

This is the author's manuscript



AperTO - Archivio Istituzionale Open Access dell'Università di Torino

Evidence and uncertainties in the management of atrial fibrillation in older persons

Original Citation:	
Availability:	
This version is available http://hdl.handle.net/2318/1844109	since 2022-12-05T12:02:41Z
Published version:	
DOI:10.23736/S0026-4806.21.07525-X	
Terms of use:	
Open Access Anyone can freely access the full text of works made available as under a Creative Commons license can be used according to the t of all other works requires consent of the right holder (author or protection by the applicable law.	erms and conditions of said license. Use

(Article begins on next page)

Minerva Medica EDIZIONI MINERVA MEDICA

ARTICLE ONLINE FIRST

This provisional PDF corresponds to the article as it appeared upon acceptance.

A copyedited and fully formatted version will be made available soon.

The final version may contain major or minor changes.

Evidence and uncertainties in the management of atrial fibrillation in older persons

Paolo ALBONI, Mario BO, Stefano FUMAGALLI, Francesco VETTA, Gianluca ISAIA, Enrico BRUNETTI, Samuele BALDASSERONI, Alessandro BOCCANELLI, Giovambattista DESIDERI, Niccolò MARCHIONNI, Renzo ROZZINI, Pierfranco TERROSU, Andrea UNGAR, Giovanni ZITO

Minerva Medica

DOI: 10.23736/S0026-4806.21.07525-X

Article type: Review Article

© 2021 EDIZIONI MINERVA MEDICA

Article first published online:

Manuscript accepted: March 29, 2021 Manuscript received: March 24, 2021

Subscription: Information about subscribing to Minerva Medica journals is online at:

http://www.minervamedica.it/en/how-to-order-journals.php

Reprints and permissions: For information about reprints and permissions send an email to:

journals.dept@minervamedica.it - journals2.dept@minervamedica.it - journals6.dept@minervamedica.it

Evidence and uncertainties in the management of atrial fibrillation in older persons Running title: Atrial fibrillation in older persons

Paolo ALBONI¹, Mario BO¹, Stefano FUMAGALLI¹, Francesco VETTA¹, Gianluca ISAIA², Enrico BRUNETTI^{2*}, Samuele BALDASSERONI¹, Alessandro BOCCANELLI¹, Giovambattista DESIDERI¹, Niccolò MARCHIONNI¹, Renzo ROZZINI¹, Pierfranco TERROSU¹, Andrea UNGAR¹, Giovanni ZITO¹

¹SICGe – Società Italiana di Cardiologia Geriatrica, Via Matteotti 7, Firenze, Italy
² Section of Geriatrics, Department of Medical Sciences, University of Turin, Città della
Salute e della Scienza – Molinette Hospital, Turin, Italy

*Corresponding author: Enrico Brunetti, Section of Geriatrics, Department of Medical Sciences, University of Turin, Città della Salute e della Scienza – Molinette Hospital, corso Bramante 88/90, 10126, Torino, Italy. E-mail: enrico.brunetti@unito.it

ABSTRACT

INTRODUCTION: Atrial fibrillation (AF) is the most common cardiac sustained arrhythmia, whose incidence and prevalence increase with age, representing a significant burden for health services in western countries. Older people contribute to the vast majority of patients affected from AF.

EVIDENCE ACQUISITION: Although oral anticoagulant therapy represents the cornerstone for the prevention of ischemic stroke and its disabling consequences, several other interventions – including left atrial appendage occlusion (LAAO), catheter ablation (CA) of AF, and rhythm control strategy (RCS) – have proved to be potentially effective in reducing the incidence of AF-associated clinical complications. Scientific literature focused on the three items will be discussed.

EVIDENCE SYNTHESIS: Practical treatment of older AF patients is presented, including approach and management of patients with geriatric syndromes, selection of the most appropriate individualized drug treatment, clinical indications and potential clinical benefit of LAAO and CA in selected older AF patients.

CONCLUSIONS: Older people carry the greatest burden of AF in real world practice. Within a shared decision making process, the patient centered approach need to be put in

COPYRIGHT© EDIZIONI MINERVA MEDICA

the context of a comprehensive assessment, in order to gain maximal net clinical benefit and avoid futility or harm.

Key words: atrial fibrillation, older patients, oral anticoagulants, left atrial appendage occlusion, AF catheter ablation

TEXT

INTRODUCTION

Atrial fibrillation (AF) is one of the most common form of cardiac arrhythmia. Both incidence and prevalence of this disorder increase with advancing age, representing a significant medical burden for health services in most western countries [1]. The currently estimated prevalence of AF in adults is between 2% and 4%, and a 2.3-fold rise is expected, owing to extended longevity in the general population and intensifying search for undiagnosed AF [2]. Indeed, prevalence of AF has been reported to be around 10% among persons aged 80 years and over in the general population [3]. Patients with AF have an increased risk of ischemic stroke, which is approximately 5-fold higher than the general population, with an incidence of approximately 5% per year. In addition, as AFrelated cardioembolic strokes are associated with higher mortality and morbidity than other strokes, the need for more effective stroke prevention in these patients should be emphasized [4,5]. Atrial fibrillation poses significant burden to patients, physicians, and healthcare systems globally. The complexity of AF requires a multifaceted, holistic, and multidisciplinary approach to the management of AF patients. Recent European Guidelines summarize and evaluate available evidence with the aim of assisting health professionals in proposing the best management strategies for an individual patient with AF [2]. The Atrial Fibrillation Better Care (ABC) approach in the 2020 European Society of Cardiology (ESC) AF Guidelines is a continuum of this approach, with the goal to further improve the structured management of AF patients, promote patient values, and finally improve patient outcomes. In this review we will discuss some common clinical uncertainties in the management of AF in older persons, dealing with the following issues: 1) practical use of oral anticoagulants; 2) percutaneous left atrial appendage occlusion; 3) catheter ablation of AF.

METHODS

Scientific literature focused on the three items discussed (use of oral anticoagulant therapy, percutaneous left atrial appendage occlusion and catheter ablation of AF) in older persons published in the last 8 years was retrieved by the authors (MB, FV, PA, GI, EB) from the MEDLINE database entering these terms and using the terms "atrial fibrillation", OR "new oral anticoagulants" OR "direct oral anticoagulants", OR "aged" OR "elderly"

OR "older" as keywords. Reviews, recommendations and expert opinions, as well as clinical trials and large observational studies in English published until December 2020 were systematically analyzed and included according to their relevance to the objective. Additional references were obtained from the reference list of the selected full-text manuscripts.

ORAL ANTICOAGULANT THERAPY (OAT) IN OLDER AF PATIENTS

Current European guidelines recommend oral anticoagulant therapy (OAT) with direct oral anticoagulants (DOACs) over vitamin K antagonists (VKAs) irrespective of age for patients with AF and a CHA_2DS_2 -VASc score ≥ 2 in men and ≥ 3 in women, and without contraindications to DOACs (mechanical prosthetic valves or moderate-to-severe mitral valve stenosis) [6,7]. Phase III DOAC randomized clinical trials (RCTs) enrolled a significant proportion of older subjects, and consistently demonstrated a greater net clinical benefit compared to VKAs also in persons aged 75 years and over, who account for the largest proportion of AF patients [8-10].

Geriatric syndromes in AF older patients

However, despite consistent evidence of clinical benefit and increasing prescription of these drugs [11], they are still widely underused, particularly in the oldest patients [12,13]. Some uncertainties in DOACs use in older patients might arise from the concern that the significant proportion of older persons enrolled in DOACs RCTs might not be fully representative of real world (RW) patients. However, with the inherent limitations of the observational design, real world studies confirm a greater net clinical benefit of DOACs compared with VKAs also in older patients, with an apparent better safety profile for apixaban and low dose dabigatran [14-18].

Moreover, geriatric syndromes such as frailty, cognitive impairment and functional dependence, which have been demonstrated to influence physicians' decision about DOACs use in older persons [19,20], were not considered in RW studies as well as in DOACs trials. Although cardiologists usually recognize frailty based on the presence of a mix of problems of motility, cognition, nutrition and inappropriate loss of body weight and muscle mass [21], there are two basic conceptualizations of frailty. The frailty "phenotype" is based on the presence of at least three of five criteria – slow gait speed, low physical activity, unintentional weight loss, self-reported exhaustion, and muscle weakness – and is associated with worsening mobility and disability, hospitalizations, and

mortality over 7 years in community-dwelling older persons [22]. This "frailty phenotype", which should not be confused with disability or comorbidity, may also be identified using other tools, such as the Simplified Fried test, the Short Physical Performance Battery (SPPB), the 5 meter gait speed, the Study of Osteoporotic Fractures (SOF) index and the simple Frail Scale [23]. On the other side, the Frailty Index [24] is a 70-item form based on the accumulation of deficits (including functional limitations and disabilities, cognitive and sensory impairment, psycho-social variables and number of diseases), whose score is associated with increased short term risk of institutionalization, mortality and hospitalization. The 7-point Clinical Frailty Scale (a semi-quantitative eyeball global judgment of frailty or vulnerability) was shown to be highly correlated with the Frailty Index and significantly associated with increased risk of death and entry into an institution [25]. The Multidimensional Prognostic Index (MPI) (including information on functional basic and instrumental activities of daily living, cognitive and nutritional status, comorbidities, medications, and social support network) has also been demonstrated to be predictive of mortality and adverse clinical outcomes [26,27]. In summary, the "frailty phenotype" based tools identify patients at risk of disability, but not of short term mortality, whereas high scores in the Frailty Index, Clinical Frailty Scale and MPI identify patients with poor health status and increased risk of mortality. Despite inherent limitations according to different frailty tools adopted, frail older patients with AF are less likely to receive an appropriate anticoagulant prescription and, at the same time, are at greater risk of embolic stroke and death [28,29]. The lack of evidence to guide optimal care for patients with AF and frailty might in part explain the gap between current guidelines and clinical practice in the management of these patients [29]. On the basis of current evidence there is general agreement that the "frailty phenotype" should not be an exclusion criterion to anticoagulate, since these patients are at increased risk of stroke and have been shown to benefit from OAC [30].

Predisposition to falls is common in frail patients and is often perceived as an important issue in starting DOACs [21,31,32]. However, patients on OAT at high risk of falls did not consistently have a significantly increased risk of major bleedings (MBs) [33-35]. Current guidelines do not require fall risk estimation in candidates to OAT, and the risk of fall per se should not be considered a contraindication to the use of DOAC [2,6,7].

Many older adults have both cognitive impairment or overt dementia and AF. Although AF is a recognized risk factor for later occurrence of cognitive impairment and dementia

[36], dementia is a well-recognized risk factor for under-use of OAT [12]. A retrospective cohort study of 2572 older patients with AF (73% aged ≥75 years) showed that after diagnosis of dementia, those who persisted on OAT had lower rates of stroke and all-cause mortality, with no significant differences in risk of MBs [37]. Although cognitive impairment and frailty were associated with increased risk of death and reduced probability of receiving OAT among older AF patients enrolled in the ORBIT-AF registry [38], there was no interaction between OAT use and cognitive impairment or frailty in their association with mortality, major bleeding and a composite end-point of stroke, systemic thromboembolism, myocardial infarction and cardiovascular death [38]. Whereas cognitive impairment at mild-to moderate stage should not be viewed as a general contraindication to DOAC therapy, especially if well-managed from a logistic point of view, in states of poor physical functioning, limited life-expectancy and high risk for competing causes of death there may be limited benefit from OAT [6,7].

Persistent uncertainties about OAT use in older AF patients

Despite recent studies reinforced the evidence of net clinical benefit of OAT, including DOACs, in extremely older community-dwelling persons (aged ≥85 years) [39], prescription of OAT to older AF patients is often a troublesome decision, involving a global evaluation of health, residual life-expectancy, functional and cognitive status, rather than a simple addition of variables within cardio-embolic and bleeding risk scales [9]. It is likely that sometimes physicians perceive OAT as "futile" or potentially harmful in patients with multi-morbidity and short life-expectancy, and, moreover, cost-effectiveness considerations might affect decision about DOACs prescription in these patients. Indeed, when considering OAT with DOACs in older persons, the high risk of competing cardiovascular and non-cardiovascular causes of death in this population should be considered. In fact, while the adjusted overall mortality in landmark phase III DOAC trials was 4.72%/year, with cardiac death contributing for 46% of deaths [40], all-cause mortality in real-world older patients are definitely higher, with difference in causespecific mortality. In a population-based, retrospective cohort study of a nationally representative sample of fee-for-service Medicare beneficiaries 65 years or older with incident atrial fibrillation diagnosed between 1999 and 2007, including 186 461 patients with AF and no recent hospitalizations for heart failure, myocardial infarction, stroke, or gastrointestinal hemorrhage, mortality was the most frequent major clinical events (19.5% at 1 year; 48.8% at 5 years), with an incidence which was 7-fold higher than that of ischemic stroke. In this sample, the risk of all-cause mortality was further increased in inpatients compared with out-patients [41]. In the ORBIT-AF registry, older patients not on OAT experienced higher mortality rates (7.42 vs 5.78%, p=0.006) over a 2.5 years followup without significant differences in thromboembolic event rates, compared with patients receiving OAT [42]. Data from the Galician Healthcare Service showed that among patients aged 80 years and older (45.6% of those with AF) two-year all-cause mortality was higher than in younger counterparts (27.8% vs 8.05%, p<0.001), as well as thromboembolic and hemorrhagic events (2.03% vs 0.9%, p<0.01 and 2.5% vs 1.7%, p=0.01, respectively) [43]. In two studies including hospital discharged older AF patients (mean age over 80 years) we documented high mortality rates, mainly for noncardiovascular causes, which were about two-fold higher in untreated patients, reflecting the higher proportion of poor health status in these latter patients [44, 45]. A recent systematic review and meta-regression analysis demonstrated that in older AF patients DOACs are superior to warfarin for stroke/thromboembolism prevention, with reduced risk of MB, thereby reinforcing the evidence that DOACs should be preferred for stroke prevention in older AF patients [46]. However, some older AF patients are at risk of increased short-term all-cause mortality, thereby diluting the undisputable benefit of DOACs. Unfortunately, by now there are not validated methods to identify those few older patients who, because of their poor general health and/or functional status, are expected not to have a net clinical benefit from anticoagulation [30].

Use of DOACs in older AF patients

Medical history and comorbidities may drive the choice of a particular DOAC in older patients. Several DOACs rankings [47-49] and expert opinions have been published to assist physicians to fit the best DOAC according to individual patient's characteristics [50-53]. Apixaban has been suggested as a reasonable first choice both in older patients and in subjects with chronic renal failure [47,52,53]. The recently updated 2019 American Geriatrics Society Beers criteria recommend a cautious use of dabigatran and rivaroxaban in AF patients aged ≥75 years because of greater risk of gastrointestinal bleeding [54]. In a recent report from the Fit-fOR-The-Aged (FORTA) classification (evaluating benefit, risk and appropriateness of drugs for older patients in everyday clinical settings), apixaban was labelled A among OATs, meaning it was seen as the drug with the most favorable risk/benefit ratio in older patients [55].

In RW clinical practice use of reduced-dose DOACs and inappropriate off-label DOAC under-dosing, particularly of apixaban and rivaroxaban, are quite common, mainly in the oldest patients and with poor health status [56-59]. In a recent review [60], we demonstrated that several conditions, including advanced age, female gender, fear of bleeding and/or previous bleeding, history of chronic kidney disease and polypharmacy, are associated with oral Factor Xa Inhibitors (FXaIs) underdosing, supporting the hypothesis that a substantial proportion of these prescriptions may be voluntary rather than casual. However, the vast majority of bleedings occurs within well-conducted OAT, and appears to be associated with patient's characteristics (e.g., advanced age, comorbidity, anemia, previous bleedings, concomitant therapy with antiplatelet drugs or non-steroidal anti-inflammatory drugs) and underlying gastrointestinal pathology, rather than with OAT intensity. Indeed, in FXaI RCTs, the rates of MB were consistently higher among patients treated with reduced dose (RD) rather than in those treated with full dose (FD). Moreover, correct use of RD had reassuringly the same efficacy and safety as FD FXaIs compared with warfarin. Real-life studies do not provide evidence of a sizeable net clinical benefit by using off-label RD FXaIs, but rather suggest an increased risk of adverse events, including hospitalizations for cardiovascular causes and stroke, particularly in patients on underdosed apixaban. Current evidence should discourage FXaIs underdosing and support their prescription according to drug-specific dosing guidelines for the most patients.

Conclusions

The availability of DOACs has dramatically increased the proportion of older AF patients receiving appropriate OAT. Because of their potential for clinical benefit, DOACs should be recommended for "fit and robust" older subjects, as well as for persons with the frailty phenotype, irrespective of age; risk of falls, cognitive impairment without functional limitations, and mild disability should not be regarded as contraindications to DOAC use in these patients. However, as for many other preventive therapies, at the moment there is no evidence of a net clinical benefit from OAT in older patients with advanced dementia, and/or with loss of functional independence, and/or short life expectancy [30]. Hopefully, further studies will provide information in this setting of patients. Individual selection of DOAC and use of recommended appropriate dose, careful clinical surveillance, periodic review of co-medications, and minimization of bleeding risk are mandatory in these patients.

PERCUTANEOUS LEFT ATRIAL APPENDAGE OCCLUSION FOR THE PREVENTION OF STROKE IN OLDER AF PATIENTS

The left atrial appendage (LAA) in the pathogenesis of thrombus

The thrombogenesis in AF is multifactorial and includes Virchow's triad which, through endothelial and endocardial damage or dysfunction, abnormal blood stasis, and altered hemostasis, platelet function and fibrinolysis, promotes the onset of thrombosis [5,61-62]. These changes are most evident in the left atrial appendage (LAA), where a remarkably low blood flow velocity plays a very important role in promoting thrombus formation. The LAA, which is the remnant of the embryonic left atrium, is a tubular blind-ended structure with several lobes and variable morphology, in contrast to the wide triangular shape of the right atrial appendage [62-65]. Unlike the right atrium, pectinate muscles are located within the LAA and do not extend into the remaining parts of the left atrium. Several forms and variants of the LAA have been described, with 1-4 lobes and 4 prevailing morphologies, called "windsock", "cauliflower", "cactus-like" and "chicken wing" [63]. Its complex structure with areas of relatively low flow velocity predisposes to stasis, mainly during AF when blood flow velocity is further reduced, as can be visualized on transthoracic or, better, transoesophageal echocardiographic examination (TOE) with evidence of spontaneous contrast (smoke effect) or on pulsed wave Doppler during AF paroxysms [63-67]. Because of these predisposing situations, it has been shown that in patients with non-valvular AF, about 91% of thrombi develop in the LAA as opposed to 57% only in patients with rheumatic AF [63,68]. In fact, literature data show that the presence of thrombi or a reduced peak flow rate in the LAA are independent predictors of increased thromboembolic risk [63,68], as well as stroke recurrence among patients with non-valvular AF. However, a recent analysis has shown that while in valvular AF more than half of the thrombi were in the left atrial cavity even in patients with non-valvular AF and a history of stroke, the chances of developing a thrombus in the left atrial cavity compared to LAA were up to 45% in the case of missed anticoagulation or ventricular dysfunction [69].

In NVAF patients with a CHA_2DS_2 -VASc score ≥ 2 for men and ≥ 3 for women, DOAC use is a class I recommendation for primary and secondary prevention of cerebral and systemic embolisms [7] with consistent greater clinical benefit over warfarin also in older patients [9,70]. LAA occlusion may be considered (Class IIb; Level of Evidence B) for stroke prevention in patients with AF and contra-indications to long-term use of OAT [6].

LAA Occluders in Clinical Trials

In recent decades, there has been a progressive development of transcutaneous LAA occlusion (LAAO) techniques along with an increasing availability of different LAAO devices, although many of them yet unapproved [71]. Therefore, most of current evidence is based on the WATCHMAN LAA occluder (Boston Scientific, Marlborough, Massachusetts), the most extensively used in RCTs and observational studies (Table I). Over the past decade, two RCTs using the WATCHMAN device, PROTECT AF (WATCHMAN Left Atrial Appendage Closure Device for Embolic Protection in Patients with Atrial Fibrillation) and PREVAIL (Prospective Randomized Evaluation of the WATCHMAN Left Atrial Appendage Closure Device in Patients With Atrial Fibrillation versus Long-Term Warfarin Therapy) [72,73] as well as registry data on patients with various LAA closure devices have demonstrated excellent safety and efficacy compared with the use of VKAs [74-76]. A recent meta-analysis demonstrated that the WATCHMAN device was non-inferior to VKAs for the prevention of stroke and systemic thromboembolism and with a significantly lower risk of hemorrhagic stroke, nonprocedure-related MB, and mortality at 5 years follow-up [77]. However, patients enrolled in these RCTs and registries received VKA-based OAT for 6 weeks to facilitate endothelialization of the device. As a consequence, the WATCHMAN device received FDA regulatory approval in the United States for use only in patients eligible for shortterm OAT, thereby excluding access to this device for patients with absolute contraindications to OAT. In contrast, ESC guidelines recommend the use of percutaneous LAA closure in patients with AF and prior life-threatening bleeding or contraindications to long-term OAT (Class IIb; Level of Evidence B) [6] using dual antiplatelet therapy (DAPT) with aspirin and clopidogrel in the postprocedural period. Currently, there are no randomized data to support this recommendation, and most of the evidence comes from prospective single-center and multicenter observational registries including the WATCHMAN and Amplatzer Occlusion Device (St. Jude Medical, Minneapolis, Minnesota) [74, 78-80].

In the prospective multicenter EWOLUTION Registry (Evaluating Real-Life Clinical Outcomes in Atrial Fibrillation Patients Receiving the WATCHMAN Left Atrial Appendage Closure Technology), including approximately 72% of patients ineligible for OAT, the post-implantation antithrombotic strategy was highly variable including use of single and dual antiplatelet therapy and VKAs. The rate of stroke observed in the entire

cohort was only 1.3%/year, and the combined end-point of ischemic stroke, transient ischemic attack (TIA), and systemic embolism was 2%/year: these rates correspond to a risk reduction of 83% and 80%, respectively, compared with that expected in untreated patients. [79-81]. Notably, these rates of ischemic and bleeding events are similar to those observed in the combined PROTECT AF and PREVAIL RCT data [77], where the WATCHMAN group received combination therapy of aspirin 81 mg daily and warfarin (target INR, 2-3) for 45 days followed by DAPT (aspirin 81 mg daily+clopidogrel 75 mg daily) for 4.5 months, and then lifetime aspirin 81 mg daily. Despite, the growing body of RW evidence on the use of single or DAPT after LAAO, the appropriate postprocedural antithrombotic regimen is not well defined and clouded by device-related thrombus (DRT), leakage, and bleeding events. In the EWOLUTION Registry, the rate of DRT, despite the highly variable antithrombotic strategy, was 4.1%/year, substantially superimposed on that of the ASAP trial and similar to the 3.7%/year observed in the combined analysis of the 4 prospective WATCHMAN FDA clinical trials (PROTECT AF, PREVAIL, CAP, and CAP-2) in which the warfarin/DAPT regimen was used [82].

A high risk of MB is typically the most common reason why patients receive LAAO [63,66]. The nonprocedural MB rate in the EWOLUTION study was 2.7%, with the lowest rates (1.1%) in patients with early discontinuation of DAPT (≤105 days) when compared with those who continued DAPT beyond that limit (3.5%) with no significant difference in the combined end point of systemic thromboembolism, TIA, and DRT between groups [79-81]. Furthermore, the benefit of LAAO in reducing stroke and MBs was also observed in high-risk subgroups, such as patients with CHA2DS2-VASc score ≥3 and history of stroke, TIA, MB, or hemorrhagic stroke. These findings strongly raise the question regarding the initial transition with OAT post-WATCHMAN closure of LAA and whether the current recommended strategy provides an additional incremental benefit over DAPT alone. Larger studies are needed to better define the role and duration of DAPT in the post-LAAO phase.

Since the vast majority of RCTs with LAAO were initiated before the widespread use of DOACs, the need to compare LAAO procedures with respect to the use of DOACs is becoming increasingly compelling. Compared with warfarin, DOACs are certainly easier to use and are associated with a greater clinical net benefit even in older subjects [9,30,70]. Transcatheter LAAO performs similarly to DOACs in RCTs against warfarin, showing a significant reduction in intracranial hemorrhage, with no statistically significant

increase in ischemic stroke and a possible reduction in all-cause mortality [77,83]. In addition, LAAO has been shown to be associated with a reduction in the risk of gastrointestinal hemorrhage [84]. The prospective, randomized, noninferiority PRAGUE-17 study aimed to compare transcatheter LAAO with DOAC therapy in 402 patients (mean age 73 years) [85]. Enrollment criteria were based on failure of DOAC treatment, significant prior bleeding, or a combination of high thromboembolic risk and high bleeding risk. Patients were randomly assigned to DOAC (mostly apixaban) or LAAO therapy. The primary endpoint was a composite outcome that included both safety and efficacy and periprocedural complications. During a 20 months period of observation, the PRAGUE-17 data demonstrated non-inferiority of LAAO compared with DOACs, with a similar rate of all-cause stroke and a lower rate of bleeding in patients receiving LAAO. Despite the numerical and methodological limitations, PRAGUE-17 has certainly the merit of reinforcing the role of LAAO in patients at high risk of bleeding or failure of drug treatment, even in the era of DOACs. Hopefully, incoming studies will provide more substantial and convincing data on the use of LAAO in subjects over 80 years old.

Clinical considerations for older AF patients

The mean age of the patients involved in the aforementioned RCTs and registries was slightly above 70 years and, although almost half of the patients enrolled were aged >75 years (Table I), there are no subanalyses related to this age group. In any case, the few data in the literature on octogenarian subjects do not show age-related differences in terms of both efficacy and safety. Even the risk of periprocedural complications does not seem to increase in relation to the age of patients showing, over time, a progressive reduction in incidence as a function of the physiological learning curve of the operators related to the greater diffusion of this technique. Recent data from the prospective observational Left-Atrium-Appendage Occluder Registry Germany, including a total of 638 patients (402, 63%, aged ≥75 years), reported similar high procedural success rate in both groups (97.6%), without significant differences of adverse events between older and younger patients. At one year follow-up, all-cause mortality was higher in patients aged ≥75 compared with younger group (13.0% vs 7.8%, p=0.04), mainly due to non-cardiovascular causes (10.6% vs 6.0%) [86].

It is also interesting to point out that there are currently no data in the literature that analyze the weight of common conditions in the elderly such as frailty syndrome, comorbidities, polypharmacy in relation to the use of LAAO devices. The evaluation of

the risk/benefit ratio of LAAO in the elderly based on the above determinants is of paramount relevance to assess the possible futility of this procedure in relation to both the estimated residual life expectancy as well as the quality of life, and the residual thromboembolic risk related to specific comorbidities. In fact, it is known that several diseases such as heart failure with severe ejection fraction impairment are associated with an increased risk of occurrence of thrombus outside the LAA, as already described by the meta-analysis of Mahajan et al [69] that showed, in patients with non-valvular AF, a prevalence of atrial thrombi outside the LAA of 11%, which doubled in older patients. Therefore, although at present potential candidates for LAAO are those with absolute or relative contraindications to OAT, or patients at high risk of recurrent ischemic stroke, severe renal dysfunction, severe gastrointestinal bleeding, in older patients, pending specific trials, it is useful that the assessment of the risk-benefit ratio is done, for each individual case by a multidisciplinary team involving the patient and care givers.

CATHETER ABLATION IN OLDER PATIENTS WITH AF

Catheter ablation (CA) is increasingly used for AF treatment since antiarrhythmic drugs have demonstrated a limited efficacy in maintaining stable sinus rhythm. Some RCTs comparing antiarrhythmic drugs for rhythm control with rate control did not demonstrate a benefit of one treatment strategy over the other [87]. However, very symptomatic patients were not enrolled. Very recently, the EAST-AFNET-4 RCT [88] demonstrated that AF patients (mean age 70 years) who were randomly assigned to early rhythm control (median time of 36 days after diagnosis) had a lower risk of death from cardiovascular causes, stroke, or heart failure hospitalization or acute coronary syndrome versus rate control (HR 0.79, 95% CI 0.66-0-94). Many observational and some small randomized studies investigated whether rhythm control therapy was more effective by utilizing antiarrhythmic drugs or AF CA and the results appear to be conflicting. However, some meta-analyses showed superiority of ablation in reducing episodes of recurrent AF and in improving quality of life [2,89], but many patients required two or more procedures. It should be stressed that in most studies the mean age of the patients was rather low, between 50 and 60 years, considering that the mean age of AF patients is approximately 75 years [2]. After single procedure, success was reported in 40-70% of the patients during a follow-up period of 12-24 months and the major risk factors for AF relapse were advanced age, long AF duration, the non-paroxysmal form of AF at baseline, structural

heart disease, increased left atrial volume and the presence of comorbidities such as hypertension, obesity. renal dysfunction, and sleep apnea [2,89]. Prospective, registry-based data show that 4-15% of patients experience periprocedural complications, 1-7% of which are defined as major (mainly stroke/TIA, symptomatic pulmonary vein stenosis, pericardial tamponade, atrioesophageal fistula, phrenic nerve injury, retroperitoneal bleeding, femoral arteriovenous fistula) [2,89]. The periprocedural mortality is approximately 0.1% [2]. Radiofrequency and cryoballoon ablation seem to have similar efficacy and complication rates, even if some studies show lower complication rates with cryoballoon ablation [2,89].

In two recent RCTs [90,91], hard endpoints such as death and stroke were investigated, besides the incidence of AF recurrence and the impact on quality of life. In the CABANA trial [90,92], 2204 symptomatic patients (median age 68 years) with paroxysmal or persistent AF were randomized to CA versus medical therapy (antiarrhythmic drugs in 88% of patients). This trial showed that CA is associated with an improvement in quality of life, a reduction in cardiovascular hospitalization, and a lower AF recurrence rate than drug therapy (50% versus 69% at 3 years follow-up). However, CA did not reduce the primary composite outcome of death, stroke, serious bleeding, or cardiac arrest compared with medical therapy. Therefore, this large trial shows that for many patients ablation is not curative and has not a prognostic impact, but it reduces AF burden and improves symptoms. In another recent trial (CASTLE-AF) [91], selected patients with paroxysmal or persistent AF, heart failure, left ventricular ejection fraction ≤35% and an implanted defibrillator were randomized to AF ablation versus medical therapy (rate or rhythm control). The study enrolled 363 patients, and the mean age was rather low (64 years). AF ablation significantly reduced the risk of death and heart failure hospitalization by 40%. There was also a benefit in all-cause mortality, which was driven by a significantly lower rate of cardiovascular death in the ablation group. Furthermore, CA reduced AF burden and improved left ventricular ejection fraction. However, the generalizability of the CASTLE-AF trial has recently been evaluated in a large heart failure patient population and this analysis showed that only a small number of patients (about 8%) met the trial inclusion criteria [93]. In the 2020 European guidelines [2], CA is recommended for rhythm control after one failed or intolerant class I or III antiarrhythmic drug to improve symptoms in patients with paroxysmal or persistent AF without major risk factors for AF recurrence (class I, level of evidence A) and in patients with persistent AF with major risk

factors for AF relapse (class I, level of evidence B). Moreover, CA should be considered in selected AF patients with heart failure with reduced ejection fraction to improve survival and reduce heart failure hospitalization (class IIa, level of evidence B).

Effects of AF ablation in the elderly

In order to maintain sinus rhythm, antiarrhythmic drugs can be difficult to manage due to unpredictable metabolism in elderly patients and intolerance of side effects. However, even the success rate of AF CA may be affected by a higher degree of atrial myopathy in older individuals. Randomized trials comparing efficacy and safety of AF ablation versus medical therapy in the elderly with AF have not been carried out. Moreover, older patients were mostly excluded from RCTs and our knowledge on this issue is very limited. Therefore, we performed a systematic review on published data dealing with AF CA through the electronic database PubMed; bibliographies of retrieved articles, review articles and textbooks were evaluated. We searched English-only articles published during the last ten years (between January 2010 and September 2020) and included those involving a group (or subgroup) of ≥ 100 elderly patients with cutoff ≥ 65 years who underwent AF CA. Twelve non-randomized articles were eligible for this review [90,94-104]. In three studies the effects of AF ablation and medical therapy were compared [90,97,104]. Blandino et al [97] investigated 412 patients aged \geq 70 years (mean age 75), admitted to two hospitals for persistent AF; 153 patients underwent radiofrequency AF ablation and 259 were treated with medications (rate or rhythm control). During a mean follow-up of 60 months, the success rate after single procedure was significantly higher in the ablation group (58% versus 43%, p=0.003) and it increased to 76% after multiple procedures, performed in 18% of patients. The periprocedural major complications rate was 6.7% and previous history of TIA/stroke was an independent predictor of postablation cerebral thromboembolic events. However, the rate of long-term adverse events was lower in the ablation group (7.7% versus 23.9%, p<0.001). Wang et al [104] investigated 1740 Chinese patients aged ≥65 years, admitted to four hospitals for AF, receiving either radiofrequency ablation or medical therapy. The propensity-matching algorithm produced 347 pairs of patients (mean age 71 years). The primary endpoint was a composite of all-cause death, non-fatal stroke and peripheral embolism and the patients in the ablation group were at significantly lower risk for these events compared with those treated conservatively (HR 0.40, 95% CI 0.19-0.85). These findings contrast with results of the large CABANA trial [90], wherein AF ablation did not reduce the primary

composite outcome of death, stroke, serious bleeding, or cardiac arrest compared with medical therapy. However, when patients were stratified according to age, patients aged <65 years (n=766) might benefit from catheter ablation (HR 0.52, 95% CI 0.27-1.00) and in those aged 65-74 years (n=1130), the two treatments had similar effects (HR 0.84, 95% CI 0.57-1.23). On the contrary, in patients aged \geq 75 years (n=308) there was a trend in favor of medical therapy (HR 1.46, 95% CI 0.80-2.67, p=0.07).

In the remaining nine studies [94-96,98-103] the effects of AF CA were investigated in older patients and in eight of these studies they were compared with those observed in younger ones. The main results are summarized in Table II. All the studies had a retrospective design. The age cutoff varied from ≥65 to ≥80 years. In eight studies the patients underwent radiofrequency ablation and in one [102] cryoballon ablation. Two studies were multicenter [101,103] and in the others only one or two centers were involved. In eight studies, success of the procedure was defined as a lack of atrial tachyarrhythmia recurrence and in two [94,100] as occurrence of infrequent relapse. In all the studies patients with symptomatic and drug refractory paroxysmal or persistent AF were enrolled. In three studies the prevalence of heart failure patients was not reported [95,101,103] and in the others it ranged between 6% and 17.8%. In one large study [103] the primary endpoint was periprocedural death, which was significantly higher in patients aged ≥ 80 years than in the younger ones (0.8% versus 0.2%, p<0.001). In the other studies the primary endpoints were periprocedural major complications rate and/or success rate. Incidence of major complications in the elderly patients enrolled in these studies ranged from 1% to 7.3%. In two studies [94100] the major complications rates increased with advancing age and in four [95,96,102,103] the differences were not statistically significant. Moser et al [101] reported a higher stroke rate in the older versus the younger cohort (1.3% versus 0.1%, p<0.01), whereas the other major complications did not statistically differ. The main predictors of major complications were kidney failure, anemia and chronic obstructive pulmonary disease [103]. The success rate was reported after both single procedure and multiple procedures in one study [96], only after single procedure in four studies [95,99,100,102] and only after multiple procedures in two studies [94,98]. After single procedure, success rates in older patients ranged from 44% to approximately 70% during a follow-up period of 18-25 months [95,96,100] and it seems to decrease to about 30% after 60 months [99]. In three studies [94,99,100] the success rates decreased with increasing age and in three [95,96,102] the differences were not

statistically significant. After multiple procedures - performed in a percentage of patients variable from 20% to 44% - the success rates in the elderly appear to be high (approximately 80%). In two studies [94,100] the success rates of ablation were reported either without and with administration of antiarrhythmicdrugs, these latter increasing the efficacy by 50-80%. However, older patients required more antiarrhythmic therapy than younger ones in order to maintain sinus rhythm. A sub-analysis carried out by Nademanee et al [98] in 75 patients with implantable devices showed that ablation markedly reduced AF burden. In two studies, good AF control was associated with better survival [98,100]. In summary, despite uneven findings among these studies carried out in elderly patients, there is some evidence that 1) periprocedural major complications rates seem to increase in the elderly, but do not appear to be much higher than in younger patients; however, a periprocedural death rate of approximately 1%, observed in a large database dealing with octogenarians, appears to be very high; 2) success rates appear to be lower in older patients than in younger counterparts; however, CA seems to be superior to medical therapy in reducing episodes of recurrent AF and in improving quality of life; 3) older patients require more antiarrhythmic drugs after ablation than the younger ones in order to maintain sinus rhythm; 4) CA does not seem to reduce the risk of stroke and other cardiovascular events, even in presence of some contrasting results; 5) comorbidities are predictive of both major complications and lower success rate.

However, the inherent limitations of observational retrospective studies do not allow us to draw clear conclusions on the indications to AF CA in older subjects for several reasons:

1) different criteria to define success of the procedure, different patient selection criteria, different follow-up duration and different presentation of the results; 2) the studies have a retrospective design, with possible selection biases and confounders; 3) most studies are single-center (tertiary referral center) and the results cannot be generalized to other institutions; 4) in most studies it is not reported how many patients were treated with antiarrhythmic drugs after ablation; 5) the prevalence of frailty was not reported, but very likely only "robust" elderly patients were selected for the invasive treatment; 6) the impact of AF ablation on the risk of disability and cognitive decline, which are important endpoints in older patients, was not adequately investigated. With regard to this latter issue, evidence of reduced incidence of dementia in mainly adult or young-older AF patients who underwent CA were reported by recent papers [105-107].

Conclusions

AF ablation appears to be a promising treatment in older subjects, but the patient selection criteria need to be better defined. In the recent European guidelines [2] this issue is not discussed. In an expert consensus statement [89], the writing group recommends that "it is reasonable to use similar indications for AF ablation in selected older people with AF as in younger patients" (Class IIa, level of evidence B), but it is not discussed how the patients should be selected.

At present, we can only speculate that CA should be considered in older "robust" patients with symptomatic, drug-refractory, paroxysmal or persistent AF, without significant heart disease and comorbid conditions, without a very dilated left atrium and a long AF history, in order to reduce symptoms and improve quality of life. Prior to undergoing CA, it is important to confirm that the patient's symptoms result from AF and to assess their severity. At present, there are no data on the effects of CA in older patients with AF and heart failure with reduced ejection fraction.

REFERENCES

- 1) Chugh SS, Havmoeller R, Narayanan K, Singh D, Rienstra M, Benjamin EJ, et al. Worldwide epidemiology of atrial fibrillation: a Global Burden of Disease 2010 Study. Circulation. 2014;129:837-47.
- 2) Hindricks G, Potpara T, Dagres N, Arbelo E, Bax JJ, Blomstrom-Lundqvist C, et al; ESC Scientific Document Group. 2020 ESC Guidelines for the diagnosis and management of atrial fibrillation developed in collaboration with the European Association of Cardio-Thoracic Surgery (EACTS) Eur Heart J 2021;42:373-498.
- 3) Go AS, Hylek EM, Phillips KA, Chang Y, Heanult LE, Selby JV, et al. Prevalence of diagnosed atrial fibrillation in adults: national implications for rhythm management and stroke prevention: the AnTicoagulation and Risk Factors in Atrial Fibrillation (ATRIA) Study. JAMA 2001;285:2370-5.
- 4) Kornej J, Börschel CS, Benjamin CJ, Schnabel RB. Epidemiology of Atrial Fibrillation in the 21st Century Novel Methods and New Insights. Circ Res 2020;127:4–20.
- 5) Ellinor PT. Introduction to the Compendium on Atrial Fibrillation, and a Few Thoughts Along The Way. Circ Res 2020;127:1–3.
- 6) Kirchhof P, Benussi S, Kotecha D, Ahlsson A, Atar D, Casadei B, et al.; ESC Scientific Document Group. 2016 ESC Guidelines for the management of atrial fibrillation developed in collaboration with EACTS. Eur Heart J 2016;37:2893–962.
- 7) Steffel J, Verhamme P, Potpara TS, Albaladejo P, Antz M, Desteghe L, et al; ESC Scientific Document Group. The 2018 European Heart Rhythm Association Practical Guide on the use of non-vitamin K antagonist oral anticoagulants in patients with atrial fibrillation. Eur Heart J 2018;39:1330-93.
- 8) Barco S, Cheung YW, Eikelboom JW, Coppens M. New oral anticoagulants in elderly patients. Best Pract Res Clin Haematol 2013;26:215-24.
- 9) Bo M, Grisoglio E, Brunetti E, Falcone Y, Marchionni N. Oral anticoagulant therapy for older patients with atrial fibrillation: a review of current evidence. Eur J Intern Med 2017;41:18-27.
- 10) Malik AH, Yandrapalli S, Aronow WS, Panza JA, Cooper HA. Meta-Analysis of Direct-Acting Oral Anticoagulants Compared With Warfarin in Patients >75 Years of Age. Am J Cardiol 2019;123:2051-7.

- 11) Camm AJ, Accetta G, Ambrosio G, Atar D, Bassand JP, Berge E, et al. Evolving antithrombotic treatment patterns for patients with newly diagnosed atrial fibrillation. Heart 2017;103:307-14.
- 12) Marzec LN, Wang J, Shah ND, Chan PS, Ting HH, Gosch KL, et al. Influence of Direct Oral Anticoagulants on Rates of Oral Anticoagulation for Atrial Fibrillation. J Am Coll Cardiol 2017;69:2475-84.
- 13) Fohtung RB, Novak E, Rich MW. Effect of New Oral Anticoagulants on Prescribing Practices for Atrial Fibrillation in Older Adults. J. Am Ger Soc. 2017;65:2405-12.
- 14) Lip GYH, Pan X, Kamble S, Kawabata H, Mardekian J, Masseria C, et al. Major bleeding risk among non-valvular atrial fibrillation patients initiated on apixaban, dabigatran, rivaroxaban or warfarin: a 'real-world' observational study in the United States. Int J Clin Pract 2016;70:752-63.
- 15) Larsen TB, Skjøth F, Nielsen PB, Kjælgaard JN, Lip GYH. Comparative effectiveness and safety of non-vitamin K antagonist oral anticoagulants and warfarin in patients with atrial fibrillation: propensity weighted nationwide cohort study. BMJ 2016;353:i3189.
- 16) Yao X, Abraham NS, Sangaralingham LR, Bellolio MF, McBane RD, Shah ND, et al. Effectiveness and Safety of Dabigatran, Rivaroxaban, and Apixaban Versus Warfarin in Nonvalvular Atrial Fibrillation. J Am Heart Assoc 2016;5:e003725.
- 17) Lip GYH, Keshishian A, Kamble S, Pan X, Mardekian J, Horblyuk R, et al. Real-world comparison of major bleeding risk among non-valvular atrial fibrillation patients initiated on apixaban, dabigatran, rivaroxaban, or warfarin. A propensity score matched analysis. Thromb Haemost 2016;116:975-86.
- 18) Noseworthy PA, Yao X, Abraham NS, Sangaralingham RL, McBane RD, Shah ND. Direct Comparison of Dabigatran, Rivaroxaban, and Apixaban for Effectiveness and Safety in Nonvalvular Atrial Fibrillation. Chest 2016;150:1302-12.
- 19) Sinnaeve PR, Brueckmann M, Clemens A, Oldgren J, Eikelboom J, Healey JS. Stroke prevention in elderly patients with atrial fibrillation: challenges for anticoagulation. J Intern Med. 2012;271:15-24.
- 20) Pugh D, Pugh J, Mead GE. Attitudes of physicians regarding anticoagulation for atrial fibrillation: a systematic review. Age Ageing 2011;40:675-83.
- 21) Fumagalli S, Potpara TS, Bjerregaard Larsen T, Haugaa KH, Dobreanu D, Proclemer A, et al. Frailty syndrome: an emerging clinical problem in the everyday

- management of clinical arrhythmias. The results of the European Heart Rhythm Association survey. Europace 2017;19:1896-902.
- Fried LP, Tangen CM, Walston J, Newman AB, Hirsch C, Gottdiener J, et al. Frailty in Older Adults: Evidence for a Phenotype. J Gerontol A Biol Sci Med Sci 2001;56:M146-M157.
- 23) Baldasseroni S, Bo M, Brambati T, Marchionni N. How much frailty is important in cardiology? G Ital Cardiol (Rome). 2019;20:210-22.
- 24) Rockwood K, Wolfson C, McDowell I. The Canadian Study of Health and Aging: organizational lessons from a national, multicenter, epidemiologic study. Int Psychogeriatr 2001;13(Suppl 1):233-7.
- 25) Rockwood K. A global clinical measure of fitness and frailty in elderly people. Can Med Assoc J 2005;173:489-95.
- 26) Pilotto A, Addante F, Franceschi M, Leandro G, Rengo G, D'Ambrosio P, et al. Multidimensional Prognostic Index based on a comprehensive geriatric assessment predicts short-term mortality in older patients with heart failure. Circ Heart Fail. 2010;3:14-20.
- 27) Pilotto A, Ferrucci L, Franceschi M, D'Ambrosio LP, Scarcelli C, Cascavilla L, et al. Development and Validation of a Multidimensional Prognostic Index for One-Year Mortality from Comprehensive Geriatric Assessment in Hospitalized Older Patients. Rejuvenation Res 2008;11:151-61.
- 28) Perera V, Bajorek BV, Matthews S, Hilmer SN. The impact of frailty on the utilisation of antithrombotic therapy in older patients with atrial fibrillation. Age Ageing 2008;38:156-62.
- 29) Wilkinson C, Todd O, Clegg A, Gale CP, Hall M. Management of atrial fibrillation for older people with frailty: a systematic review and meta-analysis. Age Ageing 2019;48:196-203.
- 30) Bo M, Marchionni N. Practical use of Direct Oral Anti Coagulants (DOACs) in the older persons with atrial fibrillation. Eur J Intern Med. 2020;71:32-8.
- 31) Rosenman MB, Simon TA, Teal E McGuire P, Nisi D, Jackson JD. Perceived or Actual Barriers to Warfarin Use in Atrial Fibrillation Based on Electronic Medical Records. Am J Ther 2012;19:330-7.
- 32) Hess PL, Mirro MJ, Diener HC, Eikelboom JW, Al-Khatib SM, Hylek EM, et al.; Atrial Fibrillation Think-Tank Participants. Addressing barriers to optimal oral

- anticoagulation use and persistence among patients with atrial fibrillation: Proceedings, Washington, DC, December 3-4, 2012. Am Heart J 2014;168:239-47.e1
- 33) Donzé J, Clair C, Hug B, Rodondi N, Waeber G, Cornuz J, et al. Risk of Falls and Major Bleeds in Patients on Oral Anticoagulation Therapy. Am J Med 2012;125:773-8.
- 34) Banerjee A, Clementy N, Haguenoer K, Fauchier L, Lip GY. Prior History of Falls and Risk of Outcomes in Atrial Fibrillation: The Loire Valley Atrial Fibrillation Project. Am J Med 2014;127:972-8.
- 35) Man-Son-Hing M, Nichol G, Lau A, Laupacis A. Choosing antithrombotic therapy for elderly patients with atrial fibrillation who are at risk for falls. Arch Intern Med 1999;159:677-85.
- 36) Singh-Manoux A, Fayosse A, Sabia S, Canonico M, Bobak M, Elbaz A, et al. Atrial fibrillation as a risk factor for cognitive decline and dementia. Eur Heart J 2017;38:2612-8.
- 37) Orkaby AR, Ozonoff A, Reisman JI, Miller DR, Zhao S, Rose AJ. Continued Use of Warfarin in Veterans with Atrial Fibrillation After Dementia Diagnosis. J Am Geriatr Soc 2017;65:249-56.
- Madhavan M, Holmes DN, Piccini JP, Ansell JE, Fonarow GC, Hylek EM, et al.; ORBIT AF Investigators. Association of frailty and cognitive impairment with benefits of oral anticoagulation in patients with atrial fibrillation. Am Heart J 2019;211:77-89.
- 39) Patti G, Lucerna M, Pecen L, Siller-Matula JM, Cavallari I, Kirchhof P, et al. Thromboembolic Risk, Bleeding Outcomes and Effect of Different Antithrombotic Strategies in Very Elderly Patients With Atrial Fibrillation: A Sub-Analysis From the PREFER in AF (PREvention of Thromboembolic Events-European Registry in Atrial Fibrillation). J Am Heart Assoc. 2017;6: e005657.
- 40) Gómez-Outes A, Lagunar-Ruíz J, Terleira-Fernández A-I, Calvo-Rojas G, Suárez-Gea ML, Vargas-Castrillón E. Causes of Death in Anticoagulated Patients With Atrial Fibrillation. J Am Coll Cardiol 2016;68:2508-21.
- 41) Piccini JP, Hammill BG, Sinner MF, Hernandez AF, Walkey AJ, Benjamin EJ, et al. Clinical course of atrial fibrillation in older adults: the importance of cardiovascular events beyond stroke. Eur Heart J. 2014;35:250-6.
- 42) Hess PL, Kim S, Fonarow GC, Thomas L, Singer DE, Freeman JV, et al.; ORBIT-AF Patients and Investigators. Absence of Oral Anticoagulation and Subsequent Outcomes Among Outpatients with Atrial Fibrillation. Am J Med 2017;130:449-56.

- 43) Rodríguez-Mañero M, López-Pardo E, Cordero A, Kredieh O, Pereira-Vazquez M, Martínez-Sande JL, et al. Clinical profile and outcomes in octogenarians with atrial fibrillation: A community-based study in a specific European health care area. Int J Cardiol 2017;243:211-5.
- Bo M, Li Puma F, Badinella Martini M, Falcone Y, Iacovino M, Grisoglio E, et al. Effects of oral anticoagulant therapy in older medical in-patients with atrial fibrillation: a prospective cohort observational study. Aging Clin Exp Res 2017;29:491-7.
- 45)Bo M, Sciarrillo I, Li Puma F, Badinella Martini M, Falcone Y, Iacovino M et al. Effects of Oral Anticoagulant Therapy in Medical Inpatients ≥65 Years With Atrial Fibrillation. Am J Cardiol 2016;117:590–5.
- 46) Bai Y, Guo S-D, Deng H, Shantsila A, Fauchier L, Ma CS, et al. Effectiveness and safety of oral anticoagulants in older patients with atrial fibrillation: a systematic review and meta-regression analysis. Age Ageing 2018;47:9-17.
- 47) Capranzano P, Miccichè E, D'Urso L, Privitera F, Tamburino C. Personalizing oral anticoagulant treatment in patients with atrial fibrillation. Expert Rev Cardiovasc Ther 2013;11:959-73.
- 48) Guo L, Li S, Wang P, Hong Y. Comparative Efficacy of Clinical Events Prevention of Five Anticoagulants in Patients With Atrial Fibrillation (A Network Meta-Analysis). Am J Cardiol 2017;119:585-93.
- 49) Mueller T, Alvarez-Madrazo S, Robertson C, Wu O, Bennie M. Comparative safety and effectiveness of direct oral anticoagulants in patients with atrial fibrillation in clinical practice in Scotland. Br J Clin Pharmacol 2019;85:422–31.
- 50) Lip GYH, Lane DA. Matching the NOAC to the Patient: Remember the Modifiable Bleeding Risk Factors. J Am Coll Cardiol 2015;66:2282-4.
- 51) Shields AM, Lip GYH. Choosing the right drug to fit the patient when selecting oral anticoagulation for stroke prevention in atrial fibrillation. J Intern Med 2015;278:1-18.
- 52) Diener H-C, Aisenberg J, Ansell J, Atar D, Breithardt G, Eikelboom J, et al. Choosing a particular oral anticoagulant and dose for stroke prevention in individual patients with non-valvular atrial fibrillation: part 1. Eur Heart J 2017;38:852-9.
- 53) Diener H-C, Aisenberg J, Ansell J, Atar D, Breithardt G, Eikelboom J, et al. Choosing a particular oral anticoagulant and dose for stroke prevention in individual patients with non-valvular atrial fibrillation: part 2. Eur Heart J 2017;38(12):860-8.

- 54) 2019 American Geriatrics Society Beers Criteria® Update Expert Panel. American Geriatrics Society 2019 Updated AGS Beers Criteria® for Potentially Inappropriate Medication Use in Older Adults. J Am Geriatr Soc 2019;67(4):674-94.
- 55) Wehling M, Collins R, Gil VM, Hanon O, Hardt R, Hoffmeister M, et al. Appropriateness of Oral Anticoagulants for the Long-Term Treatment of Atrial Fibrillation in Older People: Results of an Evidence-Based Review and International Consensus Validation Process (OAC-FORTA 2016). Drugs Aging 2017;34:499-507.
- 56) Staerk L, Gerds TA, Lip GYH, Ozenne B, Bonde AN, Lamberts M, et al. Standard and reduced doses of dabigatran, rivaroxaban and apixaban for stroke prevention in atrial fibrillation: a nationwide cohort study. J Intern Med 2018;283:45-55.
- 57) Nielsen PB, Skjøth F, Søgaard M, Kjældgaard JN, Lip GY, Larsen TB. Effectiveness and safety of reduced dose non-vitamin K antagonist oral anticoagulants and warfarin in patients with atrial fibrillation: propensity weighted nationwide cohort study. BMJ 2017;356:j510.
- 58) Ruiz Ortiz M, Muñiz J, RañaMíguez P, Roldán I, Marín F, Asunción Esteve-Pastor M, et al.; FANTASIIA study investigators Inappropriate doses of direct oral anticoagulants in real-world clinical practice: prevalence and associated factors. A subanalysis of the FANTASIIA Registry. Europace 2018;20:1577-83.
- 59) Steinberg BA, Shrader P, Thomas L, Ansell J, Fonarow GC, Gersh BJ, et al.; ORBIT-AF Investigators and Patients. Off-Label Dosing of Non-Vitamin K Antagonist Oral Anticoagulants and Adverse Outcomes: The ORBIT-AF II Registry. J Am Coll Cardiol 2016;68:2597-604.
- 60) Bo M, Corsini A, Brunetti E, Isaia G, Gibello M, Ferri N, et al. Off-label use of reduced dose direct oral factor Xa-inhibitors in subjects with atrial fibrillation: a review of clinical evidence. Eur Heart J Cardiovasc Pharmacother. 2020:pvaa103.
- 61) Di Biase L, Santangeli P, Anselmino M, Mohanty P, Salvetti I, Gili S, et al. Does the left atrial appendage morphology correlate with the risk of stroke in patients with atrial fibrillation? Results from a multicenter study. J Am Coll Cardiol. 2012;60:531-8.
- 62) Watson T, Shantsila E, Lip GY. Mechanisms of thrombogenesis in atrial fibrillation: Virchow's triad revisited. Lancet 2009;373:155–66.
- 63) Patti G, Pengo V, Marcucci R, Cirillo P, Renda G, Santilli F, et al. The left atrial appendage: from embryology to prevention of thromboembolism. Eur Heart J. 2017;38:877-87.

- 64) Khurram IM, Dewire J, Mager M, Maqbool F, Zimmerman SL, Zipunnikov V, et al. Relationship between left atrial appendage morphology and stroke in patients with atrial fibrillation. Heart Rhythm 2013;10:1843–9.
- 65) Ren JF, Callans DJ, Marchlinski FE. What is the natural relationship between left atrial appendage morphology and history of stroke? J Am Coll Cardiol 2013;61:689–90.
- 66) Piccini JP, Sievert H, Patel MR. Left atrial appendage occlusion: rationale, evidence, devices, and patient selection. Eur Heart J. 2017;38:869-76.
- 67) Handke M, Harloff A, Hetzel A, Olschewski M, Bode C, Geibel A. Left atrial appendage flow velocity as a quantitative surrogate parameter for thromboembolic risk: determinants, and relationship to spontaneous echocontrast and thrombus formation a transesophageal echocardiographic study in 500 patients with cerebral ischemia. J Am Soc Echocardiogr 2005;18:1366–72.
- 68) Al-Saady NM, Obel OA, Camm AJ. Left atrial appendage: structure, function, and role in thromboembolism. Heart 1999;82:547-54.
- 69) Mahajan R, Brooks AG, Sullivan T, Lim HS, Alasady M, Abed HS, et al. Importance of the underlying substrate in determining thrombus location in atrial fibrillation: implications for left atrial appendage closure. Heart. 2012;98:1120–6.
- 70) Vetta F, Locorotondo G, Vetta G. Anticoagulation therapy in the elderly with non-valvular atrial fibrillation: a double-edged sword. Geriatric Care, 2017;3:6371.
- 71) Schellinger PD, Tsivgoulis G, Steiner T, Köhrmannf M. Percutaneous Left Atrial Appendage Occlusion for the Prevention of Stroke in Patients with Atrial Fibrillation: Review and Critical Appraisal. Journal of Stroke 2018;20:281-91.
- 72) Holmes DR, Reddy VY, Turi ZG, Doshi SK, Sievert H, Buchbinder M, et al.; PROTECT AF Investigators. Percutaneous closure of the left atrial appendage versus warfarin therapy for prevention of stroke in patients with atrial fibrillation: a randomised non-inferiority trial. Lancet. 2009;374:534–42.
- 73) Holmes DR Jr, Kar S, Price MJ, Whisenant B, Sievert H, Doshi SK, et al. Prospective randomized evaluation of the watchman left atrial appendage closure device in patients with atrial fibrillation versus long-term warfarin therapy: the PREVAIL trial. J Am Coll Cardiol. 2014;64:1–12.
- 74) Tzikas A, Gafoor S, Meerkin D, Freixa X, Cruz-Gonzalez I, Lewalter T, et al. Left atrial appendage occlusion with the AMPLATZER Amulet device: an expert consensus step-by-step approach. EuroIntervention. 2016;11:1512–21.

- 75) Huang H, Liu Y, Xu Y, Wang Z, Li Y, Cao K, et al. Percutaneous left atrial appendage closure with the Ambre device for stroke prevention in atrial fibrillation: a prospective, multicenter clinical study. JACC Cardiovasc Interv. 2017;10:2188–94.
- 76) Turagam MK, Velagapudi P, Kar S, Holmes D, Reddy VY, Refaat MM, et al. Cardiovascular therapies targeting left atrial appendage. J Am Coll Cardiol. 2018;72:448–63.
- 77) Reddy VY, Doshi SK, Kar S, Gibson DN, Price MJ, Huber K, et al.; PREVAIL and PROTECT AF Investigators. 5-year outcomes after left atrial appendage closure: from the PREVAIL and PROTECT AF trials. J Am Coll Cardiol. 2017;70:2964–75.
- 78) Reddy VY, Möbius-Winkler S, Miller MA, Neuzil P, Schuler G, Wiebe J, et al. Left atrial appendage closure with the Watchman device in patients with a contraindication for oral anticoagulation: the ASAP study (ASA Plavix Feasibility Study With Watchman Left Atrial Appendage Closure Technology). J Am Coll Cardiol. 2013;61:2551–2556.
- 79) Boersma LV, Ince H, Kische S, Pokushalov E, Schmitz T, Schmidt B, et al.; EWOLUTION Investigators. Efficacy and safety of left atrial appendage closure with WATCHMAN in patients with or without contraindication to oral anticoagulation: 1-year follow-up outcome data of the EWOLUTION trial. Heart Rhythm. 2017;14:1302–8.
- 80) Boersma LV, Schmidt B, Betts TR, Sievert H, Tamburino C, Teiger E, et al; EWOLUTION Investigators. Implant success and safety of left atrial appendage closure with the WATCHMAN device: peri-procedural outcomes from the EWOLUTION registry. Eur Heart J. 2016;37:2465–74.
- 81) Boersma LV, Ince H, Kische S, Pokushalov E, Schmitz T, Schmidt B, et al.; following investigators and institutions participated in the EWOLUTION study. Evaluating Real-World Clinical Outcomes in Atrial Fibrillation Patients Receiving the WATCHMAN Left Atrial Appendage Closure Technology: Final 2-Year Outcome Data of the EWOLUTION Trial Focusing on History of Stroke and Hemorrhage. Circ Arrhythm Electrophysiol. 2019;12:e006841.
- 82) Dukkipati SR, Kar S, Holmes DR, Doshi SK, Swarup V, Gibson DN, et al. Device-related thrombus after left atrial appendage closure. Circulation. 2018;138:874–85.
- 83) Holmes DR Jr, Doshi SK, Kar S, Price MJ, Sanchez JM, Sievert H, et al. Left atrial appendage closure as an alternative to warfarin for stroke prevention in atrial fibrillation: a patient-level meta-analysis. J Am Coll Cardiol 2015;65:2614-23.

- 84) Price MJ, Reddy VY, Valderrábano M, Halperin JL, Gibson DN, Gordon N, et al. Bleeding outcomes after left atrial appendage closure compared with long-term warfarin: a pooled, patient-level analysis of the WATCHMAN randomized trial experience. JACC Cardiovasc Interv 2015;8:1925-32.
- 85) Osmancik P, Herman D, Neuzil P, Hala P, Taborsky M, Kala P, et al.; PRAGUE-17 Trial Investigators. Left atrial appendage closure versus direct oral anticoagulants in high-risk patients with atrial fibrillation. J Am Coll Cardiol 2020;75:3122-35.
- 86) Nasara AE, Brachmann J, Lewalter T, Akin I, Sievert H, Nienaber CA, et al. Comparison in patients < 75 years of age-versus-those > 75 years on one-year-events with atrial fibrillation and left atrial appendage occluder (from the Prospective Multicenter German LAARGE Registry). Am J Cardiol 2020;136:81-6.
- 87) Testa L, Bondi-Zoccai GGL, Dello Russo A, Bellocci F, Crea F. Rate-control vs rhythm-control in patients with atrial fibrillation: a meta-analysis. Eur Heart J 2005;26:2000-6.
- 88) Kirchhof P, Camm J, Goette A, Brandes A, Eckardt L, Elvan A, et al; EAST-AFNET 4 Trial Investigators. Early rhythm-control therapy in patients with atrial fibrillation. N Engl J Med 2020:383;1305-16.
- 89) Calkins H, Hindricks G, Cappato R, Kim YH, Saad EB, Aguinaga L, et al. 2017 HRS/EHRA/ECAS/APHRS/SOLAECE expert consensus statement on catheter and surgical ablation of atrial fibrillation: executive summary. Europace 2018;20:157-208.
- 90) Packer DL, Mark DB, Robb RA, Monahan KH, Bahnson TD, Poole JE, et al; CABANA Investigators. Effect of catheter ablation vs antiarrhythmic drug therapy on mortality, stroke, bleeding, and cardiac arrest among patients with atrial fibrillation: the CABANA randomized clinical trial. JAMA 2019;321:1261-74.
- 91) Marrouche NF, Brachmann J, Andresen D, Siebels J, Boersma L, Jordaens L, et al.; CASTLE-AF Investigators Catheter ablation for atrial fibrillation with heart failure. N Engl J Med 2018;378:417-27.
- 92) Poole JE, Bahnson TD, Monahan KH, Johnson G, Rostami H, Silverstein AP, et al.; CABANA Investigators and ECG Rhythm Core Lab Recurrence of atrial fibrillation after catheter ablation or antiarrhythmic drug therapy in the CABANA trial. J Am Coll Cardiol 2020;75:3105-18.

- 93) Noseworthy PA, Van Houten HK, Gersh BJ, Packer DL, Friedman PA, Shah ND, et al. Generalizability of the CASTLE-AF trial: catheter ablation for patients with atrial fibrillation and heart failure in routine practice. Heart Rhythm 2020;17:1057-65.
- 94) Leong-Sit P, Zado E, Callans DJ, Garcia F, Lin D, Dixit S, et al. Efficacy and risk of atrial fibrillation ablation before 45 years of age. Circ Arrhythm Electrophysiol 2010;3:452-7.
- 95) Tan HW, Wang XH, Shi HF, Yang GS, Zhou YL, Gu JN, et al. Efficacy, safety and outcome of catheter ablation for atrial fibrillation in octogenarians. Int J Cardiol 2010;145:147-8.
- 96) Santangeli P, Di Biase L, Mohanty P, Burkhardt JD, Horton R, Bai R, et al. Catheter ablation of atrial fibrillation in octogenarians: safety and outcomes. J Cardiovasc Electrophysiol 2012;23;687-93.
- 97) Blandino A, Toso E, Scaglione M, Anselmino M, Ferraris F, Sardi D, et al. Long-term efficacy and safety of two different rhythm control strategies in elderly patients with symptomatic persistent atrial fibrillation. J Cardiovasc Electrophysiol 2013;24:731-8.
- 98) Nademanee K, Amnueypol M, Lee F, Drew CM, Suwannasri W, Schwab MC, et al. Benefits and risks of catheter ablation in elderly patients with atrial fibrillation. Heart Rhythm 2015;12:44-51.
- 99) Bunch TJ, May HT, Bair TL, Jacobs J, Crandall BG, Cutler M, et al. The impact of age on 5-year outcomes after atrial fibrillation catheter ablation. J Cardiovasc Electrophysiol 2016;27:141-6.
- 100) Kautzner J, Peichl P, Sramko M, Cihak R, Aldhoon B, Wichterle D. Catheter ablation of atrial fibrillation in elderly population. J Geriatr Cardiol 2017;14:563-8.
- 101) Moser JM, Willems S, Andresen D, Brachmann J, Eckardt L, Hoffmann E, et al. Complication rates of catheter ablation of atrial fibrillation in patients aged ≥ 75 years versus < 75 years Results from the German Ablation Registry. J Cardiovasc Electrophysiol 2017;28:258–65.
- 102) Heeger CH, Bellmann B, Fink T, Bohnen JE, Wissner E, Wohlmuth P, et al. Efficacy and safety of cryoballoon ablation in the elderly: a multicenter study. Int J Cardiol 2019;278:108-13.
- 103) Romero J, Ogunbayo G, Elayi SC, Darrat Y, Rios SA, Diaz JC, et al. Safety of catheter ablation for atrial fibrillation in the octogenarian population. J Cardiovasc Electrophysiol 2019;30:2686-93.

- 104) Wang H, Du X, Guo L, Guo X, Chen Y, Xia S, et al. Ablation versus medical therapy for atrial fibrillation in the elderly: a propensity score-matched comparison. Med Sci Monit 2019;25:9875-81
- 105) Schwarz N, Kuniss M, Nedelmann M, Kaps M, Bachmann G, Neumann T, et al. Neuropsychological decline after catheter ablation of atrial fibrillation. Heart Rhythm 2010;7:1761–7.
- 106) Herm J, Schirdewan A, Koch L, Wutzler A, Fiebach JB, Endres M, et al. Impact of atrial fibrillation burden on cognitive function after left atrial ablation Results of the MACPAF study. J Clin Neurosci 2020;73:168-72.
- 107) Kim D, Yang PS, Sung JH, Jang E, Yu HT, Kim TH, et al. Less dementia after atheter ablation for atrial fibrillation: a nationwide cohort study. Eur H J 2020;41:4483-93.

Conflicts of interest.— The authors certify that there is no conflict of interest with any financial organization regarding the material discussed in the manuscript.

Authors' contributions.— All authors contributed equally to the manuscript and read and approved the final version of the manuscript.

TABLES

Table I.— Main clinical trials for left atrial appendage closure with the Watchman device.

	PROTECT-AF	PREVAIL	EWOLUTION
Study Design	Randomized	Randomized	Registry
Randomization	2:1	2:1	NA
Control	Warfarin	Warfarin	NA
Number of Patients	707	407	1025
Age, mean±s.d.	72±8.9	74±7.4	73.4±9
Aged ≥75 years, %	43	54	50.8
CHA ₂ DS ₂ -VASc score,	3.3±1.4	3.8±1.2	4.5±1.6
mean±s.d.			
Stroke/TIA, %	18	28	30.5
Implant success, %	91	95	98.5
7-days severe peri-procedural	8.7	4.2	2.7
events, %			

Abbreviations: CHA₂DS₂-VASc = congestive heart failure, hypertension, age ≥75 (doubled), diabetes mellitus, prior stroke or transient ischemic attack (doubled), vascular disease, age 65–74, female sex; EWOLUTION = Registry on WATCHMAN Outcomes in Real-Life Utilization; PREVAIL = Evaluation of the WATCHMAN Left Atrial Appendage Closure Device in Patients With Atrial Fibrillation Versus Long Term Warfarin Therapy; PROTECT-AF = Watchman Left Atrial Appendage Closure Technology for Embolic Protection in Patients With Atrial Fibrillation; s.d. = standard deviation; TIA = transient ischemic attack.

Table II – Main clinical trials for left atrial appendage closure with the Watchman device.

Author	Patient number	Mea n age (yrs)	Major complicatio ns (%)	Follo w-up (mont hs)	Success rates after single procedure (%)	Success rates after multiple procedures (%)
Leong- Sit P [94]	308 ≥ 65 yrs 570 55-64 yrs 438 45-54 yrs 232 < 45 yrs	/ / /	2.6% 2.0% 1.7% 0% p=0.01	~30	/ / / /	\$ 53% ¶ 82% \$ 65% ¶ 88% \$ 68% ¶ 88% \$ 76% ¶ 87% p<0.001 NS
Tan HV [95]	49 ≥ 80 yrs 151 70-79 yrs 177 60-69 yrs	84 75 66	2% 2.6% 1.7% NS	18	70% 72% 74% NS	/ / /
Santang eli P [96]	103 ≥ 80 yrs 2651 < 80 yrs	85 62	1% 0.9% NS	~18	69% 71% NS	87%
Nadema nee K [98]	261 ≥ 75 yrs	79	7.3%	36	/	83%
Bunch TJ [99]	46 ≥ 80 yrs 305 71-80 yrs 328 61-70 yrs 170 51-60 yrs 74 ≤ 50 yrs	83 74 66 56 43	/ / / /	60	27% 32% 43% 52% 45% p=0.01	/ / / /
Kautzne r J [100]	394 ≥ 70 yrs 2803 < 70 yrs	73 57	5.3% 3.2% p=0.03	25	§ 44% ¶ 78% § 58% ¶ 83% p=0.0001p=0.01	/ /
Moser JM [101]	227 ≥ 75 yrs 4222 < 75 yrs	77 62	*1,3% # 4.4% *0.1% # 2.7% p<0.01 NS	/	/	/
Heeger CH [102]	104 ≥ 75 yrs 104 < 75 yrs	77 63	6.7% 6.7% NS	36	59% 49% NS	/
Romero J [103]	3482 ≥ 80 yrs 82.637 < 80yrs	83 60	3.6% ф 0.8% 2.8% ф 0.2% NS p<0.001	/	/	/

COPYRIGHT© EDIZIONI MINERVA MEDICA

Abbreviations: yrs = years, NS: not significant

/ not reported, § without and ¶ with antiarrhythmic drugs administration after ablation,

*periprocedural stroke, # other major complications, \$\phi\$ periprocedural death