofilaxis. Crioablación. Ablación con radiofrecuencia.

**What is known about the epidemiology of atrial fibrillation (AF) in Mexico? Can it be considered a public health problem?**

AF incidence and prevalence increase with age. Its prevalence is near to 2% in the general population, but it could be as high as 10% in those over 75 years1,2. Before the Mexican Registry of AF (ReMeFA) was published a study was conducted in Mexico in 2007, finding that, for a population of 105,338,982 people the prevalence of cardiac arrhythmias was 2.4%, with tachyarrhythmias being the most common with 56% (1,402,453 people), of which, AF was the most frequent arrhythmia, occupying 60.7% of tachycardia (or a total of 851,489 cases)3.

Today, we estimate that in Mexico, there are more than one and a half million people with AF, with a prevalence ranging from 0.43% in the 40 to 49 age group to 8.48% in those over 80 years old, for an average of 1.58% in a population over 40 years of age. Permanent or chronic AF represents 51.5% (corresponding to 438,134, Mexicans). The ReMeFA study was the first national multicenter registry, with clinical follow-up of 1 year, in 1201 subjects, on the comparison of AF treatment with a rhythm control strategy or with rate control. This study was carried out with the collaboration of 71 cardiologists and electrophysiologists. At 1 year follow-up, an incidence of 3% of ischemic cerebral vascular disease (CVD) was observed in the rate control strategy; significantly higher than 1% in the rhythm control strategy (p = 0.04)4. Worldwide, CVD is the second leading cause of death and the leading cause of disability. CVD has become a health problem as a result of increased life expectancy and lifestyle changes, representing one of the leading causes of death in Mexico5. According to the Brain Attack Surveillance project in Durango, it is estimated that in Mexico the annual incidence of CVD is 232.3 cases per 100,000 inhabitants over 35 years of age, while its prevalence is eight cases of CVD per 1000 inhabitants, a figure that increases to 18 cases per 1000 in people over 65 years of age. It is important to note that in recent years, CVD has occurred in younger people as a result of the continuing increase in risk factors, including unhealthy lifestyles and obesity. In a Pan American Health Organization report, indicators of premature vascular mortality (in people under 70) showed that in Mexico the rate in nondiabetics was 10.7/100,000, compared to 3.3 and 5/100,000 in Canada and the USA, respectively6. Based on these results, we consider AF to be the most frequent tachyarrhythmia in Mexico with a high percentage of CVD, so it should be considered a public health problem in Mexico.

**Importance of early diagnosis**

AF is an independent risk factor associated with mortality, increasing it twice in men and 1.5 times in women; mortality due to embolic events can decrease with oral anticoagulation but other causes of cardiovascular death such as heart failure or sudden death continue to be frequent despite adequate treatment that is the reason why an early diagnosis is of utmost importance since AF can be asymptomatic (silent AF), and patients have it inadvertently, delaying proper treatment. The diagnosis of AF requires to be an event lasting at least 30 s and to be observed on an electrocardiogram (ECG), rhythm strip, or cardiac monitor, characteristi- cally with the irregularity of RR intervals without clearly identifiable P waves or with visible “f” waves of fibrillation. An early electrocardiographic recording is cost effective for documenting chronic forms of AF, particularly in populations older than 65 years with a
prevalence of up to 2.3%, obtaining a “necessary to treat” number of 70 to find one with AF\textsuperscript{1}. As for paroxysmal AF, the longer the record, the more likely it is to find silent events. Now the technology has evolved, so in Mexico, we already have 48-h recorders and implantable loop recorders whose duration is up to 3 years. The more we use these devices in high-risk patients, more likely the chance of found AF and being able to start appropriate treatment earlier\textsuperscript{4}.

**Antiarrhythmics available in Mexico for rhythm control: how and when?**

### Recent onset AF: conversion to sinus rhythm in an unstable patient

If the AF paroxysm is associated with “angina pectoris,” pulmonary edema, low blood pressure or shock, urgent electrical cardioversion should be practiced. It is recommended that the shock should be with the highest available energy 200 J biphasic or 360 J monophasic. It is not suggested to proceed in stages by increasing from lower energies. The reason for this is to reduce the number of shocks, use a lower cumulative dose of energy, and reduce the anesthetic time. For thromboembolic prophylaxis, unfractionated heparin (bolus according to body weight followed by infusion) should be administered, followed by oral anticoagulation\textsuperscript{1}. Although embolism risk might be increased because of the emergency nature of the condition.

### Stable patient

Assuming that the corresponding thromboembolic prevention measures have been taken and that the heart rate controlled with the isolated or combined use of beta-blockers, calcium antagonists, or digital, the clinician should assess whether it is reasonable to administer any antiarrhythmic drug to restore sinus rhythm. It is known that up to 50% of AF paroxysms may spontaneously remit within 24-48 h\textsuperscript{7}. If AF persists after this period, pharmacological cardioversion with amiodarone (oral or preferably intravenous), propafenone or flecainide is indicated. Intravenous amiodarone is given at a loading dose of 5-7 mg/kg in 30-60 min, followed by a maintenance dose of 1.2-1.8 g/day until 10 g\textsuperscript{1}. Completed. The oral dose of propafenone is 600 mg in a single dose and that of flecainide is 300 mg in a single dose. Sinus rhythm conversion occurs in 80-90% of cases within the first few hours\textsuperscript{8}. It should emphasize that sotalol, dronedarone, and digital are not indicated for conversion to sinus rhythm. If the episode becomes persistent despite the use of antiarrhythmics, electrical CV is indicated, preceded by a transesophageal echocardiogram to rule out intracavitary thrombus\textsuperscript{1,9}.

### Maintaining sinus rhythm

Once the conversion to sinus rhythm has achieved, the clinician should assess whether it is appropriate to use an antiarrhythmic daily for the maintenance of sinus rhythm or whether it is preferable not to give preventive antiarrhythmic and choose a strategy of treating the episode with the “pill in your pocket” strategy\textsuperscript{1,10}. For the maintenance of sinus rhythm, it is indicated to use one of the following antiarrhythmics: propafenone, flecainide, sotalol, dronedarone, or amiodarone. In the absence of structural heart disease, the use of propafenone or flecainide is recommended\textsuperscript{10}. Sotalol may use in the presence of ischemic heart disease. Dronedarone is indicated only for cases of paroxysmal AF with preserved left ventricular ejection fraction, in the absence of heart disease and with preserved systolic ventricular function. Amiodarone is considered a second-line drug due to its side effects; however, it is the most effective alternative for maintaining sinus rhythm\textsuperscript{1}. In the case of heart failure, the use of amiodarone is recommended. For the last three drugs (sotalol, dronedarone, and amiodarone), the duration of the QT interval should be monitored\textsuperscript{11}. A single dose of 600 mg propafenone or 300 mg flecainide is recommended for the “pill in your pocket” strategy\textsuperscript{1,8,10}. Sotalol, dronedarone, and amiodarone are contraindicated if the QT interval is prolonged\textsuperscript{11}. Caution should be exercised due to the possibility that these two drugs may unmask the electrocardiographic signs of Brugada syndrome or convert AF into an atrial flutter with a paradoxical increase in ventricular response (< 1% of cases)\textsuperscript{12}.

### Recurrent AF (paroxysmal and persistent)

Unlike the first episode approach (or very sporadic recurrent cases), for recidivating cases of paroxysmal and persistent presentation, it is indicated to use antiarrhythmics for prevention. The therapeutic options are propafenone, flecainide, sotalol, dronedarone, and amiodarone. It should emphasize that dronedarone is only indicated to prevent recurrence of paroxysmal or persistent AF that have lasted < 6 months of evolution, in the absence of heart disease and with preserved left ventricular function. AF ablation (radiofrequency or cryoballoon energy) should be considered as a first-line alternative for drug-refractory or symptomatic cases (at least one antiarrhythmic Class Ic or III)\textsuperscript{1,3}.
**Persistent AF lasting more than a year**

This category was established to identify patients who may benefit from a rhythm control strategy because AF is permanent of those with a chance to convert to sinus rhythm. There are two therapeutic options: (1) facilitated electrical cardioversion with prior use of antiarrhythmics\(^\text{14}\) and (2) AF ablation\(^\text{1}\). It is reasonable to proceed with facilitated electrical cardioversion with antiarrhythmic drugs as the first measure because if successful, although with early relapse, it demonstrates that the patient can maintain sinus rhythm and would be a suitable candidate for catheter ablation\(^\text{1,13,14}\).

**Immediate post-cardioversion recurrence**

Electrical cardioversion is one of the cornerstones for rhythm control in AF. However, immediate recurrence or therapeutic failure, described in up to 26% of cases, limits its clinical application\(^\text{15}\). To increase the response rate, antiarrhythmics must give before the electric shock\(^\text{13,14}\). The use of verapamil, amiodarone, or sotalol has been reported to decrease the incidence of immediate recurrence\(^\text{13-16}\). Other drugs such as ibutilide (not widely available in Mexico), vernakalant (not available in Mexico), and ranolazine (available in our country) have also shown benefit in this area\(^\text{1,17}\).

**Postponed cardioversion (facilitated by antiarrhythmic)**

It is indicated for persistent AF, mainly when the temporal progression is unknown or when a high probability of immediate recurrence is assumed. Amiodarone 600 mg/day administered for 1 month (total dose 16.8 g) is indicated for a better outcome. Pharmacological cardioversion has been observed to occur during loading in 16-18% of cases\(^\text{1}\). The success of electrical cardioversion is 88%. Besides, the ventricular response of the heart rate during AF is reduced from 100 ± 25 to 87 ± 27.5 beats/min (p ≤ 0.001) by a negative dromotropic effect on the atrioventricular (AV) node\(^\text{18}\).

**How to manage rate control in permanent AF? What is the role of AV node ablation with pacemaker implant?**

Much has been said about (AF) that can be summed up in three brief sentences: it is the most common arrhythmia, the easiest to diagnose, and the most difficult to treat\(^\text{19-21}\). Another no less ominous peculiarity is that AF is a progressive disease\(^\text{22}\) and that itself is a condition that contributes to its perpetuation\(^\text{23}\). In other words, the sooner we try to revert and achieve sinus rhythm, the higher the chances of success (to keep the patient in sinus rhythm)\(^\text{24}\).

**Permanent (chronic) AF**

Permanent AF is the one in which recovery of sinus rhythm is not possible\(^\text{1,19}\). The distinguishing feature of this phase of AF is the uncontrollable variability of the ventricular rate. It depends on the AV conduction and not on the sinus node function; it is the autonomic nervous system – sympathetic and vagal – that determines the AV conduction velocity and thus the ventricular frequency\(^\text{25}\). It is common to consider ventricular rate analysis only with a resting EKG record, however, this is not quite right because of the circadian heart rate variations. On the other hand, vagal tone during the early morning hours can delay AV conduction and cause considerable and sufficient ventricular pauses to cause low brain perfusion with its consequences. The therapeutic possibilities are: pharmacological and interventional\(^\text{26,27}\).

**Pharmacological treatment**

The main limitation of drugs is because they slow nodal conduction it can produce very severe bradycardia, without avoiding abnormally fast frequencies\(^\text{28}\). Antiarrhythmic drugs such as amiodarone are ineffective, as, by definition, sinus rhythm is not intended to be restored\(^\text{29}\). Beta-adrenergic blockers may delay AV conduction, but decrease the force of ventricular contraction\(^\text{1}\).

**Interventional treatment**

Once it has been demonstrated that the patient has a very high heart rate variability and maintains a rhythm above 140/min, heart failure is an imminent threat\(^\text{1,30}\). Ablation of the AV junction and placement of a variable frequency ventricular pacemaker (VVIR) are the indicated option. The use of anticoagulants is imperative even in patients who have regained sinus rhythm after isolating the pulmonary veins, so there is no argument against it\(^\text{31,32}\). Radiofrequency thermal injury of the AV junction causes an irreversible blockage. The injured tissue can be the AV node or the His bundle and can be achieved either from the tricuspid ring or from the left ventricle\(^\text{1}\). The success of this procedure is very close to 100%, and the possibility of recurrence is practically null. The
placement of a ventricular pacemaker is a routine procedure in any institution, with low risk and ventricular function improved by obtaining regularity of rate. In a series of patients in the Unit of Arrhythmias of Experimental Medicine of the UNAM in the General Hospital of Mexico, 177 ablations of the AV junction and placement of ventricular pacemaker have been carried out. All patients showed a ventricular rate variability (ventricular function normal) > 140 bpm, when the normal is above 100 bpm. Many of them, during 6 min of walking, could not perform more than 250 m. In 159 of the patients, ablation of the AV Union achieved from the right atrium, and in 17 (10%), it had to be done from the left ventricle. In no case, there was a recovery of AV conduction. This study concludes that ablation of the AV node is affordable and feasible in cases of permanent AF. Isolation of pulmonary veins should not be performed as and attempt to recover sinus rhythm, even if other options have been exhausted. Anticoagulation is mandatory in almost all patients, with AF, regardless of its type.

What is the clinical benefit and what is the purpose of pulmonary veins isolation in AF

In general, there is no definitive cure for AF; the therapeutic goal is to control symptoms, delay disease progression, and prevent a cardiovascular event. Electrical isolation of the pulmonary veins when there is recurrence with drug treatment may be the most effective strategy for maintaining sinus rhythm and keeping the individual asymptomatic. Invasive electrophysiological treatment is relatively recent; it began when it was discovered that premature atrial contractions from the pulmonary veins were responsible for initiating AF; which led to the establishment of the selective elimination of these ectopic foci as a therapeutic objective. At present, the strategy is broader, trying to make electrical isolation of all pulmonary veins from the antrum and not from the ostium to avoid side effects such as pulmonary stenosis. Other cases of more advanced disease require different ablation strategies such as supplemental lesion in the left or right atrium, or even both, as well as in the superior vena cava or cavotricuspid isthmus. AF is a progressive disease, starting with tachycardia of the pulmonary veins (they usually arise from there, but they can be originated in other sites) that initiate AF; however, AF produces more AF with a remodeling, not only anatomical but also electrical process of the atria. If AF is prolonged enough, it becomes a bialtrial disease with fibrosis, electrical remodeling, and dilation of both atria that causes rotor systems that support it, making it finally permanent. Technology and knowledge have evolved with results of radiofrequency catheter ablation (RFCA) of 74% of patients in sinus rhythm at 1 year of follow-up. AF ablation is recommended in paroxysmal, persistent, and persistent AF of long duration refractory or intolerant to antiarrhythmic drugs; it may also be considered as the first line in symptomatic paroxysmal AF. The therapeutic objective is to create a series of lesions that prevent AF by eliminating the triggering extrasystoles or modifying the substrate that maintains it. At present, ablation strategies depend on the type of AF; if it is paroxysmal AF, the success rate is higher, since the isolation of the pulmonary veins is sufficient to maintain sinus rhythm. On the other hand, if it is persistent AF, the success rates are lower; in these cases, the therapeutic strategy is broader, requiring different ablation lines and searching for rotors not only in the left atrium but also in the right atrium, and even in other thoracic veins such as the coronary sinus, caval veins, or Marshall’s vein. This complexity leads to a significant reduction of the long-term success rate, requiring two or more procedures to make it more likely that the patient maintains sinus rhythm. Because of these results, patients with paroxysmal AF are now preferred for early intervention. Scientific evidence shows that the main factor for maintaining sinus rhythm is achieving complete electrical isolation of the pulmonary veins. In advancing stages the posterior wall, also plays an essential role in the maintenance in the maintenance of sinus rhythm. The techniques employed can be two, with RFCA using irrigation catheters or with cryoballoon ablation (CBA); the latter was limited only for paroxysmal AF but, nowadays it is safe to perform it in persistent AF with the advantage of being a less operator-dependent, with a faster learning curve and above all, fewer complications than RFCA, with comparable results in comparative studies. In centers of high experience, it can give results of up to 85% of patients free of AF at a 12 month follow-up. In Mexico, in the series published by the Instituto Nacional de Cardiología (Clínicas Mexicanas de Cardiología) of RFCA, in a period of 8 years, in patients with paroxysmal AF, there is 78% success in a 12-month follow-up in a total of 121 patients. CBA is the first experience in Mexico from 2013 to 2014 in a multicenter study (unpublished data from Hospital Ángeles Interlomas, CMN Siglo XXI, CMN 20 de Noviembre and Servicios de Salud del Estado de Puebla) with 52 patients, exclusively with paroxysmal AF, was successful in 78% of cases with 18-month follow-up.
When and how should antithrombotic prophylaxis be given in the subject with AF? Antiplatelet drugs, Vitamin K antagonist (VKA), direct oral anticoagulants (DOAC) and left atrial appendage occluders (LAAO)

**AF is a cause of stroke**

The presence of AF has long been associated with the development of cerebral and systemic (pulmonary, limb, coronary, renal, and visceral) embolism. Initially, only AF secondary to valvular disease, usually rheumatic heart disease, was considered thrombogenic, but since the Framingham study, AF of non-rheumatic origin is also recognized as a cause of embolism.

The prevention of embolisms in “valvular” AF should perform with VKA

For embolic risk purposes, “valvular” AF is considered to be the one associated with moderate or severe mitral stenosis or in the presence of a mechanical valve prosthesis. Although acetylsalicylic acid was initially used in patients with rheumatic heart disease, subjects with valvar AF should now be anticoagulated with VKA, either acenocoumarin or warfarin. The dose is that necessary to achieve an international normalized ratio (INR) between 2.0 and 3.0, except for patients with mechanical valve prostheses that require INR between 2.5 and 3.5. DOAC should not be used in valvar AF until the results of studies supporting this practice are available. To improve the time in therapeutic intervals, it is recommended to: (1) establish anticoagulation clinics and (2) self-monitoring of the INR with portable devices.

**CHA2DS2-VASc scale and options for prevention of embolisms in “non-valvular” AF**

For patients with AF not associated with mitral stenosis or a mechanical valve prosthesis, a choice can be made between anticoagulant drugs, VKA or DOAC. Antiplatelet agents have the weakest effect in preventing embolism. In the joint analysis of randomized studies, the relative risk reduction of stroke by anabolic-androgenic steroids (AAS) compared to placebo was calculated at 19% while with VKA, it was 64%. It is important to note that based on the results of the ACTIVE-W study, dual antiaggregation therapy (e.g. AAS and clopidogrel) is not recommended over oral anticoagulation.

The decision of which drug should be used in the prevention of cerebral infarction can be based on the use of the CHA2DS2-VASc scale (Table 1). For individuals with no points, (no risk factors, considered “low risk” by not observing any embolic event in a follow-up year) in general it is possible to choose not to give treatment; in those with a score of one (“intermediate risk” of 0.6% of an embolic event per year) if it is male or two if it is female, they benefit more with oral anticoagulation with VKA or DOAC. The HAS-BLED or ATRIA scales can be used to assess the risk of bleeding.

**LAAO**

LAAO are an interventional option for the prevention of embolism that so far is only indicated for patients with high embolic risk and who have some contraindication to receive VKA or DOAC. Outside of this select group of patients, implanting these devices as substitutes for anticoagulation do not yet have sufficient evidence. The most recent results on cost-benefit analysis using dedicated statistical models (e.g. Markov’s stochastic decision model) have yielded contradictory results. However, several studies are ongoing and are expected to produce positive results for occluders. Like any invasive procedure, its efficacy in preventing stroke should weigh against possible complications of its implant.

**Final remarks**

AF, in its different forms, is considered to be the most frequent tachyarrhythmia in Mexico and should be considered as a public health problem. Its treatment...
includes “rhythm control” with a few antiarrhythmic drugs available in Mexico for this purpose. Ventricular rate control can be achieved with drugs or some intervention-al procedures, included AV junction ablation with a VVIR pacemaker implant. The role of pulmonary vein isolation is undoubted for clinical relief of symptoms with many ongoing studies on the possible effect on morbidity-mor-bidity. Thromboprophylaxis is a key and integral part of the management of any patient with AF. Recently, CENETEC (National Center for the Technical Excellence in Health, Health Ministry of Mexico) published guidelines on the antithrombotic treatment of AF\(^5\).39.

### Acknowledgments

- Endorsed by: Mexican National Association of Cardiologists (ANCAM), Mexican Electrophysiological and Pacing Society (SOMEEC) and Mexican Society of Cardiology (SMC).
- Avalado por: Asociación Nacional de Cardiòlogos de México (ANCAM), Sociedad Mexicana de Electrofisiología y Estimulación Cardiaca (SOMEEC) y Sociedad Mexicana de Cardiología (SMC).

### Conflicts of interest

The authors declare that they do not have any conflicts of interest in this paper.

### Table 1. Risk factors for cerebral infarction included in the “CHA\(_2\)DS\(_2\)-VASc” scale and hemorrhagic risk factors included in the “HAS-BLED” scale

<table>
<thead>
<tr>
<th>CHA(_2)-DS(_2)-Vasc</th>
<th>Score</th>
<th>HAS-BLED</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>C (Congestive heart failure) = Left-sided heart failure</td>
<td>1</td>
<td>H = Hypertension</td>
<td>1</td>
</tr>
<tr>
<td>H = Hypertension</td>
<td>1</td>
<td>A = Impaired liver or kidney function</td>
<td>1 each</td>
</tr>
<tr>
<td>A (Age) = ≥ 75 years</td>
<td>2</td>
<td>S (Stroke) = Cerebral vascular disease</td>
<td>1</td>
</tr>
<tr>
<td>A (Age) = Age 65-74 years</td>
<td>1</td>
<td>B (Bleeding) = Bleeding</td>
<td>1</td>
</tr>
<tr>
<td>D = Diabetes \textit{mellitus}</td>
<td>1</td>
<td>L (Labil INR) = Highly variable INR (outside therapeutic intervals)</td>
<td>1</td>
</tr>
<tr>
<td>S (Stroke) = Previous stroke</td>
<td>2</td>
<td>E (Elderly)</td>
<td>1</td>
</tr>
<tr>
<td>S = Sex category</td>
<td>1</td>
<td>D (Drugs) = Drugs or alcohol</td>
<td>1 each</td>
</tr>
<tr>
<td>V = Peripheral vascular disease</td>
<td>1</td>
<td></td>
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</tbody>
</table>

INR: international normalized ratio; AF: atrial fibrillation.

### Funding

This investigation has not received any financial support.

### ethical disclosures

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data. The authors declare that no patient data appear in this article.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

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