Left atrium remodeling predicts late recurrence of paroxysmal atrial fibrillation after second generation cryoballoon ablation

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Abstract

Background: Atrial fibrillation (AF) is the most common arrhythmia worldwide. Nowadays, AF ablation is a valuable treatment option. It has been shown that the left atrium (LA) diameter is a predictor of AF recurrence after cryoballoon ablation (CBA). Since it does not reflect the true LA size, we compared the role of different LA anatomical parameters using echocardiography for the prediction of AF recurrence after CBA.

Methods: We retrospectively included 209 patients (mean age 56.1 ± 13.6 years, male 62%) with paroxysmal AF undergoing CBA. A transthoracic echocardiography was performed in all patients.

Results: At a mean follow-up of 16.9 ± 6.3 months, AF recurred in 25.4% of the patients. LA anterior - posterior diameter (LAD), LA minimum volume (LAmin) and early AF recurrence were independent predictors of recurrence. Based on receiver operating characteristics, cut-off values for LAD and, LAmin were 41 mm, 23.69 mL, respectively. The negative predictive values for recurrence were 73% and 87.3% respectively. In patients with AF recurrence, a significant proportion (30.2%) showed LA longitudinal remodeling (LA superior – inferior diameter) even though classically measured LAD was normal.

Conclusions: Longitudinal LA remodeling plays an additional role for predicting AF recurrence after CBA, in patients without LAD dilation. Moreover, LAmin had a high negative predictive value and was an independent predictor of AF recurrence. Therefore, a more complete LA anatomical assessment allows a better prediction of AF recurrences after CBA.

Keywords: Atrial fibrillation, Cryoballoon ablation, Left atrium, Echocardiography

Background

Atrial fibrillation (AF) is the most common cardiac arrhythmia, with an increasing frequency worldwide, and it is associated with an elevated risk for stroke, heart failure and mortality. Its prevalence in developed countries is currently estimated to 1.5–2% of the general population [1]. Resinusalisation is one of the main goals in AF patients. Several ablation strategies have proven to be efficient in the treatment of AF [2, 3]. Among them, one of the most promising and effective approaches is the second generation cryoballoon ablation (CBA) (Arctic Front Advance, Medtronic). However, up to 20% of patients experience AF recurrence after CBA [4, 5].

Recent data have shown that an increase of the left atrium (LA) anterior - posterior diameter (LAD) assessed by echocardiography is a powerful predictor of recurrence after CBA [4, 6]. A possible explanation for the recurrence of AF could be the presence of myocardial fibrosis and remodeling of the LA [7]. However, LAD does not reflect the true size of the LA [8, 9]. We hypothesized that a more complete analysis of the LA dimensions could be a
better predictor of AF recurrence after CBA. Therefore, we compared the role of different anatomical parameters of the LA measured by echocardiography for predicting the recurrence of AF after CBA.

Methods
Patient characteristics
We retrospectively included 209 patients having undergone CBA for paroxysmal AF in our center from January 2014 to February 2016. Baseline demographic characteristics can be found in Table 1. CHA2DS2-VASc score was calculated according to European Society of Cardiology Guidelines [1].

The study was approved by the local Ethical Committee and was carried out in accordance with the ethical principles for medical research involving human subjects established by Helsinki’s Declaration, protecting the privacy of all participants, as well as the confidentiality of their personal information. All patients provided written informed consents.

Transthoracic echocardiography
A comprehensive transthoracic echocardiography (TTE) (using GE Vingmed Ultrasound, Vivid E9, Horten, Norway; Phillips Epiq, Philips, Andover, Massachusetts) was performed in all patients, according to the recommendations [10, 11]. Standard parasternal long and short axis views and apical two-, three- and four-chamber views were available in all patients. Left ventricular (LV) dimensions were measured using M–Mode in the parasternal long axis view. LAD was measured using anatomical M-mode and two–dimensional assessment (2D) in the parasternal long axis view. LA maximum (LAmx) and minimum (LAmi) volumes were measured in two- and four-chamber views, using the biplane area-length method (Fig. 1). LA volumes were indexed based on the patient’s body surface area. Left atrium superior – inferior diameter was measured in apical two - and four chamber views (Fig. 2). Doppler mitral inflow peak early diastolic (E), peak atrial systolic velocities (A) and deceleration time were measured. Tissue Doppler imaging at the level of the mitral annulus was used to measure peak early diastolic septal and lateral e’, as well as peak late diastolic a’, corresponding to the P-wave on the electrocardiogram (ECG) [10, 12].

To determine the reproducibility of the LAD, LA superior – inferior diameter, LAmx and LAmi, the measurements were repeated in 20 randomly selected patients by an additional investigator and by the same primary investigator 1 week later. During the repeated analysis, the investigators were blinded to the results of all previous measurements.

Pre-procedural management
The antiarrhythmic drugs (AADs) were discontinued at least 3 days before the procedure, except for amiodarone, which was stopped one month before.

Cryoballon ablation procedure
Our standard ablation procedure has been previously reported in detail [13]. Briefly, after obtaining LA access, through a steerable 15 Fr sheath (FlexCath Advance®, Medtronic®), an inner lumen mapping catheter (ILMC) (Achieve®, Medtronic®) was advanced in each pulmonary vein (PV) ostium and baseline electrical information was gathered. Optimal vessel occlusion was considered as achieved upon selective contrast injection showing total contrast retention with no backflow into the atrium. Once occlusion was documented, cryothermal energy was started for at least 180 s. Usually, the PVs were treated as follows: first, the left superior PV (LSPV), then the left inferior (LIPV), right inferior (RIPV) and right superior (RSPV). PV activity was recorded with the ILMC at a proximal site within the ostium prior to ablation in each vein. If PV potentials (PVPs) were visible during energy delivery, time to isolation was recorded as the time from the start of the cryoenergy application until the PVPs completely disappeared or were dissociated from LA activity. In cases of phrenic nerve palsy (PNP), recovery of diaphragmatic contraction was carefully monitored for 30 min. Further additional cryoenergy applications were not applied if the veins were isolated after the initial freeze. If needed, pacing from the distal and/or proximal coronary sinus was performed to recognise far field atrial signals from PVPs recorded on the mapping catheter. During the whole procedure, activated clotting time was maintained > 250 s. In order to avoid phrenic nerve palsy, diaphragmatic stimulation was achieved by pacing the phrenic nerve during septal pulmonary veins ablations.

Postablation management
Patients were discharged the day after ablation if the clinical status was stable. Following the procedure, all the patients were continuously monitored with ECG telemetry for at least 18 h. Oral anticoagulation was not discontinued for the ablation and continued for at least 2 months following the CHA2DS2-VASc score. During the blanking period (BP), AADs were continued. The decision to continue AADs after the BP or to perform a repeat procedure was taken if a first episode of recurrence of AF occurred.

Follow-up
All the included patients underwent physical examination and a 24 h Holter recording at 1, 3, 6 and 12 months after the ablation. Additional Holter
| Characteristic                      | Study population (n = 209) | AF recurrence – (n = 156) | AF recurrence + (n = 53) | P value |
|------------------------------------|---------------------------|--------------------------|--------------------------|---------|
| Age (y)                            | 56.1 ± 13.5               | 55.6 ± 13.5              | 57.4 ± 13.7              | 0.335   |
| Male gender (n, %)                 | 130 (62.2)                | 98 (62.8)                | 32 (60.4)                | 0.753   |
| Body mass index (kg/m²)            | 26.5 ± 4.7                | 26.3 ± 4.5               | 27 ± 5.3                 | 0.549   |
| Tobacco use (n, %)                 | 52 (24.8)                 | 35 (28.8)                | 17 (32)                  | 0.281   |
| Hypertension (n, %)                | 86 (41.1)                 | 59 (37.8)                | 27 (50.9)                | 0.094   |
| Diabetes (n, %)                    | 16 (7.7)                  | 9 (5.8)                  | 7 (13.2)                 | 0.079   |
| Dyslipidemia (n, %)                | 75 (35.9)                 | 55 (35.3)                | 20 (37.7)                | 0.747   |
| History of heart failure (n, %)    | 6 (2.9)                   | 4 (2.6)                  | 2 (3.8)                  | 0.651   |
| Coronary artery disease (n, %)     | 15 (7.2)                  | 10 (6.4)                 | 5 (9.4)                  | 0.464   |
| History of TIA/A/CVA (n, %)        | 19 (9.1)                  | 15 (9.6)                 | 4 (7.5)                  | 0.653   |
| Failed drugs before ablation       |                           |                          |                          |         |
| Flecainide (n, %)                  | 46 (22)                   | 31 (19.8)                | 15 (28.3)                | 0.177   |
| Propafenone (n, %)                 | 1(0.47)                   | 0 (0)                    | 1(1.8)                   | 0.083   |
| Amiodarone (n, %)                  | 9 (4.3)                   | 6 (3.8)                  | 3 (5.6)                  | 0.555   |
| Sotalol (n, %)                     | 24 (11.4)                 | 16 (10.2)                | 8 (15)                   | 0.316   |
| Metoprolol (n, %)                  | 2 (0.9)                   | 2 (1.2)                  | 0 (0)                    | 0.412   |
| Bisoprolol (n, %)                  | 78 (37.3)                 | 57 (36.5)                | 21 (39.6)                | 0.620   |
| Nebivolol (n, %)                   | 6 (3.8)                   | 5 (3.2)                  | 1 (1.8)                  | 0.632   |
| ACE inhibitors before ablation     | 26 (12.4)                 | 20 (12.8)                | 6 (11.3)                 | 0.809   |
| ARBs before ablation               | 11 (5.2)                  | 9 (5.7)                  | 2 (3.7)                  | 0.592   |
| Prior ablation                     | 39 (18.6)                 | 29 (18.5)                | 10 (18.8)                | 0.547   |
| - AVNRT                          | 13 (6.2)                  | 12 (7.7)                 | 1 (1.9)                  | 1.000   |
| - Right atrial flutter             | 13 (6.2)                  | 8 (5.1)                  | 5 (9.4)                  | 1.000   |
| - AF                            | 11 (5.3)                  | 8 (5.1)                  | 3 (5.7)                  | 1.000   |
| - Other                          | 2 (1.0)                   | 1 (0.6)                  | 1 (1.9)                  | 1.000   |
| CHA2DS2_VASc score                | 1.3 ± 1.4                 | 1.2 ± 1.3                | 1.6 ± 1.7                | 0.134   |
| Oral anticoagulation (n, %)        | 60 (28.7)                 | 41 (26.3%)               | 19 (35.8%)               | 0.552   |
| Aspirin (n, %)                     | 23 (11)                   | 17 (10.9)                | 6 (11.3)                 | 0.874   |
| Follow-up duration (months)        | 16.8 ± 6.3                | 16.9 ± 5.7               | 16.6 ± 7.7               | 0.795   |
| Re-do ablation                    | 31 (14.8)                 | 0 (0)                    | 31 (58.4)                |         |
| Medication in BP                   |                           |                          |                          |         |
| - Flecainide (n, %)                | 62 (29.6)                 | 42 (26.9)                | 20 (37.7)                | 0.122   |
| - Propafenone (n, %)               | 2 (0.95)                  | 1 (0.64)                 | 1 (1.8)                  | 0.415   |
| - Amiodarone (n, %)                | 9 (4.3)                   | 6 (3.8)                  | 3 (5.6)                  | 0.561   |
| - Sotalol (n, %)                   | 49 (23.4)                 | 34 (21.8)                | 15 (28.3)                | 0.310   |
| - Metoprolol (n, %)                | 1(0.4)                    | 1(0.6)                   | 0 (0)                    | 0.562   |
| - Bisoprolol (n, %)                | 111 (53.11)               | 81 (51.9)                | 30 (56.6)                | 0.497   |
| - Nebivolol (n, %)                 | 4 (1.9)                   | 4 (2.5)                  | 0 (0)                    | 0.241   |
| - ACE inhibitors (n, %)            | 27 (12.9)                 | 21 (13.4)                | 6 (11.3)                 | 0.710   |
| - ARBs (n, %)                      | 6 (2.8)                   | 6 (6)                    | 0 (0)                    | 0.150   |

Medication after BP

- Flecainide (n, %) 25 (11.9) 17 (10.8) 8 (15) 0.398
- Propafenone (n, %) 2 (0.9) 0 (0) 2 (3.7) 0.060
- Amiodarone (n, %) 6 (2.8) 3 (1.9) 3 (5.6) 0.154
monitoring was performed if arrhythmic symptoms occurred. All documented AF episodes > 30 s were considered as a recurrence. A BP of 3 months was considered for the study.

**Statistical analysis**

Continuous variables are expressed as mean ± standard deviation. Categorical variables are expressed as percentages. Comparisons of continuous variables were done with a Student t-test or Mann-Whitney U-test and binomial variables with a chi-square or Fisher test as appropriate. Receiver-operator characteristic (ROC) curves were constructed to evaluate the performance of variables in predicting AF recurrence and to calculate adjusted cut-off values, specificity and sensitivity of the parameters (using Youden’s Index) in the prediction of AF recurrence. Freedom from AF recurrence was estimated by Kaplan-Meier method and time-dependent comparisons by the log-rank test. In order to evaluate potential predictors of AF recurrence, a separate Cox proportional hazard model was used. Intraclass correlation coefficient (ICC) was used to determine the intra- and inter-observer variability. The calculated ICCs were judged as follows: 0.50 to 0.74 poor to moderate, 0.75 to 0.92 good and > 0.93 excellent. Statistical significance was considered with a P-value < 0.05. Statistical analyses were conducted using IBM SPSS Statistic for Windows, Version 24.0 (Armonk, NY: IBM Corp.)

| Characteristic | Study population (n = 209) | AF recurrence – (n = 156) | AF recurrence + (n = 53) | P value |
|----------------|---------------------------|--------------------------|-------------------------|---------|
| - Sotalol (n, %) | 26 (12.4) | 18 (11.5) | 8 (15) | 0.478 |
| - Metoprolol (n, %) | 1 (0.4) | 1 (0.6) | 0 (0) | 0.562 |
| - Bisoprolol (n, %) | 62 (29.6) | 50 (23.9) | 12 (22.6) | 0.211 |
| - Nebivolol (n, %) | 6 (2.8) | 5 (3.2) | 1 (1.8) | 0.628 |
| - ACE inhibitors (n, %) | 27 (12.9) | 21 (13.4) | 6 (11.3) | 0.710 |
| - ARBs (n, %) | 6 (2.8) | 6 | 0 | 0.150 |
| Recurrence in BP | 8 (3.8) | 0 (0) | 8 (15.1) | < 0.001 |

*TIA Transient ischaemic attack, *CVA Cerebrovascular accident, *ACE inhibitors, angiotensin – converter enzyme inhibitors, *ARBS angiotensin receptor blockers, *AVNRT atrioventricular nodal reentry tachycardia, *AF atrial fibrillation, *BP blanking period

**Fig. 1 a.** Illustration of the left atrium anterior – posterior diameter measurement in M-Mode, using the parasternal long axis view. **b.** Left atrium anterior – posterior diameter measurement in 2D parasternal long axis view. **c.** Left atrium volume measurement in four – chamber view. **d.** Left atrium volume measurement in two – chamber view.
Results
At a mean follow-up of 16.8 ± 6.3 months after the BP, 53 (25.4%) patients presented recurrences of AF.

Baseline population characteristics and recurrences
Baseline clinical and demographic characteristics of the study population are listed in Table 1. There was a significant difference between the groups with and without recurrence for patients who presented early AF recurrence.

Echocardiographic characteristics are presented in Table 2. In brief, there was a significant difference in patients with and without recurrence respectively for LV end-diastolic diameter (50.3 ± 7.9 mm vs. 47.2 ± 6.5 mm, \( p = 0.013 \)), LV mass index (83.7 ± 28.4 g/m\(^2\) vs. 76.6 ± 26.3 g/m\(^2\), \( p = 0.045 \)), LAD (42.2 ± 7.2 mm vs. 39 ± 6.1 mm, \( p = 0.013 \)), LMax (67.3 ± 27.8 mL vs. 57.7 ± 21.2 mL, \( p = 0.036 \)), Lamin (36.7 ± 20.1 mL vs. 27.5 ± 14.9 mL, \( p = 0.004 \)) and Lamin indexed (18.1 ± 9.3 mL/m\(^2\) vs. 14.1 ± 7.4 mL/m\(^2\), \( p = 0.007 \)).

Procedural data
All the patients underwent the procedure with the large 28-mm CBA. At the beginning of the procedure, 203 (97.1%) patients were in sinus rhythm. The mean total procedure and fluoroscopy time were 66.9 ± 20.2 min, respectively 13.8 ± 7.5 min. The mean number of freeze-thaw cycles was 1.2 ± 0.4 in the LSPV, 1.1 ± 0.3 in the LIPV, 1.19 ± 0.44 in the RSPV, 1.2 ± 0.5 in the RIPV. The mean minimal temperatures obtained were −48.7 ± 8.6 °C in LSPV, −48.7 ± 8.6 in LIPV, −49.2 ± 10.8 in RSPV, −46 ± 14.6 in RIPV. There were no significant differences regarding the procedural data between the two groups. Procedural data can be found in Table 3.

Global versus longitudinal LA remodeling and AF recurrences
Fig. 3 illustrates that using only the LAD, with an adjusted cut-off value of 41 mm, led missing up to 30.2% of AF recurrences. In the patients with recurrence without LAD dilation, a significant number of patients had longitudinal LA remodeling (increased superior – inferior LA diameter, with an adjusted cut-off value of 41 mm).

LA volumes and AF recurrences
Multivariate regression analysis showed that LAD, Lamin and early AF recurrence were independent predictors of AF recurrence after the BP, as illustrated in Table 4.

ROC analysis revealed that for a cut-off value of 23.69 ml for Lamin it was associated with a high sensitivity for predicting AF recurrence after the BP (79.5% sensitivity, 46% specificity). For the above mentioned adjusted cut-off values, Lamin had a high negative predictive values for AF recurrences 87.3%.

In contrast, for LAD, sensitivity and specificity were 56.3% and 66%, respectively, with a lower negative predictive value of 73% compared to Lamin.

Reproducibility
Intraobserver variability was excellent for all the parameters. For LAD the ICC was 0.972, for LA superior – inferior diameter ICC = 0.95, for LMax ICC = 0.964 and for Lamin ICC = 0.940. Interobserver reliability was good for LAD (ICC = 0.906), LMax (ICC = 0.907) and Lamin (0.860) and excellent for LA superior – inferior diameter (ICC = 0.96).

Discussion
The main findings of our study are: 1) recurrence of AF after the BP occurred in 25.4% of all subjects with paroxysmal AF. 2) In the group of patients with AF recurrence, longitudinal remodeling (increased superior – inferior LA diameter) was also present in patients with normal LAD. 3) In the global population of patients undergoing CBA, LA minimum volume was an independent predictor of AF recurrences after the BP. These
| Characteristic | Study population (n = 209) | AF recurrence – (n = 156) | AF recurrence + (n = 53) | P value |
|---------------|---------------------------|--------------------------|-------------------------|---------|
| Heart rate during echocardiography | 75.6 ± 14.5 | 76.3 ± 14.1 | 75.6 ± 14.5 | 0.080 |
| Left ventricle | | | | |
| - LVEF\(^a\), % | 54.4 ± 5.2 | 54.6 ± 4.7 | 53.8 ± 6.4 | 0.343 |
| - LV\(^b\) EDD\(^c\), mm | 48 ± 7 | 47.2 ± 6.5 | 50.3 ± 7.9 | 0.013 |
| - LV ESD\(^d\), mm | 30.4 ± 5.8 | 30.4 ± 5.9 | 30.5 ± 5.7 | 0.939 |
| - IVSd\(^e\), mm | 9.5 ± 1.7 | 9.5 ± 1.78 | 9.6 ± 1.7 | 0.763 |
| - PWd\(^f\), mm | 9.3 ± 1.6 | 9.1 ± 1.4 | 9.6 ± 1.8 | 0.131 |
| - LV mass, g | 140.4 ± 72.8 | 136.8 ± 70.1 | 159.9 ± 80 | 0.073 |
| - LV mass index, g/m\(^2\) | 78.4 ± 26.9 | 76.6 ± 26.3 | 83.7 ± 28.4 | 0.045 |
| - LV EDV\(^g\), mL | 99 ± 32.5 | 96.4 ± 30.5 | 109.8 ± 39.6 | 0.223 |
| - LV ESV\(^h\), mL | 46.8 ± 20.8 | 45.4 ± 20.6 | 52.4 ± 21.6 | 0.196 |
| Left atrium | | | | |
| - LA\(^i\) antero-posterior diameter, mm | 39.8 ± 6.5 | 39 ± 6.1 | 42.2 ± 7.2 | 0.013 |
| - LA antero-posterior diameter indexed, mm/m\(^2\) | 20.4 ± 3.4 | 20.1 ± 3.3 | 20.4 ± 3.4 | 0.182 |
| - LA superior- inferior diameter, mm | 48.8 ± 7.6 | 48.3 ± 7.6 | 50.3 ± 7.4 | 0.095 |
| - LA maximum volume, mL | 60 ± 23.3 | 57.7 ± 21.2 | 67.3 ± 27.8 | 0.036 |
| - LA maximum volume indexed, mL/m\(^2\) | 30.4 ± 10.8 | 29.5 ± 10 | 33.5 ± 12.6 | 0.067 |
| - LA minimum volume, mL | 29.8 ± 16.8 | 27.5 ± 14.9 | 36.7 ± 20.1 | 0.004 |
| - LA minimum volume indexed, mL/m\(^2\) | 15.1 ± 8 | 14.1 ± 7.4 | 18.1 ± 9.3 | 0.007 |
| Doppler | | | | |
| - E- wave velocity, m/s | 0.6 ± 0.1 | 0.6 ± 0.1 | 0.6 ± 0.1 | 0.768 |
| - A – wave velocity, m/s | 0.5 ± 0.1 | 0.5 ± 0.1 | 0.4 ± 0.1 | 0.405 |
| - E/A (ratio) | 1.4 ± 0.5 | 1.3 ± 0.5 | 1.4 ± 0.5 | 0.314 |
| - DTE\(^j\), ms | 180.8 ± 44.4 | 180.2 ± 45.4 | 182.7 ± 41.6 | 0.482 |
| - TDI \(e\) septal, cm/s | 7.4 ± 2.1 | 7.3 ± 2 | 7.8 ± 2.5 | 0.366 |
| - TDI \(e\) lateral, cm/s | 9.4 ± 4.9 | 9.6 ± 5.3 | 9 ± 2.8 | 0.965 |
| - E/e’ avg. (ratio) | 8.6 ± 2.9 | 8.7 ± 2.9 | 8.4 ± 3.2 | 0.893 |
| - TDI \(a\) septal, cm/s | 7.4 ± 2 | 7.6 ± 2 | 6.8 ± 2 | 0.053 |
| - TDI \(a\) lateral, cm/s | 8 ± 2.4 | 8.1 ± 2.4 | 7.5 ± 2.4 | 0.111 |
| Mitral regurgitation (n, %) | 97 (46.4) | 70 (44.8) | 27 (50.9) | 0.594 |
| - Mild (n, %) | 90 (43.1) | 64 (41) | 26 (49.1) |
| - Moderate (n, %) | 7 (3.3) | 6 (3.8) | 1 (1.9) | 0.677 |
| Aortic regurgitation (mild) (n, %) | 24 (11.9) | 20 (12.8) | 4 (7.5) | 0.298 |
| Tricuspid regurgitation (n, %) | 106 (50.7) | 77 (49.3) | 29 (54.7) | 0.338 |
| - Mild (n, %) | 96 (45.9) | 70 (44.8) | 26 (49) | 0.338 |
| - Moderate (n, %) | 9 (4.3) | 7 (4.4) | 2 (3.7) |
| - Severe (n, %) | 1 (0.4) | 0 (0) | 1 (1.8) |
| TAPSE\(^l\), mm | 23.1 ± 4 | 23 ± 3.7 | 23.5 ± 4.9 | 0.253 |
| IVC\(^m\), mm | 14.3 ± 4 | 14.1 ± 3.9 | 15.1 ± 4.2 | 0.258 |
| TR\(^n\) gradient, mmHg | 249 ± 7.5 | 251 ± 7.7 | 241 ± 6.9 | 0.783 |

\(^a\) LVEF left ventricle ejection fraction, \(^b\) LV left ventricle, \(^c\) EDD end – diastolic diameter, \(^d\) ESD end – systolic diameter, \(^e\) IVSd Interventricular septum end-diastolic diameter, \(^f\) PWd posterior wall end-diastolic diameter, \(^g\) EDV end – diastolic volume, \(^h\) ESV end – systolic volume, \(^i\) LA left atrium, \(^j\) DTE deceleration time of E – wave, \(^k\) TDI tissue Doppler imaging, \(^l\) TAPSE tricuspid annulus plane systolic excursion, \(^m\) IVC inferior vena cava, \(^n\) TR tricuspid regurgitation
| Characteristic                        | Study population (n = 209) | AF recurrence – (n = 156) | AF recurrence + (n = 53) | P value |
|--------------------------------------|----------------------------|--------------------------|--------------------------|---------|
| Procedural time, min                 | 66.9 ± 20.2                | 67 ± 20.3                | 66.5 ± 19.9              | 0.918   |
| Fluoroscopy time, min                | 13.8 ± 7.5                 | 13.4 ± 7.2               | 14.7 ± 8.3               | 0.452   |
| Freezes in LSPV                        | 1.2 ± 0.4                  | 1.2 ± 0.4                | 1.2 ± 0.4                | 0.656   |
| Freezes in LIPV                       | 1.1 ± 0.3                  | 1.1 ± 0.4                | 1.1 ± 0.3                | 0.215   |
| Freezes in RSPV                        | 1.1 ± 0.4                  | 1.1 ± 0.4                | 1.2 ± 0.4                | 0.086   |
| Freezes in RIPV                        | 1.2 ± 0.5                  | 1.2 ± 0.4                | 1.2 ± 0.5                | 0.366   |
| LSPV freeze duration, s              | 223.6 ± 78.9               | 224.8 ± 80.5             | 220 ± 74.6               | 0.782   |
| LIPV freeze duration, s              | 208.2 ± 65.5               | 211.1 ± 69               | 199.7 ± 53.7             | 0.412   |
| RSPV freeze duration, s              | 208.5 ± 79                 | 206 ± 75                 | 215.9 ± 89.8             | 0.647   |
| RIPV freeze duration, s              | 225.4 ± 89.1               | 223 ± 89.4               | 232.5 ± 88.8             | 0.147   |
| Min temp in LSPV, °C                | −48.7 ± 8.6                | −48.7 ± 9.5              | −48.6 ± 5                | 0.325   |
| Min temp in LIPV, °C                | −45.4 ± 8.6                | −45.6 ± 9.5              | −44.9 ± 5.4              | 0.192   |
| Min temp in RSPV, °C                | −49.2 ± 10.8               | −49.3 ± 11.9             | −47 ± 14.6               | 0.246   |
| Min temp in RIPV, °C                | −46 ± 14.6                 | −45.4 ± 16.3             | −47.9 ± 6.5              | 0.698   |
| Pulmonary veins variants              | 102 (48.8)                 | 79 (50.6)                | 23 (43.3)                | 0.347   |
| - Left common ostium                  | 58 (27.8)                  | 42 (26.9)                | 16 (30.2)                | 0.647   |
| - Right middle pulmonary vein        | 29 (13.9)                  | 25 (16.0)                | 4 (7)                    | 0.124   |
| - Right common ostium                | 8 (3.8)                    | 7 (4.5)                  | 1 (1.9)                  | 0.640   |
| - Other                               | 7 (3.3)                    | 5 (3.2)                  | 2 (3.8)                  | 0.686   |

*LSPV left superior pulmonary vein, "LIPV left inferior pulmonary vein, "RSPV right superior pulmonary vein, "RIPV right inferior pulmonary vein, "Min temp, minimum temperature

Fig. 3 Proportions of patients with AF recurrences according to the adjusted cut-off values for LA superior – inferior diameter (41 mm) and LA anterior – posterior diameter (41 mm), showing that 30.2% of patients with recurrence had longitudinal remodeling, in the absence of an increased anterior-posterior diameter
results suggest that AF recurrence after ablation also occurs in the absence of global LA remodeling assessed by LAD.

This study evaluates the role of additional anatomical parameters of the LA using two-dimensional echocardiography for the prediction of recurrence of AF after the BP in patients who have undergone CBA.

The incidence of recurrence of AF after the BP in our study was of 25.4%, which is consistent with results of previous studies. Gerede et al. [6] have reported a rate of recurrence after the BP of 31.3%. In a recent study published by Coutino et al. [5] they showed a rate of recurrence after the BP of 25.2%. Previous studies have shown that the presence of recurrence during the BP is an independent predictor of late AF recurrence, which was also confirmed by our analysis [14, 15].

Previous studies have shown that the enlargement of LAD is an independent predictor of recurrence of AF after the BP, suggesting that the remodeling of the LA is a leading cause for late recurrences of AF [4, 6]. This is consistent with our results that report LAD as a predictor of recurrence of AF after CBA.

However, the remodeling of the LA can occur non-uniformly, correlating with reduced success of different ablation techniques [9, 16] and furthermore, LAD non-uniformly, correlating with reduced success of different ablation techniques [9, 16] and furthermore, LAD non-uniformly, correlating with reduced success of different ablation techniques. The study evaluates the role of additional anatomical parameters of the LA using two-dimensional echocardiography for the prediction of recurrence of AF after the BP in patients who have undergone CBA.

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Study limitations

This is a single center retrospective study, therefore results may not be generalized. Larger prospective clinical studies are warranted to confirm our findings. Moreover, the present study addressed the prognostic value of LA anatomy only in paroxysmal AF and cannot be extrapolated to a persistent AF population undergoing CBA.

Interestingly, our study showed that L.Amin was an independent predictor of AF recurrence after the BP, and that it was associated with a high sensitivity for predicting recurrence for the adjusted cut-off values (79.5%).

Limited data is available regarding the value of LA minimum volume and its prognostic value for different cardiovascular events. L.Amin volume was associated with the occurrence of AF in an elderly cohort, as showed by Fatema et al. [17] and it was the most accurate parameter reflecting anatomical remodeling associated with paroxysmal AF in a recent study performed by Schaaf et al. [7]. In several studies, LA minimum volume was associated with diastolic dysfunction starting from incipient stages [18, 19]. However, we did not find a significant difference regarding the diastolic function between the groups with and without recurrence. These findings suggest that LA minimum volume represents a more subtle form of LA remodeling that could also trigger the occurrence of AF recurrence after ablation.

Whether LA minimum volume may have a better prognostic value in predicting late AF recurrences following cryoballoon ablation requires further investigations.

As suggested by Canpolat et al. [20], CBA may have a positive impact on LA reverse remodeling. Since the remodeling is a continuous process, measurements performed at a single moment may not reflect the truth and a temporal trend of the LA volumes should be assessed. However, data regarding LA reverse remodeling is limited and there are no clear definitions for it, therefore it should be evaluated in further studies.
Future perspectives
As mentioned earlier, it is widely accepted that anatomical remodeling is associated with recurrence of AF after ablation [4, 6, 9, 16, 21, 22]. 2D echocardiography can provide useful information for the detection of patients at risk to develop recurrences of AF, but LA volumes are still based on only two orthogonal planes and related to geometrical assumptions. Our results suggest that LA longitudinal remodeling could play an additional role in the prediction of AF recurrences, as the expansion of the LA can be asymmetrical. Therefore, advanced imaging techniques such as three – dimensional (3D) echocardiography can overcome these limitations and provide more accurate and reproducible LA measurements [23]. Moreover, several studies showed that LA strain could predict recurrence of AF after catheter ablation. Therefore, strain imaging may bring additional information regarding the more subtle structural and functional left atrium remodeling in such patients and could be used for the follow-up assessment of possible reverse atrial remodeling [24–26].

Conclusions
The results of our study showed that LA longitudinal remodeling, assessed by the increase of LA superior – inferior diameter plays an additional role in AF recurrence after CBA. Moreover, LA minimum volume was an independent predictors of AF recurrences after the BP, suggesting that less advanced forms of LA remodeling, not detected by the dilation of LAD can trigger AF recurrences. Further large – scale prospective randomized studies are needed to confirm our results.

Abbreviations
AADs: Antiarrhythmic drugs; AF: Atrial fibrillation; BP: Blanking period; CBA: Cryoballoon ablation; ECG: Electrocardiogram; ICC: Intraclass correlation coefficient; ILMC: Inner lumen mapping catheter; LA: Left atrium; LAD: Left atrium anterior – posterior diameter; LAMax: Left atrium maximum volume; LAMin: Left atrium minimum volume; LIPV: Left inferior pulmonary vein; LSPV: Left superior pulmonary vein; LV: Left ventricle; PNP: Phrenic nerve palsy; PV: Pulmonary vein; PVPs: Pulmonary vein potentials; RSPV: Right inferior pulmonary vein; ROC: Receiver – operator characteristic; RIPV: Right superior pulmonary vein; TTE: Transthoracic echocardiography.

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Availability of data and materials
The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions
All authors contributed equally in analyzing and interpreting the patients’ data and writing the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate
The study was approved by the local Ethical Committee and was carried out in accordance with the ethical principles for medical research involving human subjects established by Helsinki’s Declaration, protecting the privacy of all participants, as well as the confidentiality of their personal information. All patients provided written informed consents.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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