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Procedural success, safety and patients satisfaction after second ablation of atrial fibrillation in the elderly: results from the German Ablation Registry

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Abstract

Background Aged patients are underrepresented in clinical trials on catheter ablation of atrial fibrillation (AF). In addition, results of outcomes after repeat ablation in the elderly are lacking. We report the results of first repeat AF ablation procedures of aged patients from a real-world multicenter prospective registry.

Methods Patients undergoing second AF ablation included in the prospective, multicenter German Ablation Registry were divided in two groups (age > 70 years (group 1) and age \leq 70 years (group 2)) and analyzed for procedural characteristics and clinical follow-up.

Results 738 patients were analyzed (108 patients in group 1, 630 patients in group 2). Significantly more aged patients had structural heart disease (56 patients (51.9%) vs. 203 patients (32.2%), p < 0.001). The majority of the patients underwent repeat pulmonary vein isolation (101 patients (93.5%) vs. 593 patients (94.1%), p = 0.98). More aged patients underwent ablation of left atrial linear lesions (78.1% vs. 57.3% of all linear lesions, p = 0.027). There was no difference in the occurrence of peri-procedural complications (7 patients (6.5%) vs. 24 patients (3.8%), p = 0.30). Recurrence of atrial arrhythmias was documented in 45/105 (42.9%) and 252/603 (41.8%) patients with available follow-up in groups 1 and 2 after a median of 447 (400; 532) and 473 (411; 544) days (p = 0.84). A comparable amount of patients were asymptomatic or reported symptom improvement after repeat ablation in both groups (80% (80/100) in group 1 and 77% (446/576) in group 2; p = 0.57). **Conclusion** Repeat ablation for AF in elderly patients can be performed with safety and efficacy comparable to younger patients.

Keywords Catheter ablation · Atrial fibrillation · Elderly · Repeat ablation

Introduction

Catheter ablation has emerged as an established approach to treat patients suffering from symptomatic atrial fibrillation (AF) [1]. The incidence of AF is increasing with age progression and the number of aged patients undergoing AF ablation is rising continuously [2]. AF ablation was demonstrated to be safe and effective with radiofrequency

Thomas Fink thomas_fink47@gmx.de current (RF) and cryoballoon based (CB) ablation in elderly patients [4–6]. Albeit PVI-based ablation approaches have a high likelihood for the need of repeat ablation procedures to prevent recurrence of atrial tachyarrhythmias [6, 7], to our knowledge, there are no data in larger patient cohorts on the characteristics, safety and efficacy of repeat ablations in aged patients, yet.

This study sought to analyze safety and efficacy of first repeat AF ablation procedures in elderly patients which were followed in the large prospective, multicenter German Ablation Registry.

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Extended author information available on the last page of the article

Methods

Registry structure and management

The German Ablation Registry (NCT01197638) is a prospective, multicenter non-profit registry under supervision of the "Institut für Herzinfarktforschung" (IHF, Ludwigshafen, Germany). The study was approved by local ethic boards of all participating centers. Project development and management, data acquisition and clinical monitoring were organized by the IHF. Fifty-five centers enrolled patients from January 2007 to January 2010. Patients gave written informed consent for procedure and registry participation. Data acquisition was conducted on a websitebased platform, as previously published [8].

Patient selection

Patients undergoing first repeat ablation due to arrhythmia recurrence after previous AF ablation were enrolled. Patients undergoing AV node ablation were excluded from the analysis. Patients were divided into two groups according to their age (group 1 > 70 years and group $2 \le 70$ years).

Ablation procedures and post-procedural care

Ablation procedures were performed according to institutional standards at the participating centers. Patients underwent transesophageal echocardiography (TEE) prior to the ablation procedure. Oral anticoagulation with vitamin-K antagonists was stopped before ablation procedure and replaced with low molecular-weight heparin. During procedures an activated clotting time (ACT) of 250-300 s was aimed. Procedures were performed under deep sedation using midazolam, sufentanil and/or continuous propofol infusion. The use of pre-interventional and intraprocedural imaging systems (cardiac computed tomography, cardiac magnetic resonance imaging, intracardiac echocardiography), the use of a 3D mapping system (CARTO or NavX) and the ablation system (radiofrequency (RF) or cryoballoon (CB) ablation) were at the discretion of the operators. The standard ablation protocol included assessment of PV isolation and re-ablation in case of PV reconnection to achieve persistent electrical disconnection of the PVs from the left atrium (LA). Additional ablation strategies including the creation of right atrial (RA) and LA linear lesions including block of the cavo-tricuspid isthmus (CTI), or ablation of complex fractionated atrial electrograms (CFAEs) were at the discretion of the operator. The post-procedural anticoagulation management and anti-arrhythmic drug therapy (AAD) were conducted according to local standards.

Complications were categorized in severe, moderate and minor complications (see Supplement Table 1 for further details).

Clinical follow-up

Follow-up was conducted according to local standards and included patient visits, ECG and Holter-ECG recordings. Additionally, a centralized telephonic follow-up was conducted after 12 months using a standardized protocol questioning the incidence of adverse events, arrhythmia recurrence, repeat ablations, symptoms and patients quality of life. Patients subjective perception of the ablation therapy was questioned and defined as successful, partly successful, or unsuccessful.

In case of arrhythmia recurrence documentation of the ECG or medical treatment was obtained. Adverse events (AE) were categorized in serious, moderate and related to the repeat ablation procedure (see Supplement Table 2).

Statistics

Continuous data are summarized as mean and standard deviation or median plus interquartile range (IQR; first and third quartile) in case of skewed data. Categorical data are presented as absolute and relative frequencies. Differences between the patient groups were compared with a Chi-square test or Mann–Whitney–Wilcoxon test. The Kaplan–Meier method was used to calculate 12-month event-rate of MACE (composite endpoint of death and myocardial infarction), MACCE (composite endpoint of death, myocardial infarction and stroke) and quality endpoints (composite endpoint of death, myocardial infarction, A log rank test was used to compare incidences of MACE, MACCE and quality endpoints. Statistical calculations were based on available data and cases at the timepoint of follow-up.

Results

Patient population and baseline characteristics

A total of 738 patients undergoing first repeat ablation were analyzed (Group 1 = 108 patients > 70 years, group 2 = 630 patients \leq 70 years). Median ages were 73 (71; 75) and 61 (53; 66) years (p < 0.001) and group 1 consisted of significantly more females (41 (38.0%) vs. 174 (27.6%) patients, p < 0.001). The underlying arrhythmia leading to repeat ablation was paroxysmal AF in 60 (55.6%) and 397 (63.0%) patients (p = 0.14) and persistent AF in 48 (44.4%) and 233 (37.0%) patients (p = 0.14) in groups 1 and 2, respectively. Significantly more patients in group 1 suffered from coronary artery disease and valvular heart disease was significantly more common in the elderly patients (31 (28.7%) and 17 (15.7%) patients vs. 88 (14.0%) and 50 (7.9%) patients, p < 0.001 for each parameter). There were more patients with dilated or hypertrophic cardiomyopathy in group 2 (0 (0.0%) vs. 28 (4.4%) patients, p < 0.001). Group 1 patients had did significantly more often a pacemaker, implantable cardioverter defibrillator (ICD) or cardiac resynchronization therapy (CRT) device (15 (13.9%) vs. 31 (4.9%) patients, p < 0.001). Table 1 gives an overview of patients baseline data.

Procedural parameters

Most patients underwent repeat ablation using RF ablation (698/738 patients, 94.6%). There was a statistical trend for a more frequent use of cryoballoon ablation in group 2 (2/108 (1.9%) patients in group 1 and 38/630 (6.0%) patients in group 2, p = 0.076). There were significantly less patients in group 1 undergoing manual catheter navigation (77/83 (92.8%) vs. 424/430 (98.6%) patients, p = 0.001) due to a significantly more frequent use of magnetic navigation in group 1 (5/83 (6.0%) vs. 5/430 (1.2%) patients, p = 0.003). In 1 patient of group 1 the method of catheter navigation was unknown and in an additional patient of group 2 ablation was performed under deep sedation in a comparable

Table 1Baseline patientcharacteristics

amount of the patients in group 1 and 2, respectively (97.6% (81/83) and 96.6% (402/416); p = 0.65). The remaining patients underwent repeat ablation without sedation in 1.2% (1/83) and 2.4% (1/416) (p=0.50), with invasive ventilation in 0% and 0.7% (3/416) (p=0.44) and an undetailed sedation method in 1 patient of each group (p=0.20).

A similar amount of patients underwent repeat PVI due to reconduction to the PVs in groups 1 and 2, respectively (101 (93.5%) vs. 593 (94.1%) patients, p = 0.98). Circumferential repeat PVI was conducted in 85 (78.7%) and 491 (77.9%) and segmental repeat PVI in 16 (14.8%) and 102 (16.2%) patients. The patients underwent additional RA and LA linear lesion ablation in 32 (29.6%) and 164 (26.0%) and CFAE ablation in 30/99 (30.3%) and 131/542 (24.2%) of the cases in groups 1 and 2 (p = 0.43 and p = 0.20). There was a significantly higher amount of patients undergoing ablation of linear lesions in the LA in the older patient group (25/32 (78.1%) vs. 94/164 (57.3%), p = 0.027). The procedural data are depicted in Table 2.

Acute procedure related complications during hospital stay

There was no significant difference in the incidence of severe, moderate and minor complications between the two groups (Table 2; Fig. 1). The total incidence of complications was 6.5% (7 patients with complications) and 3.8% (24 patients with complications) in groups 1 and 2 (p=0.30). Details of the incident complications are shown in Table 2

Variable	>70 years $(n = 108)$	\leq 70 years (n = 630)	<i>p</i> value
Age (years)	73 [71; 75]	61 [53; 66]	0.029
Female gender	41 (38.0)	174 (27.6)	0.029
Type of AF			
Paroxysmal	60 (55.6)	397 (63.0)	0.14
Persistent	48 (44.4)	233 (37.0)	0.14
LVEF normal ^a	76 (91.6)	467 (87.5)	0.37
CAD	31 (28.7)	88 (14.0)	< 0.001
Cardiomyopathy	0 (0.0)	28 (4.4)	0.026
Dilated cardiomyopathy	0 (0.0)	18 (2.9)	_
Hypertrophic cardiomyopathy	0 (0.0)	10 (1.6)	_
Valvular disease	17 (15.7)	50 (7.9)	0.009
Diabetes mellitus	10 (9.3)	40 (6.3)	0.27
Cardiac pacemaker	15 (13.0)	31 (4.0)	< 0.001
ICD	1 (0.9)	6 (1.0)	0.98
CRT	1 (0.9)	0 (0.0)	0.016

Values are medians [25th, 75th percentile], or n (%)

AF atrial fibrillation, *CAD* coronary artery disease, *CRT* cardiac resynchronization therapy, *ICD* implantable cardioverter defibrillator, *LVEF* left ventricular ejection fraction, *RF* radiofrequency ^aDefined as LVEF > 50%

Table 2 Procedural data

Variable	>70 years ($n = 108$)	\leq 70 years (n = 630)	p value
Procedure duration (min)	150 [120; 205]	160 [115; 210]	0.77
Fluoroscopy duration (min)	24 [16; 37]	27 [18; 45]	0.088
Radiation dose (cGy cm ²)	2666 [1426; 6135]	3390 [1809; 6238]	0.079
Mode of recurrence			
AF	108 (100)	630 (100)	1.0
Atrial tachycardia	0 (0.0)	2 (0.3)	0.56
Ablation technology			
RF	106 (98.1)	592 (94.0)	0.076
Cryoballoon	2 (1.9)	38 (6.0)	0.076
Ablation strategy			
Repeat PVI	101 (93.5)	593 (94.1)	0.98
CFAE ablation	30 (30.3) ^a	131 (24.2) ^a	0.20
Linear lesions	32 (29.6)	164 (26)	0.43
Left atrial linear lesions	25/32 (78.1)	93/164 (57.3)	0.027
Right atrial linear lesions	15/32 (46.9)	98/164 (59.8)	0.18
CTI	14/15 (93.3)	91/98 (92.9)	0.95
Rhythm at start of procedure			
Sinus rhythm	78 (72.2)	425 (67.5)	0.33
AF	30 (27.8)	205 (32.5)	0.33
Severe complication	1 (0.9)	3 (0.5)	0.56
Major bleeding requiring intervention	1 (0.9)	3 (0.5)	0.56
Moderate complication	4 (4.4)	10 (1.8)	0.12
Transitoric ischemic attack	0 (0.0)	0 (0.0)	1.0
Cardiopulmonary resuscitation	0 (0.0)	1 (0.2)	0.68
Aneurysma spurium, AV fistula	2 (1.9)	6 (1.0)	0.40
Clinically relevant pericardial effusion/car- diac tamponade	1 (1.1)	2 (0.4)	0.34
High-degree AV block	0 (0.0)	1 (0.2)	0.68
Pulmonary vein stenosis, not significant	1 (1.1)	0 (0.0)	0.014
Minor complication	2 (1.9)	11 (1.7)	0.90
Minor bleeding	2 (1.9)	10 (1.6)	0.84
New AV block I° or II°	0 (0.0)	1 (0.2)	0.68

Values are medians [25th, 75th percentile], or %

AV atrioventricular, CTI cavo-tricuspid isthmus, PVI pulmonary vein isolation

^aData only available for 99 and 542 patients

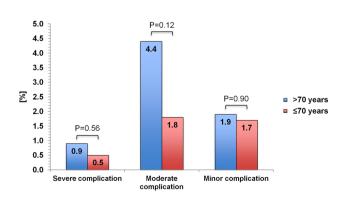


Fig. 1 Incidences of periprocedural complications

and Fig. 1. No patient died during the procedure or the hospital stay.

Arrhythmia recurrence and patients satisfaction after repeat ablation

Clinical follow-up was available for 107 and 611 patients in groups 1 and 2 (99.1% and 97.0% of the total patient population, p = 0.36). Median follow-up duration was 447 (400; 532) days in group 1 and 473 (411; 544) days in group 2 (p = 0.10). Rhythm follow-up with ECG documentation was available in 105 patients of group 1 (97.2%) and 603 patients of group 2 (95.7%). Recurrence of atrial tachyarrhythmias was documented in 45/105 (42.9%) and 252/603 (41.8%)

of the patients in each group, which was statistically not significant (p = 0.84). A similar amount of patients in each group underwent another repeat ablation procedure during the follow-up period (19/105 (18.1%) and 104/603 (17.2%) in groups 1 and 2, p = 0.83) after a median of 285 (144; 376) and 208 (103; 331) days after the first repeat procedure (p=0.050). At the end of the follow-up 40.8% (42/103) and 33.2% (191/576) of the patients were still on AAD therapy (class I, III or IV). In group 1 significantly less patients were on class I AAD, but significantly more patients on class III AAD (10/103 (9.7%) vs. 104/576 (18.1%) patients on class I, p = 0.037; 28/103 (27.2%) vs. 86/576 (14.9%) patients on class III AAD, p = 0.002). In total, a comparable 35.0% (36/103) and 42.0% (333/603) of the patients in groups 1 and 2 were without documented arrhythmia recurrence and off AAD therapy at the end of the follow-up (p=0.17).

At the end of the follow-up a comparable amount of patients stated to be free of arrhythmia-related symptoms or to have symptom improvement in both groups (80/100 patients (80%) and 446/576 patients (77%) in groups 1 and 2; p = 0.57). Additionally, a comparable amount of patients with available data stated the ablation therapy to be successful and partly successful (13/21 (61.9%) and 96/157 (61.1%) patients in groups 1 and 2, p = 0.95 and 6/21 (28.6%) and 38/157 (24.2%) patients in groups 1 and 2, p = 0.66 stated to have successful and partly successful ablation therapy, respectively). Table 3 and Fig. 2 give an overview about the follow-up data.

Patient survival and long-term safety

Kaplan–Meier estimates of 366-day incidence for MACE (death and myocardial infarction) was 0.9% and 0.2% (95% CI 0.36–91.65, log-rank p=0.16, HR 5.73), for MACCE (death, myocardial infarction and stroke) was 0.9% and 0.5% (95% CI 0.20–18.41, log-rank p=0.57, HR 1.92) and for quality endpoints (death, myocardial infarction, stroke and major bleeding) was 1.9% and 1.3% (95% CI 0.31–6.79, log-rank p=0.64, HR 1.44).

One death with unknown reason occurred during the follow-up period in each patient group (0.9% and 0.2% of all patients, p=0.69; Table 4). A total of 16 severe AE occurred in both groups during follow-up with an overall incidence of severe AE of 1.0% vs. 2.6% in groups 1 and 2 (1 major bleeding in group 1, 2 myocardial infarctions in group 2, 6 strokes in group 2 and 7 major bleedings in group 2, p=0.32). In detail, there occurred numerically more strokes in group 2 but the difference was not significantly different (0/103 (0%) vs. 6/577 (1%) of the patients, p=0.30). Moderate and late ablation procedure related AEs occurred in 5.4% (5/103)/0.0% (0/103) and 8.1% (38/472)/0.5% (3/579) of the patients in group 1 and 2, which was statistically not significantly different (p=0.39 and 0.46).

More patients in group 1 were still on oral anticoagulants at the end of the follow-up (55/103 (53%) vs. 226/576 (39%) patients, p = 0.007), which did not result in a significantly different amount of major or minor bleedings in the patient

Table 3 Clinical follow-up

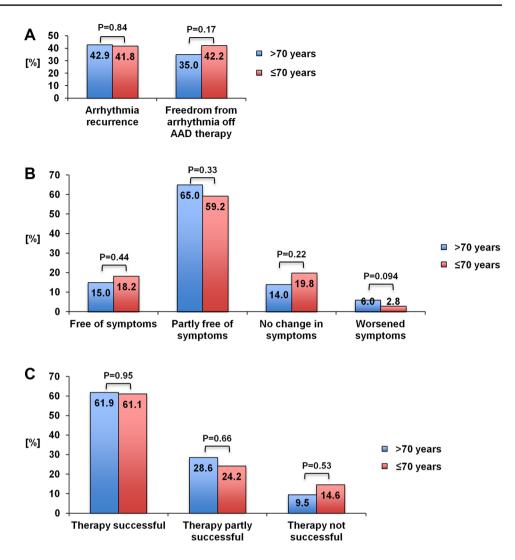
Variable	> 70 years $(n = 107)^{a}$	\leq 70 years (n=611) ^a	p value
FU duration	447.0 [400.0; 532.0]	473.0 [411.0; 544.0]	0.10
Arrhythmia recurrence	45/105 (42.9)	252/603 (41.8)	0.84
Second repeat ablation procedure	19/105 (18.1)	104/603 (17.2)	0.83
Timepoint of second repeat ablation (<i>n</i> days after first repeat ablation)	285 [144; 376]	208 [103; 331]	0.050
Rehospitalization during FU	53/103 (51.5)	249/578 (43.1)	0.11
AAD therapy at end of FU			
Class I	10/103 (9.7)	104/576 (18.1)	0.037
Class III	28/103 (27.2)	86/576 (14.9)	0.002
Class IV	4/103 (27.2)	14/576 (14.9)	0.40
Digitalis	6/103 (5.8)	20/576 (3.5)	0.25
None	65/103 (59.2)	386/576 (66.8)	0.13
Anticoagulation at end of FU			
ASS	28/103 (27.2)	156/576 (26.6)	0.90
Clopidogrel	4/103 (3.9)	11/576 (1.9)	0.21
Oral anticoagulation	55/103 (53.4)	226/576 (39.2)	0.007
Heparin, unfractionated	0 (0.0)	1/576 (0.2)	0.67
Heparin, low molecular weight	3/103 (2.9)	1/576 (0.2)	< 0.001

Values are medians [25th, 75th percentile], or n (%)

A FU follow-up, AAD anti-arrhythmic drug

^aPatient number diverges in case of missing follow-up data

Fig. 2 Clinical follow-up



cohorts with 0.9% (1/108) and 0.5% (3/630) major bleedings (p = 0.56) and 1.9% (2/108) and 1.6% (10/629) minor bleedings (p = 0.84) in groups 1 and 2, respectively.

The total incidence of re-hospitalisation during follow-up was not different between the two groups (51.5% and 43.1%, p=0.11). The median time from discharge after repeat ablation to re-hospitalisation was 214 (126; 357) and 202 (94; 346) days (p=0.74) and the reason for hospitalisation was stated as cardiovascular cause in 73.1% and 71.9% in groups 1 and 2 (p=0.86). Table 4 gives an overview on follow-up safety data.

Comparison of patients > 75 and \leq 75 years of age

Subgroup analysis of patients > 75 years of age (n=23) as compared to patients \leq 75 years of age (n=715) revealed a significantly higher incidence of CAD (9/23 (39.1%) and 110/715 (15.4%) patients, p=0.002), history of myocardial infarction (4/23 (17.4%) and 42/715 (5.9%) patients, p=0.025) and of cardiac device carriers (9/23 (39.1%) and 37/715 (5.2%) patients, p < 0.001). There was no significant difference between procedural parameters, occurrence of adverse events and follow-up parameters in patients > 75 and \leq 75 years of age except shorter duration of fluoroscopy during procedures in patients > 75 years (19 (15; 32) vs. 27 (28; 44) min, p = 0.049) and a higher amount of patients being on oral anticoagulation during clinical follow-up in the older patient group (16/22 (72.2%) and 265/657 (40.3%) patients with available follow-up, p = 0.002).

Discussion

This is the first report on real-world outcomes of first repeat AF ablation procedures in elderly patients, assessed in a large, prospectively enrolled multicenter patient cohort. Our main finding is that repeat ablation can be conducted safely in the elderly. Furthermore, repeat ablation results in a comparable procedural success, patients Table 4Follow-up safetyparameters

Variable	>70 years ($n = 107$)	\leq 70 years (n=611)	p value
Death during FU	1 (0.9)	1 (0.2)	0.69
Death, unknown reason	1 (0.9)	1 (0.2)	0.69
Rehospitalisation	53 (51.1)	249 (43.1)	0.11
Duration to rehospitalisation (days)	214 [126; 357]	202 [94; 346]	0.74
Cardiovascular reason for rehospitalisation ^a	38/53 (73.1)	179/249 (71.9)	0.86
Severe AE ^b	1 (1.0)	15 (2.6)	0.32
Myocardial infarction	0 (0.0)	2 (0.3)	0.55
Stroke	0 (0.0)	6 (1.0)	0.30
Major bleeding requiring intervention	1 (1.0)	7 (1.2)	0.83
Moderate AE ^c	5 (5.4)	38 (8.1)	0.39
Syncopy	0 (0.0)	6 (1.1)	0.30
Systemic embolism	0 (0.0)	1 (0.2)	0.67
Pulmonary embolism	1 (1.0)	0 (0.0)	0.018
Deep venous thrombosis	0 (0.0)	2 (0.4)	0.55
Cardiopulmonary resuscitation	0 (0.0)	3 (0.5)	0.46
Minor bleeding	0 (0.0)	4 (0.7)	0.40
Groin problem	3 (3.2)	14 (2.8)	0.85
Coronary revascularisation	2 (1.9)	11 (1.9)	0.98
PCI	2/2 (100)	9/10 (90)	0.64
Coronary bypass grafting	0 (0.0)	1/10 (10)	0.64
Ablation specific AE ^d	0 (0.0)	3 (0.5)	0.46
Phrenic nerve palsy	0 (0.0)	2 (0.3)	0.55
Atrio-esophageal fistula	0 (0.0)	1 (0.2)	0.68

Values are medians [25th, 75th percentile], or %

AE adverse event, FU follow-up, TIA transitoric ischemic attack

^aData only available for 52 and 249 patients

^bData only available for 103 and 577 patients

^cData only available for 92 and 472 patients

^dData only available for 103 and 579 patients

satisfaction and symptom improvement in aged patients as compared to younger patients.

Catheter ablation in the elderly: current knowledge

The majority of patients investigated in large trials of AF ablation are of younger age [6, 7] and data on AF ablation in older patients are sparse. There is no unique definition of an elderly patient but most studies used a cut-off value between 65 and 75 [4–6, 9–12].

The majority of available studies reported on patient cohorts with a limited number of individuals [4, 9, 11, 13]. A larger number of elderly patients was reported by the German Ablation Registry which analysed the first attempt of AF ablation in patients older than 75 years [5]. Two studies reported on the feasibility of more complex ablation strategies for the treatment of AF in elderly including extensive ablation of CFAEs [14, 15].

Repeat ablations in the elderly patient

To achieve long-lasting clinical success rates after AF catheter ablation repeat ablation procedures are often necessary, in particular, in patients with persistent AF [6, 7]. In contrast to the positive results of the above mentioned studies on AF ablation in elderly patients, Bunch et al. found an association between higher age and the likelihood of AF recurrence after catheter ablation [16]. Therefore, data addressing the procedural outcome and long-term data of repeat ablation for AF in the elderly are needed.

PV-reconnection was seen in a comparable number of patients in both groups and was the main driver of arrhythmia recurrence after the initial ablation procedure [6, 7]. However, aged patients in our study underwent significantly more often ablation of linear lesions within the LA in addition to repeat PVI as compared to the younger patients. Since advanced age is associated with a higher amount of atrial fibrosis in AF patients [17] this might explain why left atrial substrate modification was performed more often in the older patients.

We found a comparable long-term effectiveness in older and younger patients, which is in line with the majority of studies on first ablation procedures for atrial fibrillation [4, 9, 12–14, 18]. Younger patients were more often on class I and older on class III AAD therapy. This difference can be attributed to the higher amount of structural heart disease in the older patients of this study population. Our registry assessed the patient's satisfaction after repeat ablation and found a comparable amount of patients that termed repeat ablation as successful or at least partly successful and a comparable amount of patients reporting on improvement in subjective symptoms in both groups. Our data show that catheter ablation even with repetitive ablation procedures is a feasible treatment for symptom control in AF also for elderly patients. This finding is of major importance since there is a high likelihood to undergo a repeat procedure after arrhythmia recurrence after catheter ablation.

Beneath these favorable results of repeat ablations in older patients our data show that both, young and aged patients had a high incidence of rehospitalisation after first repeat ablation. Nearly half of the patients had hospital admission during the first 12 months after repeat ablation and nearly 20% of the patients underwent a second repeat ablation procedure. This demonstrates the ongoing need to improve the long-term efficacy of AF ablation and the need for development of new approaches in AF treatment [19].

Safety of repeat ablation

Although in most studies AF ablation in older patients was not associated with a higher complication rate, data about the safety of AF ablation in the elderly are still conflicting [3, 9-12, 20]. Studies found a higher incidence of periprocedural thromboembolic events in aged patients [5, 18]. Data from the US Nationwide Inpatient Sample demonstrated a higher incidence of periprocedural complications in patients older than 80 years [20].

We did not find a statistically significant incidence of overall periprocedural complications, in particular of major complications, in the elderly as compared to younger patients. Contrary to a previous report, which associated extensive linear ablation with an increased risk of cardiac tamponade [21], the incidence of cardiac tamponade was not higher in older patients in our study, although older patients had more often extensive ablations including left atrial linear lesions. In summary, our study shows, that repeat ablation, even when more extensive ablation is performed, can be performed with a comparable complication rate in the elderly.

Limitations

Data acquisition was conducted in a registry based on voluntary reports of procedural results and adverse events and in a non-randomized fashion. Additionally, follow-up was conducted according to local standards and we cannot exclude an influence of heterogenous methods and quality of data acquisition on study results. Nevertheless, the German Ablation Registry represents real-world data in a large patient cohort and we report the so far largest patient population in which repeat ablation and the influence of ageing on safety and procedural efficacy was studied. Since recent technological innovations like contact-force catheters [22], the use of the second-generation CB [23], laser balloon [24] or other advanced ablation techniques like rotor-mapping [25] were not available during the enrolment period of the registry, the influence of these technological advances has to be addressed in further studies. Nevertheless, we show that AF ablation in the elderly is effective and safe even without implementation of these novel technologies.

Conclusions

Repeat catheter ablation for AF can be safely performed in elderly patients with comparable results to patients in younger ages regarding periprocedural complications, long-term efficacy and patient satisfaction.

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Compliance with ethical standards

Conflict of interest Dr. Metzner received speaker's honoraria from Medtronic. Dr. Willems received speaker honoraria from Abott, Boston Scientific, Boehringer Ingelheim, Bristol Myers Squibb, Bayer Vital, Acutus and study funding from Abott, Boston Scientific and Acutus. Dr Eckardt consulting fees/speaker honoraria from Bayer Health Care, Daiichi Sankyo, Pfizer, Bristol-Myers, Squibb, Boehringer Ingelheim, Johnson&Johnson, Medtronic, Boston Scientific, Abbott, Novartis and research support by the DFG the German Heart Foundation. Dr Brachmann received consulting fees/honoraria from Biotronik, Biosense Webster, Medtronic, Boston Scientific, and St. Jude Medical. Dr Rillig received travel grants/lecture fees from Biosense, Hansen Medical, Medtronic, EP Solutions and St. Jude Medical and participated at the Boston scientific EP-fellowship. The other authors report no conflict of interest.

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