TRANSLATIONAL RESEARCH

In Silico TRials guide optimal stratification of ATrIal FIbrillation patients to Catheter Ablation and pharmacological medicaTION: the i-STRATIFICATION study

Albert Dasí [1]*, **Claudia Nagel** [2], **Michael T.B. Pope** [3,4],
Rohan S. Wijesurendra [3,5], **Timothy R. Betts** [3], **Rafael Sachetto** [6],
Axel Loewe [2], **Alfonso Bueno-Orovio** [1], and **Blanca Rodriguez**[1]*

[1]Department of Computer Science, University of Oxford, Wolfson Building, Parks Road, Oxford OX1 3QD, UK; [2]Institute of Biomedical Engineering, Karlsruhe Institute of Technology (KIT), Karlsruhe, Germany; [3]Department of Cardiology, Oxford University Hospitals NHS Foundation Trust, Oxford, UK; [4]Department for Human Development and Health, University of Southampton, Southampton, UK; [5]Oxford Centre for Clinical Magnetic Resonance Research, Division of Cardiovascular Medicine, Radcliffe Department of Medicine, University of Oxford, Oxford, UK; and [6]Departamento de Ciência da Computação, Universidade Federal de São João del Rei, São João del Rei, MG, Brazil

|	
Aims	Patients with persistent atrial fibrillation (AF) experience 50% recurrence despite pulmonary vein isolation (PVI), and no consensus is established for secondary treatments. The aim of our i-STRATIFICATION study is to provide evidence for stratifying patients with AF recurrence after PVI to optimal pharmacological and ablation therapies, through *in silico* trials.
Methods and results	A cohort of 800 virtual patients, with variability in atrial anatomy, electrophysiology, and tissue structure (low-voltage areas, LVAs), was developed and validated against clinical data from ionic currents to electrocardiogram. Virtual patients presenting AF post-PVI underwent 12 secondary treatments. Sustained AF developed in 522 virtual patients after PVI. Second ablation procedures involving left atrial ablation alone showed 55% efficacy, only succeeding in the small right atria (<60 mL). When additional cavo-tricuspid isthmus ablation was considered, Marshall-PLAN sufficed (66% efficacy) for the small left atria (<90 mL). For the bigger left atria, a more aggressive ablation approach was required, such as anterior mitral line (75% efficacy) or posterior wall isolation plus mitral isthmus ablation (77% efficacy). Virtual patients with LVAs greatly benefited from LVA ablation in the left and right atria (100% efficacy). Conversely, in the absence of LVAs, synergistic ablation and pharmacotherapy could terminate AF. In the absence of ablation, the patient's ionic current substrate modulated the response to antiarrhythmic drugs, being the inward currents critical for optimal stratification to amiodarone or vernakalant.
Conclusion	*In silico* trials identify optimal strategies for AF treatment based on virtual patient characteristics, evidencing the power of human modelling and simulation as a clinical assisting tool.

---

* Corresponding authors. Tel: 01865 610806. *E-mail address:* blanca.rodriguez@cs.ox.ac.uk (B.R.); *E-mail address:* albert.dasiimartinez@cs.ox.ac.uk (A.D.)

Graphical Abstract
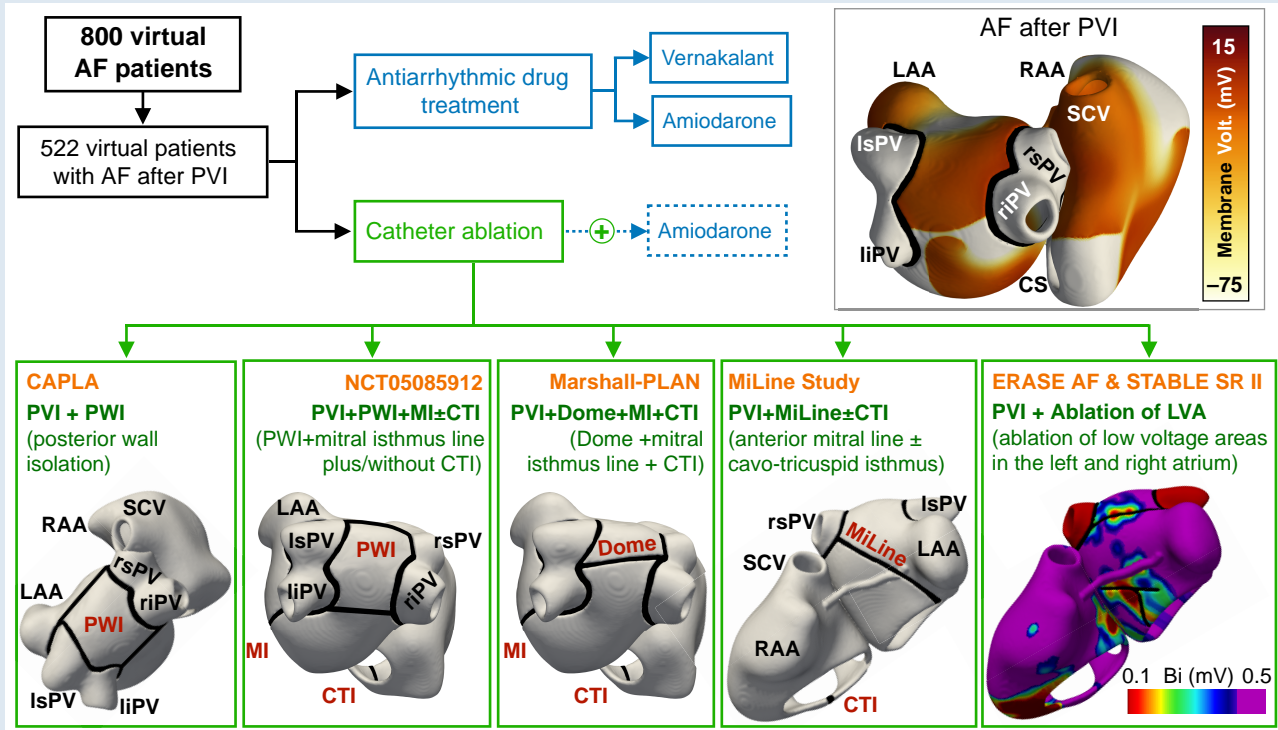
Keywords   *In silico* trials • Atrial fibrillation • Stratification • Virtual patients • Catheter ablation • Pharmacological therapy

Translational perspective

This study highlights the power of *in silico* trials based on human modelling and simulation on large virtual patient cohorts for understanding cardiac arrhythmia mechanisms and selecting appropriate therapies. Excellent agreement with clinical studies lends credibility to *in silico* trials, which identify key patient characteristics dictating treatment success and mechanisms underlying therapy failure. After appropriate validation, the proposed decision algorithm for stratifying patients with atrial fibrillation to optimal treatments could be implemented in the clinic. This methodological framework could be readily translated to any cardiac disease.

Introduction

Catheter ablation is the most effective therapy for patients with paroxysmal atrial fibrillation (AF).[1] Since most ectopic triggers originate in the pulmonary veins,[2] pulmonary vein isolation (PVI) is the cornerstone of ablation therapy. Challenges remain for persistent AF, where patients experience 50% recurrence after PVI,[3] suggesting an additional substrate for arrhythmia beyond pulmonary venous ectopic triggers. Nevertheless, since additional empirical ablation has not demonstrated incremental benefit in large clinical trials (STAR AF II[3] or CAPLA[4]), PVI remains the recommended ablation strategy at the first procedure.[5] In patients who experience AF recurrence despite

durable PVI,[6] there is no consensus regarding the optimal further treatment strategy.

The ablation of atrial low-voltage areas (LVAs; areas of bipolar voltage <0.5 mV in sinus rhythm) has shown superiority to PVI for the first time in ERASE-AF.[7] However, only 36% of patients recruited had LVAs,[7] indicating a limitation to the applicability of LVA ablation. Novel approaches, namely, MiLine[8] (anterior mitral line) and Marshall-PLAN[9] (sequential elimination of the Marshall bundles, PVI, and right and left atrial lines of block), obtained promising results in preliminary studies. Although apparently effective and approved for testing in larger trials, these strategies rely on extensive ablation lesions, increasing the risk of complications and generation of a proarrhythmic substrate. Moreover, an empirical application of these strategies is a counter to the paradigm of patient-specific ablation of pathological conduction patterns, which has proved very effective in the RECOVER AF study.[5] Thus, the ability to accurately stratify patients with AF recurrence after PVI to those requiring extensive vs. minimal substrate modification, those benefitting from empirical vs. functional ablation, and those who may be optimally treated only with additional antiarrhythmic drug therapy, would be of tremendous clinical value but remains a major challenge.

Due to their multi-scale nature and scalability, human *in silico* trials (i.e. clinical trials conducted in large cohorts of virtual patients using human-based computer modelling and simulation) can assist in identifying and explaining individual patient characteristics that guide optimal therapy selection.[10,11] The aim of our i-STRATIFICATION study is to provide evidence for the optimal stratification of patients with AF

recurrence after PVI to 12 state-of-the-art pharmacological and ablation therapies, through *in silico* trials.

Methods

The methodology used is explained in detail in the Supplementary Material, and the files used to conduct simulations are publicly available at https://zenodo.org/records/10562550.

Overview of study design

A total of 800 virtual patients with AF with durable PVI underwent a non-pulmonary vein-based AF induction protocol. The subset with sustained arrhythmia despite PVI was then subjected to 12 independent further treatments for AF (*Graphical abstract*). This served to identify subgroups of patients who, based on their clinical characteristics, were most likely to respond to a specific treatment. These clinical features were used to inform a decision algorithm for the optimal stratification of patients with AF to specific catheter ablation and/or antiarrhythmic drug therapies.

Study population

The cohort of 800 virtual patients with AF was designed to have variability in anatomy, electrophysiology, and tissue structure, based on human data and recreating the heterogeneity encountered in clinical practice (*Figure 1*). The criteria for the calibration of the population of models using experimental and clinical data are described in the Supplementary Material.

In line with STABLE-SR-II,[12] one-half of the cohort was modelled without LVAs. These 400 virtual patients with AF were developed combining 40 widely variable ionic current profiles with 10 population-representative atrial anatomies.

Anatomical variability

The atrial anatomies in the virtual cohort were selected to span volume ranges associated with AF recurrence (right and left atrial volume ranges of 80–148 and 65–112 mL, respectively[13]; Supplementary material online, *Figure S1*). Thus, 10 atrial anatomies [with right and left atrial volumes of 127 ± 51 and 105 ± 39 mL, respectively; mean $\pm$ standard deviation (SD)] were derived from a human bi-atrial statistical shape model,[14] based on 47 (40% female) clinical computed tomography (CT) and magnetic resonance imaging (MRI) data sets.[14] The anatomies used are described in Supplementary material online, *Table S1*.

Electrophysiological variability

Forty atrial cardiomyocyte profiles were identified to capture the variability in the action potential characteristics of right atrial trabeculae from 149 (37% female) patients with persistent AF.[15] The population of 40 atrial cardiomyocyte models was created by introducing variability in the ionic current densities of a baseline human atrial cellular model[16] (the scaling factors used are available in Supplementary material online, *Table S2*). Each of these 40 cardiomyocyte models, representative of the right atrial tissue, was scaled to reflect electrophysiological heterogeneities in six atrial regions, as described in Supplementary material online, *Table S3*. Together, the seven action potential models configured a unique ionic current profile (*Figure 1*). Every virtual patient presented a unique combination of ionic current profile and atrial anatomy, resulting in 400 human bi-atrial models with variability in anatomy and electrophysiology. Regional heterogeneities in conduction velocity and anisotropy were also included (see Supplementary material online, *Tables S3 and S4*), setting the baseline plane wave velocity to 80 cm/s in the bulk tissue.[11] After considering variability in the ionic current profile (e.g. $\pm 50\%$ variation in I_{Na} density), the population of virtual patients had a longitudinal conduction velocity in healthy tissue (i.e. not defined as an LVA) ranging between 72.3 and 110.9 cm/s [93.6 $\pm$ 16.5 cm/s; median $\pm$ interquartile range (IQR); see Supplementary Material for detailed explanation).

Structural variability

To develop the other half of the cohort, each of the 400 virtual patients was duplicated into a version that considered an LVA. The low-voltage area occupied at least 15% of the left atrium, as this LVA extension has been associated with AF recurrence in patients with persistent AF.[17] Moreover, the same degree of LVA was modelled in the right atrium, since structural remodelling occurs in both atria.[18]

To incorporate LVAs, the bi-atrial electro-anatomical maps of 20 (76% persistent, 41% female) patients with AF[19] were registered to the endocardial surface of the bi-atrial statistical shape model (see Supplementary Material for detailed explanation). Then, the bipolar voltage data of all 20 patients were superimposed into the same atrial geometry to build a probabilistic LVA map, i.e. identifying those atrial regions most likely to undergo LVA remodelling. These most probable areas were assigned to be LVAs until 15%[17] of the left and right atria were remodelled. Finally, the LVA distribution created in the bi-atrial statistical shape model was extended to the 10 atrial anatomies. *Figure 1* shows the 15% bi-atrial LVA extension on a representative anatomy. Low-voltage areas were simulated as regions of 30% decreased longitudinal conductivity, increased anisotropy (i.e. 8:1 longitudinal to transversal conductivity ratio), and 50, 40, and 50% reductions in I_{CaL}, I_{Na}, and I_{K1}, respectively.[11] After applying this remodelling on top of the individual electrophysiological properties of each virtual patient, the slowest possible longitudinal and transversal conduction velocities observed in the LVAs were 50.1 and 10.5 cm/s, respectively. A convergence analysis is available in Supplementary material online, *Figures S2 and S3*.

Pulmonary vein isolation and atrial fibrillation induction

To simulate AF recurrence after PVI, the pulmonary veins of virtual patients were isolated before applying the AF induction protocol. Pulmonary vein isolation was modelled as in Roney *et al.*,[20] considering a wide circumferential ablation. For consistency across different atrial anatomies, the circumferential ablation was applied at the minimum distance from the veins that enabled their complete isolation. In order to eliminate the intracellular electrotonic load and closely reproduce ablation lesions clinically, the latter were modelled *in silico* by removing the respective elements from the atrial mesh.

After the virtual application of PVI, spiral wave re-entries were imposed in the atria as the initial conditions of the simulation and AF dynamics were analysed for 7 s of activity.[11] This protocol mimicked AF initiation from non-specific triggers outside the pulmonary veins, and a detailed description of the protocol is provided in the Supplementary Material.

Simulated 12-lead electrocardiograms (ECGs) were computed in virtual patients with sustained (>7 s) AF. The ECG was computed for 30 s (i.e. the duration used clinically for AF diagnosis) to ascertain the proportion of virtual patients who, being initially defined as patients with sustained (>7 s) AF, presented uninterrupted arrhythmia over a meaningful duration in the clinics.

Intervention

Virtual patients with uninterrupted arrhythmia (i.e. >7 s of organized flutter or AF) after PVI were independently subjected to nine ablation strategies and two antiarrhythmic drugs. An additional drug was applied after ablation if arrhythmia sustained for all ablation procedures (*Graphical abstract*).

The ablation strategies considered: (i) posterior wall isolation (PWI)[4]; (ii) PWI plus mitral isthmus (MI) line (PWI + MI); (iii) PWI + MI plus cavotricuspid isthmus (CTI) line (PWI + MI + CTI); (iv) anterior mitral line (MiLine)[8]; (v) MiLine + CTI; (vi) Marshall-PLAN[9]; and (vii–ix) ablations of LVAs in three steps: only isolating LVAs in the left atrium, as in STABLE-SR-II[12] and ERASE-AF,[7] together with CTI block, and with CTI block and right atrial LVA ablation. A drug could be applied on top of the ablation procedure (i.e. synergistic drug and ablation therapy[21]): amiodarone 1.5 μM, as it is recommended for long-term rhythm control of AF.[22]

The pharmacological arm included: (i) amiodarone 3.0 μM; and (ii) vernakalant 30 μM (considered in this study as a potent Class IC agent[23]). Drug action was simulated as a simple pore-block model, considering the doses within the therapeutic plasma concentration.[11] *Table 1* shows the percentage of ionic current block at clinically relevant concentrations.

The three-dimensional monodomain equation of the transmembrane voltage and all ECG calculations were solved using the high-performance open-source software MonoAlg3D,[24] with meshes of 400 μm edge length.[25]
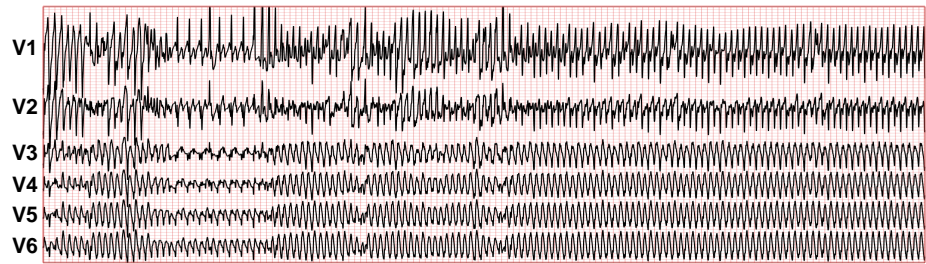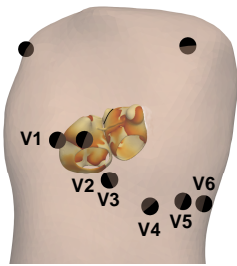
Results

Primary outcome

Sustained arrhythmia (i.e. organized flutter or AF) occurred in 522 (65%) virtual patients despite PVI (243 and 279 with the absence and presence
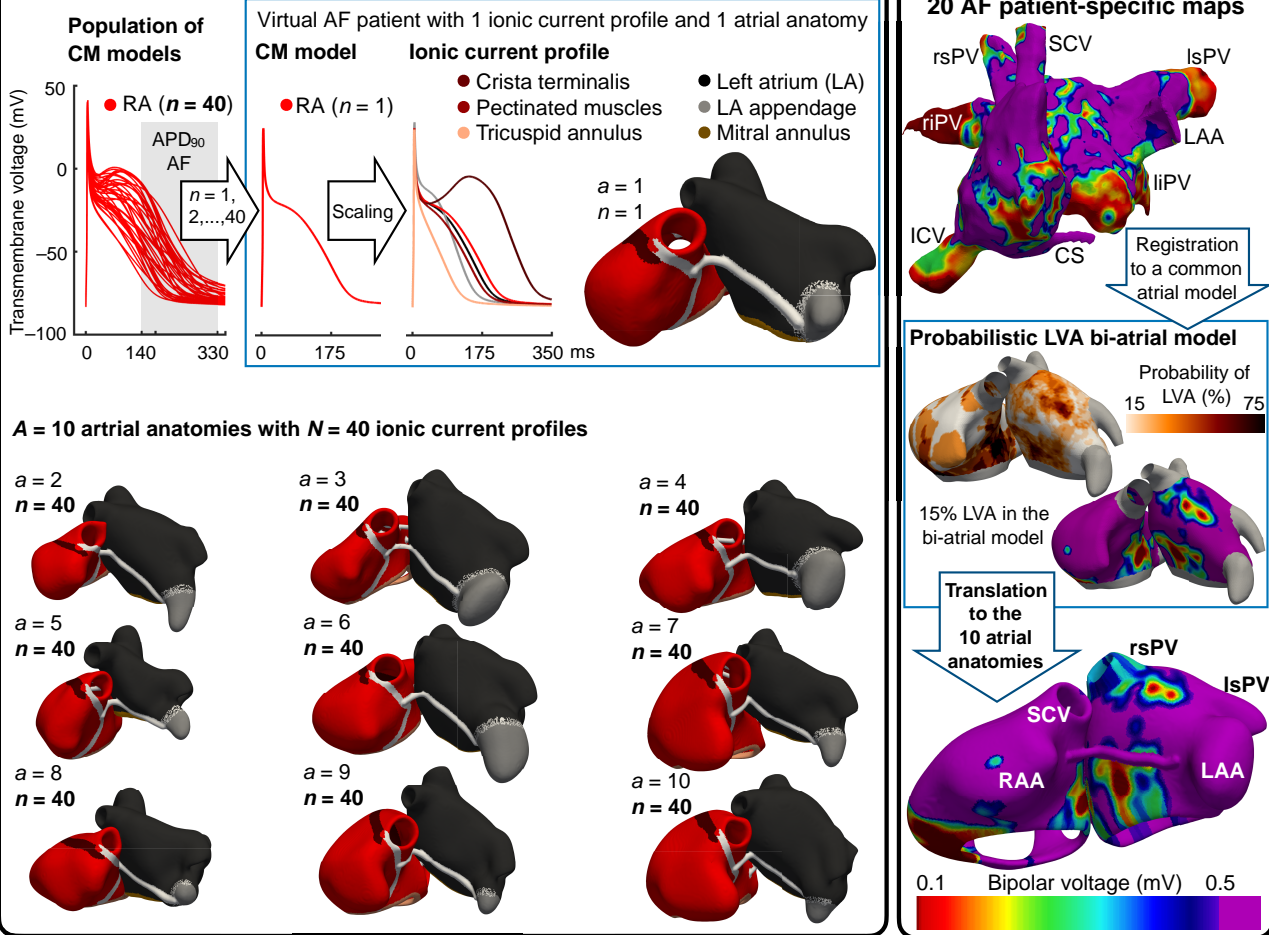
Figure 1 Construction of the cohort of 800 virtual patients. (Top) Simulated 30 s electrocardiogram during AF of a representative virtual patient (virtual atrial inside the torso). (Bottom) Population of 400 virtual patients with the absence of LVAs: 40 CM models, representative of right atrial tissue, are considered. The 40 ionic current profiles, with a consideration of regional heterogeneities, are used to populate 10 atrial anatomies. Every virtual patient presents a unique combination of ionic current profile and atrial anatomy, resulting in 400 human atria. The cohort is extended to 800 virtual patients by adding LVAs into each of the 400 atria, using a probabilistic LVA map derived from 20 patients with AF. A representative anatomy with 15% LVA extension in the right and left atria is illustrated. AF, atrial fibrillation; CM, cardiomyocyte; CS, coronary sinus; RA–LA, right and left atria; RAA–LAA, RA and LA appendage; rs–ri–ls–li–PV, right superior, right inferior, left superior, and left inferior pulmonary vein; SCV–ICV, superior and inferior cava vein.

of LVAs, respectively; the type of arrhythmia is explained below). All virtual patients with sustained arrhythmia over 7 s also presented uninterrupted arrhythmia for 30 s on the ECG (a representative example is shown in *Figure 1*). The average ECG dominant frequency was 6.9 ± 3.4 Hz, consistent with clinical measurements.[26]

These virtual patients with sustained (>7 s) AF despite PVI underwent 12 second treatments for AF, resulting in 7.064 multiscale simulations (over 50 000 h of computing time). *Table 2* summarizes the efficacy obtained *in silico* and in the clinical trial for each treatment.

Table 1 Ionic current block (%) exerted by antiarrhythmic drugs

Treatment (concentrations)	Ionic current block (%)									Ref.
	I_{Kur}	I_{Kr}	I_{to}	I_{K1}	I_{Ks}	I_{CaL}	I_{NaK}	I_{NCX}	I_{Na}	
Amiodarone (1.5 µM)	—	40	—	—	30	—	—	30	20	Dasí et al.[10]
Amiodarone (3.0 µM)	—	70	30	20	50	—	40	50	50	Dasí et al.[10]
Vernakalant (30 µM)	70	60	70	—	—	20	—	—	70	Dasí et al.,[10] Wettwer et al.[23]

Table 2 Efficacy of second strategies according to the presence of LVAs

Treatment	Number of patients free from arrhythmia after treatment, n (%)				
	In silico trials			Clinical trials/studies	
	Absence LVA (n = 243)	Presence LVA (n = 279)	Total (n = 522)	n (%)	Name
Amiodarone	160 (66)	140 (50)	**300 (57)**	65 (61)	AMIO-CAT[27]
Vernakalant	135 (56)	163 (58)	**298 (57)**	60 (52)	AVRO[28]
PWI	10 (4)	0 (0)	**10 (2)**	89 (52)	CAPLA[4]
PWI + MI	127 (52)	146 (52)	**273 (52)**	38 (76)	Ahn et al.[29]
PWI + MI + CTI	187 (77)	217 (78)	**404 (77)**	147 (74)	Li et al.[30]
				36 (80)	Davong et al.[31]
Marshall-PLAN	182 (75)	170 (61)	**352 (67)**	39 (62)	Marshall-PLAN[9]
				54 (72)	
MiLine	130 (53)	152 (54)	**282 (54)**	69 (63)	MiLine study[8]
MiLine + CTI	186 (77)	206 (74)	**392 (75)**	38 (66)	Kim et al.[32]
LVA$_{LA}$	—	**156 (56)**	—	29 (57)	ERASE-AF[7]
				46 (65)	STABLE-SR-II[12]
LVA$_{LA}$ + CTI	—	**222 (80)**	—	—	—
LVA$_{LA-RA}$ + CTI	—	**279 (100)**	—	—	—

Comparison between *in silico* and clinical trials.
The total number of virtual patients responding to a given treatment is highlighted in bold. Since LVA ablation is applied only in those patients with presence of LVA, the total number appears in the third column.
CTI, cavo-tricuspid isthmus ablation; LVA, low-voltage area; LVA$_{LA}$–LVA$_{RA}$, ablation of LVA in the left (LA) and right atrium (RA); MI, mitral isthmus ablation; MiLine, anterior mitral line; PWI, posterior wall isolation.

In the cohort of 552 virtual patients with AF after PVI, additional PWI had no incremental benefit, which is in agreement with CAPLA,[4] and arrhythmia stopped only in 2% of cases. Higher success rates were observed when PWI was applied together with MI (PWI + MI) and CTI ablation (PWI + MI + CTI): 52 and 77%, respectively. Similar efficacies were derived from the virtual application of MiLine and MiLine + CTI: 54 and 75%, which is in agreement with 63–66% reported clinically.[8,32] Virtual Marshall-PLAN prevented 67% of AF cases, comparable with 72% observed in human patients.[9] Finally, LVA ablation was applied in the 279 virtual patients with LVA remodelling, and 56, 80, and 100% efficacies were observed for LVA ablation in the left atrium (LVA$_{LA}$), together with CTI block (LVA$_{LA}$ + CTI) and LVA ablation in the right and left atria (LVA$_{LA}$ + CTI + LVA$_{RA}$), respectively. With regard to the anti-arrhythmic drug arm, both amiodarone and vernakalant had a success rate of 57%, as observed clinically.[22] The credibility of *in silico* trials is supported by the overall agreement with the efficacy reported in clinical trials.

Arrhythmia despite pulmonary vein isolation

In the absence of LVAs, 243 virtual patients had inducible arrhythmia despite PVI. In 137 (57%) patients, this was an organized atypical flutter, while in 106 (43%) patients, it was AF. An average of 2.92 ± 0.62 rotors sustained AF through a wide range of complex dynamics (11% of virtual patients showed only stationary rotors, 70% had one meandering rotor and others stationary, 19% had two or more unstable rotors, and 0% presented wave break-ups). Atrial fibrillation was facilitated by a shorter effective refractory period (ERP: 172.3 ± 13.0 vs. 230.6 ± 47.7 ms; virtual patients with AF vs. flutter, respectively; mean ± SD) and larger right atria (i.e. AF proportion increased by 0.21% per mL increase in right atrial volume). The latter enabled sustained arrhythmia in virtual patients with faster conduction velocity (95.0 ± 6.1 vs. 75.3 ± 5.7 cm/s; mean wave velocity in the bulk tissue observed in virtual patients with right atrial volume bigger vs. smaller than 90 mL, respectively).
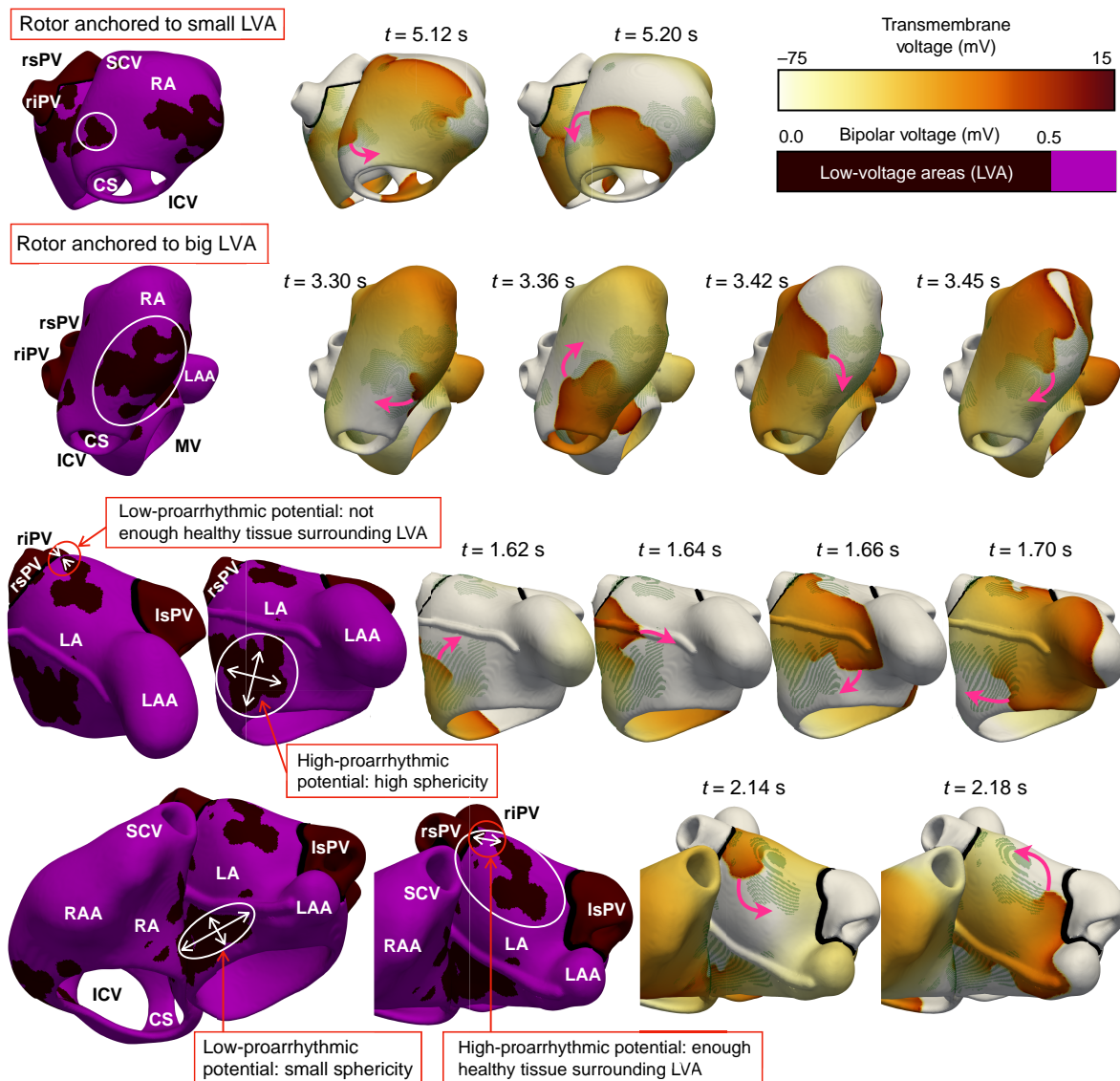
Figure 2 Proarrhythmic potential of LVAs. Re-entrant drivers anchored to smaller LVAs in virtual patients with short refractoriness and slow conduction velocity. Virtual patients with a higher conduction velocity showed rotors within bigger LVAs. Rotors anchored only to LVAs that presented enough healthy myocardium in their surroundings. Higher proarrhythmic potential derived from more spherical rather than elongated LVA shapes. CS, coronary sinus; MV–TV, mitral and tricuspid valve; LVAs, low–voltage areas; RA–LA, right and left atria; RAA–LAA, RA and LA appendage; rs–ri–ls–li–PV, right superior, right inferior, left superior, and left inferior pulmonary vein; SCV–ICV, superior and inferior cava vein.

Low-voltage area remodelling increased the proportion of virtual patients sustaining AF to 60% (169 of the 279), while the remaining 40% had organized flutter. A higher number of rotors (3.13 ± 0.95) and an increase in AF complexity (18% presented stationary rotors, 43% had one meandering rotor, 32% had two or more unstable rotors, and 7% showed wavelets or breakthroughs) owed to a higher amount of pathological conduction patterns located in LVAs. *Figure 2* illustrates a representative LVA in the left and right atria associated with lower and higher proarrhythmic potential.

Atrial fibrillation drivers (i.e. sources of electrical activity with the fastest activation time) were observed within LVAs only in virtual patients with the shortest ERP (168.5 ± 12.1 ms). When these patients additionally presented slower conduction velocity (75.3 ± 5.7 cm/s), rotors anchored to smaller LVAs (*Figure 2*, top). On the other hand, rotors were observed in bigger LVAs (*Figure 2*, middle) for virtual patients with a higher conduction velocity (95.0 ± 6.1 cm/s). Throughout the population, rotors only anchored to dense and more spherical LVAs that presented healthy myocardium in their surroundings (*Figure 2*, bottom). In virtual patients with a longer ERP (237.8 ± 46.6 ms), drivers were observed in healthy myocardium, and LVAs contributed to AF perpetuation by creating localized irregular activity (i.e. changes of wavefront direction[5]). In these patients, AF maintenance and the success of ablation therapy were determined by the right and left atrial volumes, as discussed below.
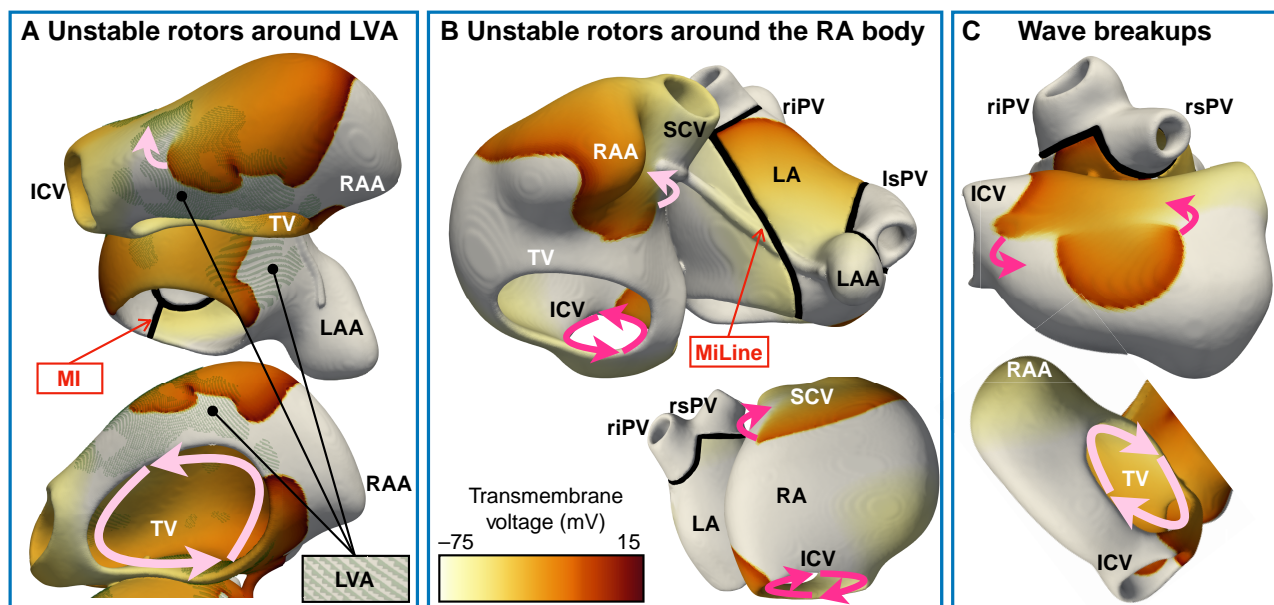
Figure 3 Snapshots of complex arrhythmias sustained by the presence of a macro-re-entrant circuit: (*A*) Unstable rotors around LVA, (*B*) Unstable rotors around the right atrial body and (*C*) wave breakups. The arrows show the direction of rotation. One arrow represents unstable rotors (i.e. rotors that vanish if the macro-re-entry is terminated), and two arrows denote a macro-re-entrant circuit. MI, mitral isthmus line of block; MiLine, anterior mitral line of block; RA–LA, right and left atria; RAA–LAA, RA and LA appendage; rs–ri–ls–li–PV, right superior, right inferior, left superior, and left inferior pulmonary vein; SCV–ICV, superior and inferior cava vein; TV, tricuspid valve.

Catheter ablation as second procedure treatment

The success of second ablation procedures was determined by the extent of bi-atrial enlargement and the presence of LVAs. Thus, positive ablation outcomes derived from targeting anatomic and functional structures, not only in the left but also in the right atrium. Indeed, the importance of additional right atrial ablation is evidenced by comparing the efficacy of strategies that considered CTI ablation (*Table 2*).

Additional benefit of cavo-tricuspid isthmus block

Fifty per cent of virtual patients presented sustained arrhythmia when only left atrial ablation was considered, regardless of the ablation extent (52, 54, and 56% arrhythmia freedom after PWI + MI, MiLine, and LVA$_{LA}$ ablation, respectively). As explained below, small differences in the efficacy of these strategies regarded whether drivers could still arise in the left atrium post-ablation. Patients with sustained arrhythmia after left atrial ablation were characterized by bigger right atria: no arrhythmia was observed for right atrial volumes <60 mL, and a 40% probability of sustained arrhythmia arose for volumes >100 mL. These patients additionally presented a short ERP (172.3 $\pm$ 13.0 ms), so that right atrial drivers could still perpetuate AF after extensive left atrial ablation.

On the other hand, 67–80% efficacy derived from additionally considering CTI block (67, 77, 75, and 80% arrhythmia freedom after Marshall-PLAN, PWI + MI + CTI, MiLine + CTI, and LVA$_{LA}$ + CTI, respectively). Noteworthy, only 10% of virtual patients where CTI block was successful had originally flutter alone. The other 90% showed a stable macro-re-entry in the right atrium, that potentiated other forms of complex arrhythmias such as unstable rotors anchored to LVAs (*Figure 3A*), meandering throughout the right atrial body (*Figure 3B*) or breaking into wavelets (*Figure 3C*). When the macro-re-entry was eliminated through CTI block, unstable rotors spontaneously vanished.

When CTI block was applied, the success of left atrial ablation depended on the ablation strategy considered, atrial ERP, and presence of LVAs. *Figure 4A–D* illustrates representative examples of unsuccessful ablation for the ablation strategies that considered CTI block. *Figure 4E* shows the number of virtual patients with left vs. right atrial drivers after extensive left atrial ablation. *Figure 4F* separates the latter between drivers anchored to LVAs and present in healthy myocardium of the right atrium.

Marshall-PLAN

Marshall-PLAN had a 75% efficacy in virtual patients with the absence of LVAs (*Table 2*). The lines of block considered by Marshall-PLAN (i.e. dome and MI) prevented all arrhythmias in virtual patients with a left atrial volume <90 mL. However, since no ablation lines were applied in the roof or anterior wall, where rotors preferentially located (*Figure 4Ai* and *Aii*), the proportion of AF increased by 0.15% per mL increase in left atrial volume. The efficacy of Marshall-PLAN worsened with LVA presence (61%, *Table 2*), since more virtual patients presented drivers anchored to LVAs in the left atrial anterior wall (*Figure 4Aii*). In total, 24 and 69 virtual patients with the absence and presence of LVAs, respectively, presented drivers in the left atrium.

MiLine + cavo-tricuspid isthmus

Compared with Marshall-PLAN, anterior mitral line prevented the formation of rotors in the anterior wall of the left atrium for both patients with the absence and presence of LVAs. However, after applying MiLine, 6 and 24 virtual patients with the absence and presence of LVAs, respectively, still presented left atrial drivers. The six virtual patients with the absence of LVAs had a left atrial volume >140 mL, which facilitated rotor anchoring in the posterior wall (*Figure 4Bi*). The 24 virtual patients with LVAs had varying left atrial sizes (100.5 $\pm$ 54.5 mL), but all of them presented a very short ERP (168.5 $\pm$ 12.1 ms). Rotors also appeared in the posterior wall, anchored to small LVAs (*Figure 4Bii*). For a longer ERP, MiLine was very effective even for large left atria.
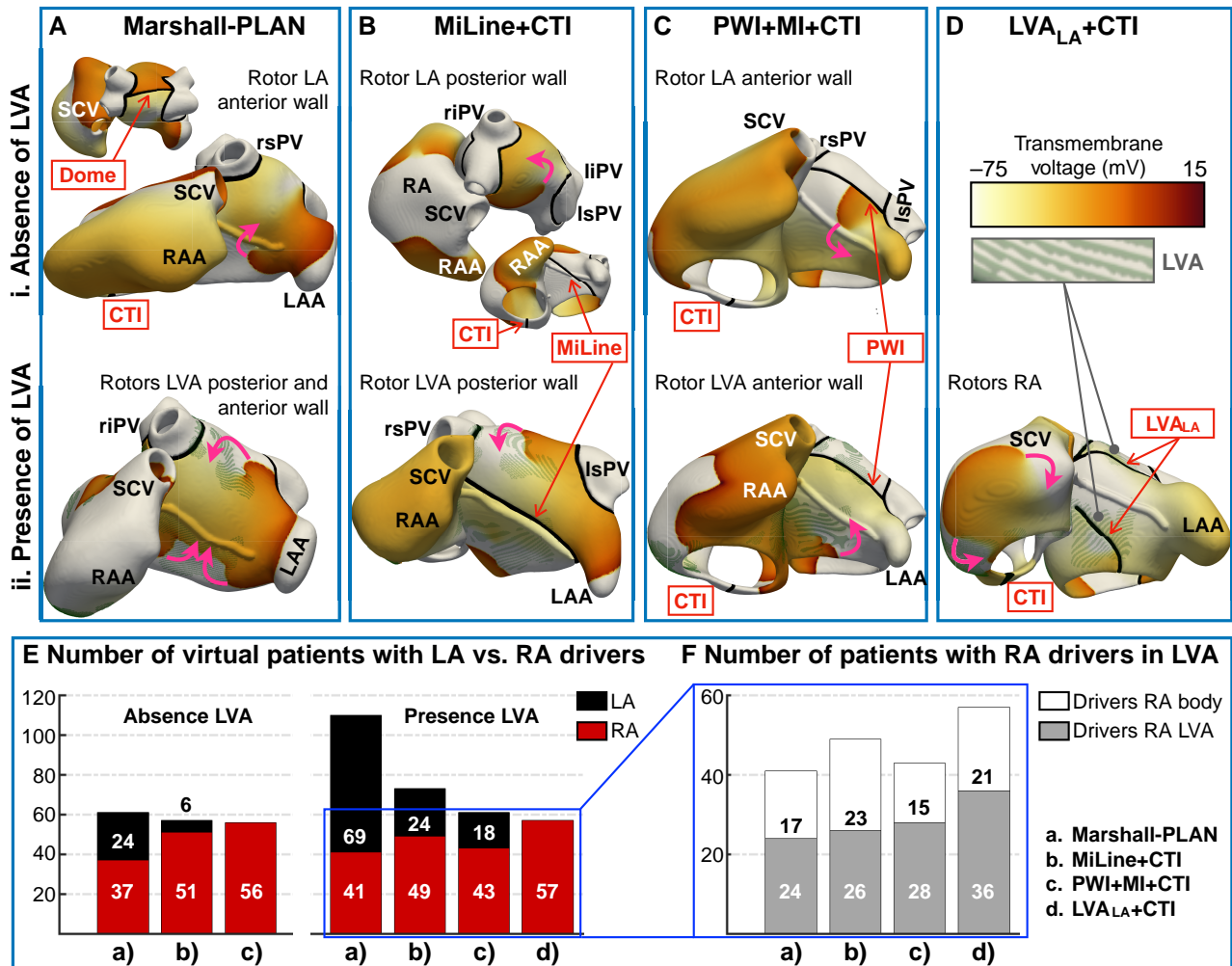
Figure 4 Examples of unsuccessful ablation for the strategies considering CTI block. (*A*) Re-entrant drivers (the arrows show the direction of rotation) in the anterior and posterior wall of the left atrium after Marshall-PLAN. (*B*) Rotors in the posterior wall after anterior mitral line (MiLine). (*C*) Drivers in the anterior wall after PWI and MI ablation (PWI + MI). (*D*) Ablation of LVAs in the left atrium (LVA$_{LA}$) prevents left atrial drivers, but right atrial drivers may exist. (*E*) Number of virtual patients presenting left atrial vs. right atrial drivers after extensive left atrial ablation. (*F*) Number of virtual patients with right atrial drivers in healthy myocardium vs. anchored to LVAs. CTI, cavo-tricuspid isthmus; LVA, low-voltage area; MI, mitral isthmus; PWI, posterior wall isolation; RA–LA, right and left atria; RAA–LAA, RA and LA appendage; rs–ri–ls–li–PV, right superior, right inferior, left superior, and left inferior pulmonary vein; SCV–ICV, superior and inferior cava vein.

Posterior wall isolation + mitral isthmus + cavo-tricuspid isthmus

When applied with CTI block, PWI plus MI ablation had a similar efficacy in virtual patients with the absence and presence of LVAs: 77 and 78%, respectively. In the absence of LVAs, drivers were mainly located in the right atrium, but unstable rotors appeared in the anterior wall of the left atrium (*Figure 4Ci*). In this subgroup, the proportion of AF increased with increasing right atrial volume (0.16%/mL). Conversely, in the subgroup of virtual patients with LVAs, 18 presented stable left atrial drivers in the anterior wall (*Figure 4Cii*).

Low-voltage area ablation in the left atrium + cavo-tricuspid isthmus

Ablation of LVAs prevented all left atrial drivers, showing the highest success rate in the subgroup of virtual patients with LVAs. Fifty-seven

patients still had sustained arrhythmia, due to right atrial drivers (*Figure 4Dii* and *E*).

Arrhythmia despite left atrial ablation and cavo-tricuspid isthmus block

Despite extensive left atrial ablation and CTI block, these four strategies had an efficacy of 67–80%, since drivers could still arise primarily in the right atrium (*Figure 4E*). In the subgroup of patients with the presence of LVAs, drivers were mostly anchored to the LVAs (*Figure 4F*). Furthermore, even when drivers located in healthy myocardium of the right atrium, LVAs helped perpetuating the arrhythmia. Consequently, a prevention of 100% was observed after ablating LVAs in the right and left atria plus additional CTI block.

In the subgroup with the absence of LVAs, 56–61 virtual patients presented right atrial drivers despite left atrial ablation and CTI block (*Figure 4E*). In this subgroup, no further ablation was considered, since
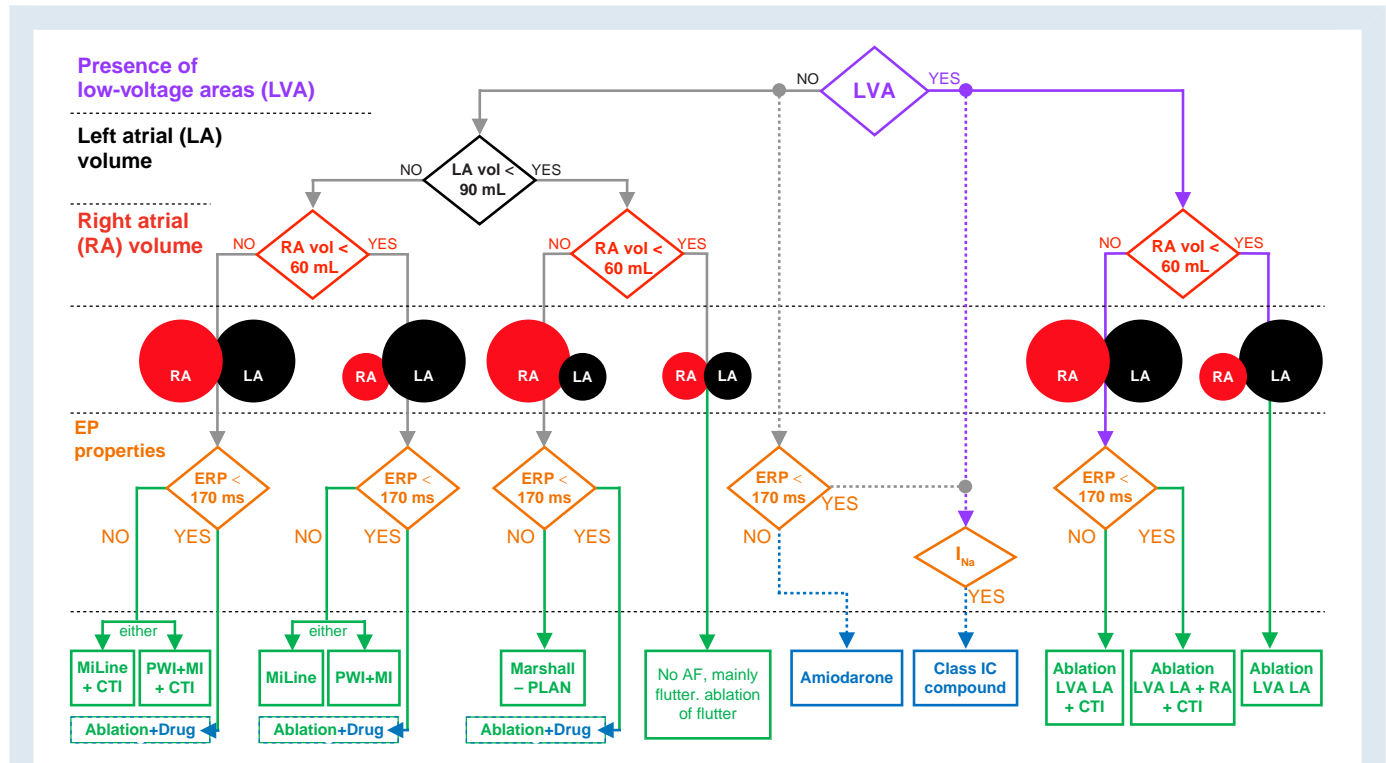
Figure 5 Decision algorithm for optimal stratification of persistent atrial fibrillation patients to catheter ablation or antiarrhythmic drug therapy. Pharmacological treatment is presented with dashed lines, since its selection is also influenced by potential concomitant patient's heart disease.[22] Abbreviations as in *Table 2*.

unstable rotors meandered throughout the right atrial body. Conversely, additional pharmacological treatment successfully stopped the arrhythmias. Low-dose amiodarone (i.e. 40% I_{Kr} inhibition) was successful in virtual patients with the absence of LVAs when applied on top of the ablation.

Antiarrhythmic drug as second procedure treatment

When antiarrhythmic drugs were virtually tested post-PVI in the absence of further ablation procedures, the ionic current substrate of the atria determined the response to pharmacological treatment.

Amiodarone

Amiodarone prevented all arrhythmia in atria presenting a longer ERP (230.7 ± 51.4 ms), resulting from higher I_{CaL} density (0.13 ± 0.08 vs. 0.08 ± 0.02 S/mF; responders vs. non-responders; median $\pm$ IQR). For a shorter ERP, the success rate of amiodarone was also influenced by the structural substrate (i.e. atrial volume and LVA infiltration). Low-voltage area infiltration hampered amiodarone efficacy (66 and 50% for the absence and presence of LVAs, respectively). Moreover, 88% (73/83) and 82% (114/139) of non-responders with the absence and presence of LVAs, respectively, were characterized by a bi-atrial volume >200 mL. Thus, amiodarone was less efficacious in virtual patients with a short ERP, big atria, and the presence of LVAs.

Vernakalant

Vernakalant increased post-repolarization refractoriness through I_{Na} block, and thus, virtual patients with I_{Na} up-regulation were less likely to respond to vernakalant (5.2 ± 1.8 vs. 9.1 ± 1.8 S/mF; responders vs. non-responders). Unlike amiodarone, the efficacy of vernakalant

was not influenced by atrial size or LVA infiltration (56 and 58% efficacy for the absence and presence of LVAs, respectively), but by the excitability (i.e. I_{Na} expression) of virtual patients with AF.

Key patient characteristics and proposed decision algorithm

In an attempt to rationalize the findings of our large-scale simulation study, *Figure 5* illustrates a possible decision algorithm for optimal stratification to pharmacological treatment and catheter ablation strategies based on the assessment of patient characteristics. This algorithm can be evaluated and updated with additional clinical evidence in successive iterations. Based on the simulation results, we propose that the algorithm first evaluates the presence of LVAs (i.e. *Figure 5*, right arm of the algorithm), since their ablation yielded the highest prevention efficacy. Virtual patients with LVAs and smaller right atria (<60 mL) benefited from LVA ablation in the left atria alone. Conversely, right atrial ablation was required for bigger right atria. Virtual patients with a right atrial volume above 60 mL and a short ERP (<170 ms) were optimally treated with LVA ablation in both the left and right atria, with additional CTI block to prevent flutter-derived more complex arrhythmias (*Figure 3*). In patients with big right atria but a longer ERP, CTI block and LVA ablation in the left atrium were sufficient to stop AF maintenance.

In the absence of LVAs (i.e. *Figure 5*, left arm of the algorithm), the optimal empirical ablation strategy was also dictated by the atrial volume. Atrial flutter was observed in virtual patients with small right (<60 mL) and left atria (<90 mL). Thus, ablation attempted to eliminate the specific re-entrant circuit. For bigger right atria (>60 mL), additional CTI ablation improved AF prevention. Accordingly, within the strategies considering CTI block, Marshall-PLAN was sufficient for small

left atria (<90 mL), but a more extensive ablation, such as MiLine + CTI or PWI + MI + CTI, was required as the left atrial volume increased. In the cases of big left atria and small right atria (<60 mL), the latter strategies without CTI line (i.e. MiLine and PWI + MI) succeed in preventing AF maintenance.

Noteworthily, virtual patients with the absence of LVAs but a short ERP (<170 ms) required synergistic pharmacological (i.e. low-dose amiodarone) and ablation therapy. The optimal stratification to ablation strategies was still dependent on the atrial volumes, as with virtual patients with a long ERP, but additional antiarrhythmic drugs were needed to prevent AF maintenance.

When administered without catheter ablation, the efficacy of pharmacotherapy was determined by the patient's electrophysiological profile, being the inward currents (i.e. I_{Na} and I_{CaL}) critical for optimal selection of amiodarone or Class IC agents (i.e. vernakalant). Due to the current inability to characterize the patient's ionic current profile on the fly, the stratification to antiarrhythmic drugs in *Figure 5* has been represented with dashed lines (see *Discussion*).

Discussion

In this large-scale simulation study, AF maintenance was evaluated in a population of 800 virtual patients presenting variability in anatomy, electrophysiology, and tissue structure. In this cohort, 522 virtual patients had inducible arrhythmia after PVI and were subjected to *in silico* trials with ablation and pharmacological treatments (7.064 multi-scale simulations obtained through >50 000 h of computing time). Accordingly, we have demonstrated overall agreement between simulated and clinical results, supporting the credibility of the *in silico* trials. Moreover, we have identified specific patient characteristics that dictate treatment success and provided mechanistic explanations for treatment failure. The digital evidence obtained from the *in silico* trials has informed a possible decision algorithm to guide patient stratification to optimal AF treatments. This methodological framework can be probed and updated with further clinical data, to evaluate further treatments for AF, or for other cardiac diseases, highlighting a route for direct translation of human modelling and simulation into clinical practice.

In silico trials using modelling and simulation for optimal atrial fibrillation treatment

Computer modelling and simulation have been widely used for testing optimal ablation therapies of AF[33] in virtual cohorts. Early work by Hwang *et al.*[34] showed the feasibility of applying virtual ablation in a cohort of 20 virtual atria reconstructed from CT images of 20 (80% persistent) patients with AF. The authors virtually tested three empirical ablations strategies and, in agreement with our study, concluded that PVI applied together with roof, floor, and anterior mitral line had the highest antiarrhythmic efficacy. This work preceded a prospective clinical trial by the same group,[35] in which 108 (78% persistent) patients with AF were randomly assigned to receive *in silico* guided or standard-of-care ablation. In the *in silico* ablation group, the patient's left atrium was reconstructed and five ablation strategies were assessed. The strategy yielding the fastest AF termination *in silico* was selected as optimal therapy and applied in the actual patient. The study obtained non-inferiority of *in silico*-guided ablation and showcased *in silico* trials for guiding the selection of optimal AF therapies.

Further simulation studies[36,37] included personalization of both the left atrial geometry and the patient-specific fibrotic distribution. Thus, the authors additionally assessed virtual ablation of structurally remodelled substrate (i.e. LVAs or fibrosis derived from late gadolinium-enhancement MRI), which showed higher success rates than empirical ablation. Both studies demonstrated that targeting AF-perpetuating areas (i.e. drivers), yielded the greatest freedom from arrhythmia (i.e. 46[37] and 40–80%[36]). In our study, we have also observed the importance of LVA ablation for driver termination. However, after complete isolation of left atrial drivers, we found that right atrial drivers sustained AF in ~40% of cases, not considered in the above-mentioned studies. The importance of right atrial ablation has been illustrated by Roney *et al.*,[38] in which bi-atrial ablation of fibrosis proved superior to left atrial ablation alone. This was further supported by Boyle *et al.*,[39] in which the optimal ablation of drivers in both atrial chambers is now being tested in a clinical trial (NCT04101539). In this study, 100 patients with AF are expected to undergo simulation-driven ablation, compared with the 800 virtual patients included in our pilot study.

While all these works aimed to develop 'digital twins' of the atria for precision medicine, they included no variability in atrial electrophysiology. Failing to consider accurate tissue repolarization properties might lead to believe (among others) that re-entrant AF drivers exclusively localize within regions of structurally remodelled tissue.[36] In our study, variability in ionic current profiles was considered, resulting in a whole spectrum of AF dynamics (i.e. stable re-entrant drivers anchored to LVAs, wave break-ups, unstable and meandering rotors, etc.) for the same atrial anatomy and LVA distribution. As shown in this population-based study, including variability in atrial electrophysiological properties is crucial for both accurately phenotyping AF dynamics and properly stratifying patients to optimal ablation and, specifically, pharmacological therapies.[10,11]

The latter concept, patient stratification, is another aspect largely disregarded in both simulation and clinical studies. The 'one-size fits all' approach that is currently adopted for rhythm control of AF,[40] and the search for one best treatment, might explain the high recurrence rates reported in patients with persistent AF. Conversely, we have shown that the efficacy of AF therapy could be improved if optimal patient stratification, based on key patient characteristics, is considered.

Efficacy of *in silico* vs. clinical trials

Overall agreement between simulated and clinical results was observed, supporting credibility of the *in silico* trials. This is important since the results of clinical trials might be confounded by discontinuous monitoring, yielding inflated success, or lack of lesion durability, leading to treatment failure due to gaps in lines rather than ineffective ablation strategy. In this sense, while simulated AF was assessed for 7 s (see *Limitations*), a 100% lesion effectiveness was guaranteed *in silico*. Thus, small differences were observed between clinical and simulated results due to procedural differences. For example, most clinical studies allowed concomitant antiarrhythmic drug use or considered first procedure treatments. Therefore, the proposed strategy comprised the efficacy derived from the elimination of pulmonary vein triggers (PVI), plus the added benefit of further lines of block (PVI+). Conversely, our *in silico* trials considered second procedure treatments, in which non-pulmonary vein triggers were imposed in every virtual patient. Thus, PVI had no additional benefit, which explained the lower efficacies generally observed *in silico*.

This was the case for the virtual application of PWI. A 52% efficacy was observed in patients with persistent AF treated with PWI in CAPLA,[4] when compared with 2% obtained in our study. The posterior wall is believed to be a focus of triggered activity[4] and thus, the high success of PVI + PWI might be attributed to a complete elimination of ectopy. In our study, we mimicked a situation in which triggers arose everywhere in the atria, which could explain the low efficacy of simulated PWI. Moreover, additional lines of block, such as CTI ablation,[29] are often performed in clinical trials evaluating PWI efficacy, which hampers a direct comparison against *in silico* PWI. In agreement with these clinical studies,[29,41] we also observed a high proportion (51%) of virtual patients with atrial flutter after PWI that benefited from additional ablation lines.

Therefore, an important increase in the success rate derived from the virtual application of PWI together with MI ablation. Efficacies of 73.5 and 80% have been reported clinically after PWI + MI in a retrospective[30] and a small prospective study.[31] In the former study,[30] 30 (15%) patients with persistent AF treated with PWI + MI additionally received CTI ablation, and in the latter study,[31] CTI block was applied in 6 (13%) patients. Consistent with the clinical results, we observed 77% efficacy when PWI + MI + CTI was considered.

A similar success rate (75%) derived from the virtual application of *anterior mitral line* and CTI ablation. When MiLine was applied alone, 282 (54%) virtual patients were free of arrhythmia. While a higher efficacy (62.7%) was observed clinically in a cohort of 110 patients with persistent AF,[8] 59 of them were treated with additional lines of block (50 with PWI and 9 roof line). Moreover, the patients in this study[8] were carefully selected, so that MiLine was only applied in those presenting large LVAs in the left atrial anterior wall. In this case, MiLine regarded a patient-tailored approach,[8] rather than an empirical ablation. Consistent with the clinical study, we observed the absence of AF drivers anchored to LVAs in the left atrial anterior wall after the virtual application of MiLine.

In this sense, a complete elimination of LVA-dependent drivers derived from *LVA ablation* in both atrial chambers. When LVA ablation was applied only in the left atrium, we observed a 56% success rate, consistent with ERASE-AF.[7] In the latter, from 51 patients with the presence of LVAs who underwent LVA ablation, 29 (57%) were free from arrhythmia. A slightly higher efficacy (65%) was observed in STABLE-SR-II,[12] where recurrence occurred in 25 out of 71 patients with the presence of LVAs. This might be attributed to the fact that not all LVA harbour proarrhythmic potential, as shown in this study, and PVI might be enough for preventing arrhythmia.

A slightly higher efficacy was also observed for Marshall-PLAN in a prospective study[9] compared with our simulations (72 vs. 67%, respectively). The ablation set of Marshall-PLAN was the hardest to reproduce, since the vein of Marshall ethanol infusion could not be accurately modelled *in silico*. In VENUS,[42] adding vein of Marshall ethanol infusion proved to reduce recurrence compared with PVI alone, which could explain the slightly lower efficacy of Marshall-PLAN observed *in silico*. Nevertheless, since we incorporated MIs ablation, comparable efficacies of 67 and 72% derived from our simulations and clinically.[9] In the prospective study, 79% freedom from arrhythmia was observed in 68 patients presenting a complete lesion set, consistent with the efficacy that we observed in virtual patients with the absence of LVAs (75%). No subgroup analysis or proportion of LVA is reported in the clinical study for comparison with our simulations.

Likewise for pharmacological therapy, the success rate obtained after virtual administration of amiodarone post-PVI (57%) agrees with the 61% (65/107) reported in AMIO-CAT,[27] a randomized clinical trial assessing whether the short-term use of amiodarone after catheter ablation could prevent early recurrence. The slightly higher efficacy observed in the clinical study[27] might be due to the fact that PVI only was performed in 75% of patients, with the other 25% receiving additional ablation lines. Overall, the excellent agreement between *in silico* and clinical trials is improved when the different procedural aspects are also taken into account.

Limitations

This study proposes an *in silico* trial framework, applied to study 12 AF treatments in a large population of 800 virtual patients. Thus, a compromise was reached between the total number of simulations (i.e. over 7.000) and the time per simulation (i.e. 7 s). However, it is unclear whether 7 s of simulated AF is representative of the follow-up time reported in clinical trials, and thus, unclear whether it could be used to predict AF recurrence clinically. Similarly, for computational feasibility,

we have used a mesh resolution of 400 μm. However, in extreme scenarios, modelling areas of very slow conduction velocity (e.g. LVAs) might require a finer element length to accurately resolve transverse conduction speeds (see Supplementary material online, *Figures S2 and S3*). While we have shown that 400 μm yields similar re-entry dynamics than finer spatial resolutions, it is important to consider that numerical-related uncertainties might still arise in those extreme cases.

Furthermore, we propose a decision algorithm in an attempt to rationalize the large amount of data generated and initiate the process of clinical translation. In its current version, the proposed decision algorithm for patient stratification largely relies on invasive electroanatomical mapping, even when the algorithm suggests the use of antiarrhythmic drugs without ablation. In this sense, the selection of pharmacological treatment would ideally be guided by non-invasive procedures, even if a redo procedure is common in patients with AF recurrence after PVI. Similarly, the characterization of the patient's ionic current profile is not routinely performed in the clinic, due to multiple ethical and procedural reasons. This would impede optimal pharmacological treatment selection according to the *in silico* results, and thus, other non-invasive biomarkers need to be identified to guide optimal pharmacological therapy. Lastly, while we calibrated and constructed our virtual atria by considering at all stages both male- and female-related data, future studies could aim at extending our approach to sex-specific decision algorithms.

Conclusions

This population-based study highlights the power of *in silico* trials based on human modelling and simulation for selecting and understanding optimal cardiac therapies. Simulations demonstrate agreement with clinical results in terms of treatment efficacy and hold the advantages of identifying specific patient characteristics that dictate treatment success and providing mechanistic explanations for treatment failure. A decision support system is build based on this evidence to guide patient stratification to optimal AF treatments. The validation of the methodological framework and the decision algorithm with further clinical data could boost the efficacy of AF treatment and reinforce the integration of *in silico* trials into clinical practice.

Supplementary material

Supplementary material is available at *Europace* online.

Conflict of interest: none declared.

Data availability

The original contributions presented in the study are included in the article, and further inquiries can be directed to the corresponding authors.

References

1. Wazni OM, Dandamudi G, Sood N, Hoyt R, Tyler J, Durrani S *et al.* Cryoballoon ablation as initial therapy for atrial fibrillation. *N Engl J Med* 2021;**384**:316–24.
2. Haïssaguerre M, Jaïs P, Shah DC, Takahashi A, Hocini M, Quiniou G *et al.* Spontaneous initiation of atrial fibrillation by ectopic beats originating in the pulmonary veins. *N Engl J Med* 1998;**339**:659–66.
3. Verma A, Jiang C, Betts TR, Chen J, Deisenhofer I, Mantovan R *et al.* Approaches to catheter ablation for persistent atrial fibrillation. *N Engl J Med* 2015;**372**: 1812–22.
4. Kistler PM, Chieng D, Sugumar H, Ling LH, Segan L, Azzopardi S *et al.* Effect of catheter ablation using pulmonary vein isolation with vs without posterior left atrial wall isolation on atrial arrhythmia recurrence in patients with persistent atrial fibrillation: the CAPLA randomized clinical trial. *JAMA* 2023;**329**:127–35.
5. Betts TR, Good WW, Melki L, Metzner A, Grace A, Verma A *et al.* Treatment of pathophysiologic propagation outside of the pulmonary veins in retreatment of atrial fibrillation patients: RECOVER AF study. *Europace* 2023;**25**:euad097.
6. Benali K, Barré V, Hermida A, Galand V, Milhem A, Philibert S *et al.* Recurrences of atrial fibrillation despite durable pulmonary vein isolation: the PARTY-PVI study. *Circ Arrhythm Electrophysiol* 2023;**16**:e011354.
7. Huo Y, Gaspar T, Schönbauer R, Wójcik M, Fiedler L, Roithinger FX *et al.* Low-voltage myocardium-guided ablation trial of persistent atrial fibrillation. *NEJM Evidence* 2022;**1**: EVIDoa2200141.
8. Bergonti M, Spera FR, Ferrero TG, Nsahlai M, Bonomi A, Boris W *et al.* Anterior mitral line in patients with persistent atrial fibrillation and anterior scar: a multicenter matched comparison—the MiLine study. *Heart Rhythm* 2023;**20**:658–65.
9. Derval N, Duchateau J, Denis A, Ramirez FD, Mahida S, André C *et al.* Marshall bundle elimination, pulmonary vein isolation, and line completion for ANatomical ablation of persistent atrial fibrillation (Marshall-PLAN): prospective, single-center study. *Heart Rhythm* 2021;**18**:529–37.
10. Dasí A, Roy A, Sachetto R, Camps J, Bueno-Orovio A, Rodriguez B. In-silico drug trials for precision medicine in atrial fibrillation: from ionic mechanisms to electrocardiogram-based predictions in structurally-healthy human atria. *Front Physiol* 2022;**13**:966046.
11. Dasí A, Pope MTB, Wijesurendra RS, Betts TR, Sachetto R, Bueno-Orovio A *et al.* What determines the optimal pharmacological treatment of atrial fibrillation? Insights from in silico trials in 800 virtual atria. *J Physiol (Lond)* 2023;**601**:4013–32.
12. Yang G, Zheng L, Jiang C, Fan J, Liu X, Zhan X *et al.* Circumferential pulmonary vein isolation plus low-voltage area modification in persistent atrial fibrillation. *JACC Clin Electrophysiol* 2022;**8**:882–91.
13. Lang RM, Badano LP, Mor-Avi V, Afilalo J, Armstrong A, Ernande L *et al.* Recommendations for cardiac chamber quantification by echocardiography in adults: an update from the American Society of Echocardiography and the European Association of Cardiovascular Imaging. *Euro Heart J—Cardiovasc Imaging* 2015;**16**: 233–71.
14. Nagel C, Schuler S, Dössel O, Loewe A. A bi-atrial statistical shape model for large-scale in silico studies of human atria: model development and application to ECG simulations. *Med Image Anal* 2021;**74**:102210.
15. Sánchez C, Bueno-Orovio A, Wettwer E, Loose S, Simon J, Ravens U *et al.* Inter-subject variability in human atrial action potential in Sinus rhythm versus chronic atrial fibrillation. *PLoS One* 2014;**9**:e105897.
16. Courtemanche M, Ramirez RJ, Nattel S. Ionic mechanisms underlying human atrial action potential properties: insights from a mathematical model. *Am J Physiol* 1998;**275**: H301–21.
17. Bergonti M, Spera FR, Ferrero TG, Nsahlai M, Bonomi A, Tijskens M *et al.* Characterization of atrial substrate to predict the success of pulmonary vein isolation: the prospective, multicenter MASH-AF II (multipolar atrial substrate high density mapping in atrial fibrillation) study. *J Am Heart Assoc* 2023;**12**:e027795.
18. Hopman LHGA, Visch JE, Bhagirath P, van der Laan AM, Mulder MJ, Razeghi O *et al.* Right atrial function and fibrosis in relation to successful atrial fibrillation ablation. *Eur Heart J—Cardiovasc Imaging* 2022;**24**:336–45.
19. Pope MT, Kuklik P, Briosa e Gala A, Leo M, Mahmoudi M, Paisey J *et al.* Spatial and temporal variability of rotational, focal, and irregular activity: practical implications for mapping of atrial fibrillation. *J Cardiovasc Electrophysiol* 2021;**32**:2393–403.
20. Roney CH, Sim I, Yu J, Beach M, Mehta A, Solis-Lemus JA *et al.* Predicting atrial fibrillation recurrence by combining population data and virtual cohorts of patient-specific left atrial models. *Circ Arrhythm Electrophysiol* 2022;**15**:e010253.
21. Gharaviri A, Pezzuto S, Potse M, Conte G, Zeemering S, Sobota V *et al.* Synergistic anti-arrhythmic effect of inward rectifier current inhibition and pulmonary vein isolation in a 3D computer model for atrial fibrillation. *Europace* 2021;**23**:i161–8.
22. Hindricks G, Potpara T, Dagres N, Arbelo E, Bax JJ, Blomström-Lundqvist C *et al.* 2020 ESC Guidelines for the diagnosis and management of atrial fibrillation developed in collaboration with the European Association for Cardio-Thoracic Surgery (EACTS): the Task Force for the diagnosis and management of atrial fibrillation of the European Society of Cardiology (ESC) developed with the special contribution of the European Heart Rhythm Association (EHRA) of the ESC. *Eur Heart J* 2020;**42**:373–498.
23. Wettwer E, Christ T, Endig S, Rozmaritsa N, Matschke K, Lynch JJ *et al.* The new anti-arrhythmic drug vernakalant: ex vivo study of human atrial tissue from sinus rhythm and chronic atrial fibrillation. *Cardiovasc Res* 2013;**98**:145–54.
24. Sachetto Oliveira R, Martins Rocha B, Burgarelli D, Meira W Jr, Constantinides C, Weber dos Santos R. Performance evaluation of GPU parallelization, space-time adaptive algorithms, and their combination for simulating cardiac electrophysiology. *Int J Numer Method Biomed Eng* 2018;**34**:e2913.
25. Boyle PM, Ochs AR, Ali RL, Paliwal N, Trayanova NA. Characterizing the arrhythmogenic substrate in personalized models of atrial fibrillation: sensitivity to mesh resolution and pacing protocol in AF models. *Europace* 2021;**23**:i3–i11.
26. Lankveld T, de Vos CB, Limantoro I, Zeemering S, Dudink E, Crijns HJ *et al.* Systematic analysis of ECG predictors of sinus rhythm maintenance after electrical cardioversion for persistent atrial fibrillation. *Heart Rhythm* 2016;**13**:1020–7.
27. Darkner S, Chen X, Hansen J, Pehrson S, Johannessen A, Nielsen JB *et al.* Recurrence of arrhythmia following short-term oral AMIOdarone after CATheter ablation for atrial fibrillation: a double-blind, randomized, placebo-controlled study (AMIO-CAT trial). *Eur Heart J* 2014;**35**:3356–64.
28. Camm AJ, Capucci A, Hohnloser SH, Torp-Pedersen C, Gelder ICV, Mangal B *et al.* A randomized active-controlled study comparing the efficacy and safety of vernakalant to amiodarone in recent-onset atrial fibrillation. *J Am Coll Cardiol* 2011;**57**:313–21.
29. Ahn J, Shin DG, Han SJ, Lim HE. Does isolation of the left atrial posterior wall using cryoballoon ablation improve clinical outcomes in patients with persistent atrial fibrillation? A prospective randomized controlled trial. *Europace* 2022;**24**:1093–101.
30. Li X, Liu T, Cui B, Zhang J, Chen Y, Wu G. Efficacy and safety ablation index-guided high-energy linear ablation for persistent atrial fibrillation: PVI plus linear ablation of mitral isthmus and posterior box isolation. *J Clin Med* 2023;**12**:619.
31. Davong B, Adeliño R, Delasnerie H, Albenque JP, Combes N, Cardin C *et al.* Pulsed-field ablation on mitral isthmus in persistent atrial fibrillation. *JACC Clin Electrophysiol* 2023;**9**: 1070–81.
32. Kim JS, Shin SY, Na JO, Choi CU, Kim SH, Kim JW *et al.* Does isolation of the left atrial posterior wall improve clinical outcomes after radiofrequency catheter ablation for persistent atrial fibrillation?: a prospective randomized clinical trial. *Int J Cardiol* 2015;**181**: 277–83.
33. Remme CA, Heijman J, Gomez AM, Zaza A, Odening KE. 25 Years of basic and translational science in EP Europace: novel insights into arrhythmia mechanisms and therapeutic strategies. *Europace* 2023;**25**:euad210.
34. Hwang M, Kwon SS, Wi J, Park M, Lee HS, Park JS *et al.* Virtual ablation for atrial fibrillation in personalized in-silico three-dimensional left atrial modeling: comparison with clinical catheter ablation. *Prog Biophys Mol Biol* 2014;**116**:40–7.
35. Shim J, Hwang M, Song JS, Lim B, Kim TH, Joung B *et al.* Virtual in-silico modeling guided catheter ablation predicts effective linear ablation lesion set for longstanding persistent atrial fibrillation: multicenter prospective randomized study. *Front Physiol* 2017;**8**:8.
36. Azzolin L, Eichenlaub M, Nagel C, Nairn D, Sanchez J, Unger L *et al.* Personalized ablation vs. conventional ablation strategies to terminate atrial fibrillation and prevent recurrence. *Europace* 2022;**25**:211–22.
37. Roney CH, Beach ML, Mehta AM, Sim I, Corrado C, Bendikas R *et al.* In silico comparison of left atrial ablation techniques that target the anatomical, structural, and electrical substrates of atrial fibrillation. *Front Physiol* 2020;**11**:11.
38. Roney C, Alonso JSL, Sim I, Ulgen O, Rodero C, Strocchi M *et al.* In silico biatrial fibrosis ablation improves atrial fibrillation outcome. *Europace* 2023;**25**:euad122.587.
39. Boyle PM, Zghaib T, Zahid S, Ali RL, Deng D, Franceschi WH *et al.* Computationally guided personalized targeted ablation of persistent atrial fibrillation. *Nat Biomed Eng* 2019;**3**:870–9.
40. Kany S, Reissmann B, Metzner A, Kirchhof P, Darbar D, Schnabel RB. Genetics of atrial fibrillation—practical applications for clinical management: if not now, when and how? *Cardiovasc Res* 2021;**117**:1718–31.
41. Saad EB, Slater C. Complete isolation of the left atrial posterior wall (box lesion) to treat longstanding persistent atrial fibrillation. *J Atr Fibrillation* 2014;**7**:1174.
42. Valderrábano M, Peterson LE, Swarup V, Schurmann PA, Makkar A, Doshi RN *et al.* Effect of catheter ablation with vein of marshall ethanol infusion vs catheter ablation alone on persistent atrial fibrillation: the VENUS randomized clinical trial. *JAMA* 2020; **324**:1620–8.