

# Safety and efficacy of long-term sodium channel blocker therapy for early rhythm control: the EAST-AFNET 4 trial

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## Aims

Clinical concerns exist about the potential proarrhythmic effects of the sodium channel blockers (SCBs) flecainide and propafenone in patients with cardiovascular disease. Sodium channel blockers were used to deliver early rhythm control (ERC) therapy in EAST-AFNET 4.

## Methods and results

We analysed the primary safety outcome (death, stroke, or serious adverse events related to rhythm control therapy) and primary efficacy outcome (cardiovascular death, stroke, and hospitalization for worsening of heart failure (HF) or acute coronary syndrome) during SCB intake for patients with ERC ( $n = 1395$ ) in EAST-AFNET 4. The protocol discouraged flecainide and propafenone in patients with reduced left ventricular ejection fraction and suggested stopping therapy upon QRS prolongation  $>25\%$  on therapy. Flecainide or propafenone was given to 689 patients [age 69 (8) years; CHA<sub>2</sub>DS<sub>2</sub>-VASc 3.2 (1); 177 with HF; 41 with prior myocardial infarction, coronary artery bypass graft, or percutaneous coronary intervention; 26 with left ventricular hypertrophy  $>15$  mm; median therapy duration 1153 [237, 1828] days]. The primary efficacy outcome occurred less often in patients treated with SCB [3/100 (99/3316) patient-years] than in patients who never received SCB [SCB<sup>never</sup> 4.9/100 (150/3083) patient-years,  $P < 0.001$ ]. There were numerically fewer primary safety outcomes in patients receiving SCB [2.9/100 (96/3359) patient-years] than in SCB<sup>never</sup> patients [4.2/100 (135/3220) patient-years, adjusted  $P = 0.015$ ]. Sinus rhythm at 2 years was similar between groups [SCB 537/610 (88); SCB<sup>never</sup> 472/579 (82)].

## Conclusion

Long-term therapy with flecainide or propafenone appeared to be safe in the EAST-AFNET 4 trial to deliver effective ERC therapy, including in selected patients with stable cardiovascular disease such as coronary artery disease and stable HF.

**Clinical Trial Registration** ISRCTN04708680, NCT01288352, EudraCT2010-021258-20, [www.easttrial.org](http://www.easttrial.org)

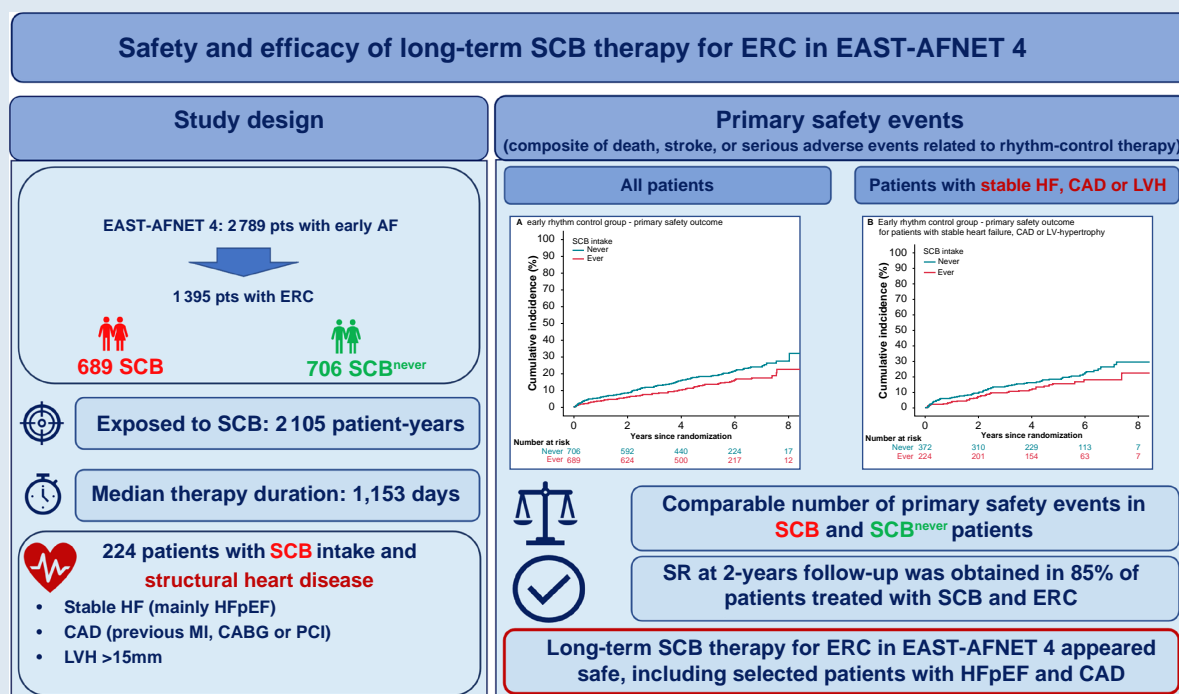
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## Graphical Abstract



SCB, sodium channel blocker; ERC, early rhythm control; HF, heart failure; CAD, coronary artery disease; MI, myocardial infarction; CABG, coronary artery bypass graft; PCI, percutaneous coronary intervention; LVH, left ventricular hypertrophy; SR, sinus rhythm.

## Keywords

Atrial fibrillation • Early rhythm control • Sodium channel blocker • Stable cardiovascular disease • Heart failure • Coronary artery disease

## What's new?

- Flecainide and propafenone were used in patients without structural heart disease and in selected patients with left ventricular hypertrophy, heart failure with preserved ejection fraction, and with revascularized coronary artery disease.
- Patients treated with flecainide or propafenone to initiate early rhythm control in EAST-AFNET 4 had fewer outcome events than patients treated with other types of rhythm control over a 5-year follow-up.
- The results might encourage the use of flecainide and propafenone in similar patients when the protocol-mandated safety precautions are followed.

## Introduction

Early rhythm control (ERC) therapy reduces cardiovascular events in patients with recently diagnosed atrial fibrillation (AF) in the EAST-AFNET 4 trial.<sup>1</sup> Beneficial effects have been observed in several sub-analyses, including in patients with heart failure (HF) and in those with a high comorbidity burden.<sup>2–8</sup> Early rhythm control therapy in the EAST-AFNET 4 trial was initially delivered using antiarrhythmic drugs in 85% of the patients.<sup>1</sup> Sodium channel blockers play a major role in antiarrhythmic drug therapy based on their effectiveness<sup>9</sup> and their low risk of extracardiac side effects.<sup>10</sup> This is even more important, considering that in the past decade no novel antiarrhythmic agent became available.<sup>11</sup> Sodium channel blocker remains underutilized, even in patients without structural heart

disease,<sup>12,13</sup> most likely due to fear of proarrhythmia.<sup>14</sup> The Cardiac Arrhythmia Suppression Trial (CAST) observed proarrhythmic effects of flecainide and encainide in patients with prior myocardial infarction, frequent ventricular premature beats, and HF with reduced ejection fraction.<sup>10,15,16</sup> These clear safety signals led to a restricted use of SCB. Whether patients with stable or revascularized coronary artery disease (CAD) and those with HF with preserved ejection fraction can be treated with SCB is not well evaluated, and current guidelines therefore slightly vary in their recommendations.<sup>17</sup> The potential underuse of SCB is specifically observed in older patients with comorbidities, patients that potentially have the most prognostic benefit from ERC therapy.<sup>4,10,18,19</sup>

To provide contemporary information on the efficacy and safety of SCB therapy, we analysed outcomes of long-term SCB therapy in the EAST-AFNET 4 patients with and without cardiovascular disease.

## Methods

The full methods of the EAST-AFNET 4 trial have been published previously.<sup>1</sup> The trial randomized 2789 patients in an international, investigator initiated, parallel-group, randomized, open, blinded outcome assessment trial design. Patients included in the trial had AF diagnosed within 12 months and at least two stroke risk factors approximating a CHA<sub>2</sub>DS<sub>2</sub>-VASc score of 2 or higher. Randomization in a one-to-one fashion to either ERC therapy ( $n = 1395$ ) or usual care (UC;  $n = 1394$ ) was performed.<sup>1</sup> Early rhythm control was selected by the site teams and consisted of antiarrhythmic drug therapy, catheter ablation, or cardioversion. The protocol discouraged SCB therapy in patients with reduced left ventricular ejection fraction (LVEF) and recommended stopping SCB therapy in patients with a QRS prolongation >25% upon therapy initiation. In patients assigned to UC, rate control

was the initial strategy and rhythm control was only initiated in patients symptomatic on optimized rate control therapy.<sup>1</sup>

The first primary efficacy outcome was a composite of death from cardiovascular cause, stroke or hospitalization with worsening of HF or acute coronary syndrome. The primary safety outcome was defined as a composite of death, stroke, or serious adverse events related to rhythm control therapy.<sup>1</sup>

All serious adverse events were prospectively captured throughout the trial. Adverse events were considered to be serious in case they resulted in death, were life-threatening, required inpatient hospitalization or prolongation of existing hospitalization, resulted in persistent or significant disability, incapacity, a congenital anomaly, or birth defect, or were judged as medically important event.<sup>1</sup>

All serious adverse events related to rhythm control therapy were centrally adjudicated as part of the primary safety outcome. The definition of 'proarrhythmia' was any arrhythmic event or an event with a potential arrhythmic background, judged as causally related to the therapeutic intervention, e.g. drug-induced proarrhythmia (torsade de pointes, ventricular tachycardia, or ventricular fibrillation), atrioventricular block, ablation-induced or drug-induced atrial arrhythmias (e.g. left atrial flutter), drug-induced bradycardia, or syncope.<sup>1</sup> Events that were judged as causally related to the therapies in the trial were considered for analysis such as drug toxicity of AF-related drug therapy, bleeding events caused by AF ablation or antithrombotic therapy, complications of ablation procedures, and others.<sup>1</sup>

Cardiovascular comorbidities were defined by the site teams at baseline and during regular follow-up visits following common clinical criteria as described in the EAST-AFNET 4 protocol (chapter 8<sup>1</sup>). In brief, stable HF was defined as presence of HF symptoms [New York Heart Association (NYHA)] class II or higher, or LVEF of <50%. Severe CAD was defined as previous myocardial infarction, coronary artery bypass graft (CABG) or percutaneous intervention (PCI); left ventricular hypertrophy (LVH) was defined as left ventricular (LV) wall thickness >15 mm (as defined via echocardiography).

All analyses reported were performed in the final, locked data set assigning patients to therapy group based on the randomization (intention-to-treat population). Data are available on reasonable request (contact: info@kompetenznetz-vorhofflimmern.de).

The protocol was approved by the ethics review boards of all institutions involved. All patients participating in the trial gave written informed consent.

## Statistics

This analysis included all 2789 patients randomized in the EAST-AFNET 4 trial and categorized patients into either SCB intake at baseline, SCB intake later during follow-up, or never SCB intake during the study period. Patients randomized to ERC ( $n = 1395$ ) were used for further analysis. As no relevant differences were observed between patients with SCB intake at baseline and SCB intake later during follow-up (see [Supplementary material online, Table S1](#)), these two groups were summarized in one group (SCB group,  $n = 689$ ) and compared to patients without any SCB intake during the study period (SCB<sup>never</sup>,  $n = 706$ ).

Patient's baseline characteristics were summarized with descriptive statistical methods. Categorical data are summarized as absolute and relative frequencies, and continuous variables were described by mean and standard deviation or median, first and third quartile.

The  $P$ -values shown are calculated from mixed linear regression models for continuous variables and mixed (ordinal) logistic regression models for categorical variables with sites included as random effect. For categorical variables with more than two categories (not ordinal), a random effect was not included.

The primary efficacy and safety outcomes of the EAST-AFNET 4 trial randomized to ERC ( $n = 1395$ ) were separately analysed for patients with SCB intake ( $n = 689$ ) or no SCB intake (SCB<sup>never</sup>,  $n = 706$ ).

For the primary efficacy outcomes and its individual components (death from cardiovascular causes, stroke, hospitalization with worsening of HF, hospitalization with acute coronary syndrome) as well as the primary safety outcomes (stroke, death and serious adverse event of special interest related to rhythm control therapy), we used multivariable Cox regression models with a time-dependent term for intake of SCB, site as a shared frailty term, for patients from the ERC group.

Additionally, the models were expanded with adjustment for age, stable HF, CAD, and type of HF by LVEF (cut-off 35%). The coefficients are expressed as hazard ratios (HRs) with a 95% confidence interval.

Furthermore, we calculated the models for the safety outcomes in patients with stable cardiovascular disease (stable severe CAD including previous myocardial infarction, CABG or PCI), stable HF, and LVH >15 mm. Statistics software R version 4.1.0. was used for all analyses.

## Results

### Baseline characteristics

Of the randomized 2789 patients included in the EAST-AFNET 4 trial, 585 (21%) patients received SCB therapy at baseline (ERC:  $n = 554$ ; UC:  $n = 31$ ), whereas 2204 patients (79%) did not. Two hundred and fifty-three patients received SCB later during the study follow-up (ERC:  $n = 135$ ; UC:  $n = 118$ ) with baselines as described in [Supplementary material online, Tables S2 and S3](#). Patients randomized to ERC ( $n = 1395$ ) were included in the analysis. Finally, overall patients with SCB intake were defined as patients with ERC treated with SCB intake (SCB,  $n = 689$ ) and compared to patients without SCB intake (SCB<sup>never</sup>,  $n = 706$ ; [Table 1](#)).

Patients with SCB intake were younger (age:  $69 \pm 8$  years vs.  $71 \pm 9$  years,  $P = 0.002$ ), were more often female [354/689 (51%) vs. 291/706 (41%),  $P < 0.001$ ], had less often stable structural heart disease such as stable HF [177/689 (26%) vs. 219/706 (31%),  $P < 0.001$ ] and severe CAD [41/689 (6.0%) vs. 202/706 (29%),  $P < 0.001$ ], and had lower CHA<sub>2</sub>DS<sub>2</sub>-VASc scores [ $3.2$  (1.3) vs.  $3.5$  (1.3),  $P < 0.001$ ] than patients without SCB intake with a similar rate of LVH [26/689 (3.8%) vs. 39/706 (5.5%),  $P = 0.37$ ; [Table 1](#)]. Differences were also observed in AF type and the number of patients in sinus rhythm at the baseline ([Table 1](#)). Detailed baseline characteristics and patient characteristics as by randomized groups are shown in [Table 1](#) and [Supplementary material online, Tables S2 and S3](#). Concomitant medical therapy showed no differences in oral anticoagulation [SCB: 625/689 (91%), SCB<sup>never</sup>: 642/700 (92%),  $P = 0.43$ ], but patients with SCB intake were less often treated with digoxin or digitoxin [16/689 (2.3%) vs. 30/700 (4.3%),  $P = 0.021$ ], mineralocorticoid receptor antagonists [25/689 (3.6%) vs. 65/700 (9.3%),  $P < 0.001$ ], diuretics [240/689 (35%) vs. 319/700 (46%),  $P < 0.001$ ], and platelet inhibitors [63/689 (9.1%) vs. 166/700 (24%),  $P < 0.001$ , [Table 1](#)].

### Duration of sodium channel blocker intake and effectiveness

Duration of SCB intake was calculated as median according to the overall duration of drug intake during the course of the study. Median treatment with propafenone or flecainide duration was 2105 patient-years and median therapy duration 1153 [237, 1828] days ([Figure 1](#), [Supplementary material online, Table S4](#)).

The number of patients in sinus rhythm at 12 months [SCB<sup>baseline</sup> 426 (88%); SCB<sup>later</sup> 111 (87%); SCB<sup>never</sup> 472 (82%)] and 24 months [SCB<sup>baseline</sup> 382 (85%); SCB<sup>later</sup> 108 (86%); SCB<sup>never</sup> 431 (79%)] was similar in patients with or without SCB intake (see [Supplementary material online, Table S5](#)).

A higher number of catheter ablations were performed in patients without SCB intake (see [Supplementary material online, Table S3](#)).

### Impact of sodium channel blocker intake on left ventricular function and New York Heart Association class

Patients with SCB intake at baseline or later had more often a normal LV function at baseline as compared to patients without SCB intake

**Table 1** Demographic and clinical characteristics of patients with and without SCB intake of patients treated with ERC

| Characteristics   | Overall, N = 1395 <sup>a</sup> | Sodium channel blocker intake Ever |                          | P-value* |
|---|--------------------------------|------------------------------------|--------------------------|----------|
|   |                                | Yes, N = 689 <sup>a</sup>          | No, N = 706 <sup>a</sup> |          |
| Age   |                                |                                    |                          | 0.002    |
| Mean $\pm$ SD   | 70 $\pm$ 8.4                   | 69 $\pm$ 8.3                       | 71 $\pm$ 8.5             |          |
| Median (IQR)  | 71 (65.0, 76)                  | 70 (65.0, 75)                      | 72 (66.0, 77)            |          |
| Gender  |                                |                                    |                          | <0.001   |
| Female  | 645/1395 (46%)                 | 354/689 (51%)                      | 291/706 (41%)            |          |
| Male  | 750/1395 (54%)                 | 335/689 (49%)                      | 415/706 (59%)            |          |
| Body mass index (calculated) (kg/m <sup>2</sup> )       |                                |                                    |                          | 0.023    |
| Mean $\pm$ SD   | 29.2 $\pm$ 5.4                 | 28.9 $\pm$ 5.2                     | 29.6 $\pm$ 5.5           |          |
| Median (IQR)  | 28.4 (25.5, 32.0)              | 28.2 (25.4, 31.5)                  | 28.7 (25.8, 32.7)        |          |
| AF type   |                                |                                    |                          | <0.001   |
| First episode   | 528/1391 (38%)                 | 244/689 (35%)                      | 284/702 (40%)            |          |
| Paroxysmal  | 501/1391 (36%)                 | 291/689 (42%)                      | 210/702 (30%)            |          |
| Persistent or long-standing persistent                  | 362/1391 (26%)                 | 154/689 (22%)                      | 208/702 (30%)            |          |
| Concomitant cardiovascular conditions                   |                                |                                    |                          |          |
| Sinus rhythm at baseline                                | 762/1389 (55%)                 | 428/689 (62%)                      | 334/700 (48%)            | <0.001   |
| Median days since AF diagnosis (IQR)                    |                                |                                    |                          | 0.86     |
| Mean $\pm$ SD   | 81.5 $\pm$ 172.5               | 79.0 $\pm$ 194.5                   | 84.1 $\pm$ 148.0         |          |
| Median (IQR)  | 36.0 (6.0, 114.0)              | 36.0 (6.0, 104.0)                  | 35.0 (6.0, 119.5)        |          |
| Absence of atrial fibrillation symptoms                 | 395/1305 (30%)                 | 180/644 (28%)                      | 215/661 (33%)            | 0.047    |
| Previous pharmacological or electrical cardioversion    | 546/1364 (40%)                 | 288/681 (42%)                      | 258/683 (38%)            | 0.83     |
| Prior AF ablation                                       |                                |                                    |                          |          |
| No  | 1395/1395 (100%)               | 689/689 (100%)                     | 706/706 (100%)           |          |
| Previous stroke or transient ischaemic attack           | 175/1395 (13%)                 | 80/689 (12%)                       | 95/706 (13%)             | 0.36     |
| At least mild cognitive impairment                      | 582/1326 (44%)                 | 267/663 (40%)                      | 315/663 (48%)            | 0.10     |
| Arterial hypertension                                   | 1230/1395 (88%)                | 606/689 (88%)                      | 624/706 (88%)            | 0.89     |
| Systolic blood pressure (mmHg)                          |                                |                                    |                          | 0.14     |
| Mean $\pm$ SD   | 137 $\pm$ 19.4                 | 136 $\pm$ 18.2                     | 137 $\pm$ 20.5           |          |
| Median (IQR)  | 135 (122.0, 150)               | 135 (124.0, 145)                   | 135 (120.0, 150)         |          |
| Diastolic blood pressure (mmHg)                         |                                |                                    |                          | 0.79     |
| Mean $\pm$ SD   | 81 $\pm$ 12.1                  | 80 $\pm$ 11.3                      | 81 $\pm$ 12.8            |          |
| Median (IQR)  | 80 (73.0, 90)                  | 80 (72.0, 90)                      | 80 (73.0, 90)            |          |
| Stable heart failure                                    | 396/1395 (28%)                 | 177/689 (26%)                      | 219/706 (31%)            | <0.001   |
| Medication at discharge                                 |                                |                                    |                          |          |
| HFrEF   | 57/396 (14%)                   | 3/177 (1.7%)                       | 54/219 (25%)             | <0.001   |
| HFmrEF  | 110/396 (28%)                  | 37/177 (21%)                       | 73/219 (33%)             | 0.28     |
| HFpEF   | 224/396 (57%)                  | 136/177 (77%)                      | 88/219 (40%)             | <0.001   |
| CHA <sub>2</sub> DS <sub>2</sub> -VASc score            |                                |                                    |                          | <0.001   |
| Mean $\pm$ SD   | 3.4 $\pm$ 1.3                  | 3.2 $\pm$ 1.3                      | 3.5 $\pm$ 1.3            |          |
| Median (IQR)  | 3.0 (2.0, 4.0)                 | 3.0 (2.0, 4.0)                     | 3.0 (3.0, 4.0)           |          |
| Chronic kidney disease of MDRD stage 3 or 4             | 172/1395 (12%)                 | 83/689 (12%)                       | 89/706 (13%)             | 0.10     |
| Severe coronary artery diseases (prev. MI, CABG or PCI) | 243/1395 (17%)                 | 41/689 (6.0%)                      | 202/706 (29%)            | <0.001   |
| Left ventricular hypertrophy on echocardiography        | 65/1395 (4.7%)                 | 26/689 (3.8%)                      | 39/706 (5.5%)            | 0.37     |
| LVEF at BL  |                                |                                    |                          | <0.001   |
| Abnormal  | 167/1364 (12%)                 | 40/680 (5.9%)                      | 127/684 (19%)            |          |
| Normal  | 1197/1364 (88%)                | 640/680 (94%)                      | 557/684 (81%)            |          |

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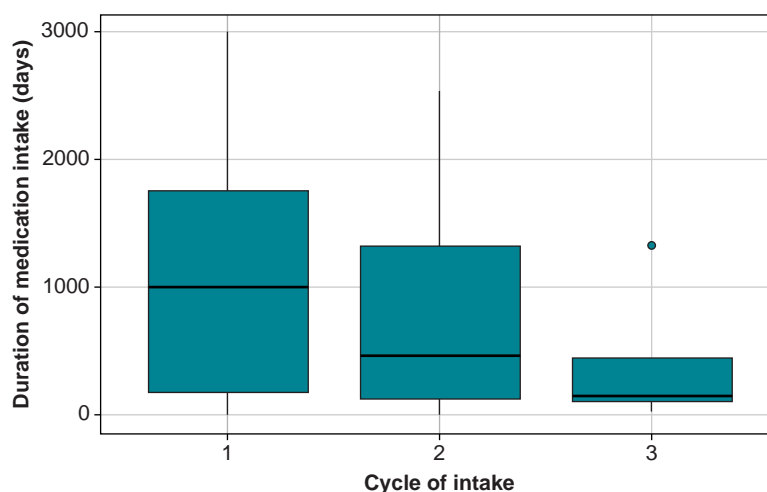
**Table 1** Continued

| Characteristics                                   | Overall, N = 1395 <sup>a</sup> | Sodium channel blocker intake Ever |                          | P-value* |
|---|--------------------------------|------------------------------------|--------------------------|----------|
|   |                                | Yes, N = 689 <sup>a</sup>          | No, N = 706 <sup>a</sup> |          |
| Oral anticoagulation with NOAC or VKA             | 1267/1389 (91%)                | 625/689 (91%)                      | 642/700 (92%)            | 0.43     |
| Digoxin or digitoxin                              | 46/1389 (3.3%)                 | 16/689 (2.3%)                      | 30/700 (4.3%)            | 0.021    |
| Beta-blockers                                     | 1058/1389 (76%)                | 537/689 (78%)                      | 521/700 (74%)            | 0.19     |
| ACE inhibitors or angiotensin II receptor blocker | 953/1389 (69%)                 | 455/689 (66%)                      | 498/700 (71%)            | 0.071    |
| Mineralocorticoid receptor antagonist             | 90/1389 (6.5%)                 | 25/689 (3.6%)                      | 65/700 (9.3%)            | <0.001   |
| Diuretic  | 559/1389 (40%)                 | 240/689 (35%)                      | 319/700 (46%)            | <0.001   |
| Statin  | 628/1389 (45%)                 | 279/689 (40%)                      | 349/700 (50%)            | <0.001   |
| Platelet inhibitor                                | 229/1389 (16%)                 | 63/689 (9.1%)                      | 166/700 (24%)            | <0.001   |
| Oral antidiabetics                                | 228/1389 (16%)                 | 102/689 (15%)                      | 126/700 (18%)            | 0.078    |
| Planned therapy for rhythm control at baseline    |                                |                                    |                          | <0.001   |
| AAD   | 1211/1395 (87%)                | 661/689 (96%)                      | 550/706 (78%)            |          |
| Ablation  | 112/1395 (8.0%)                | 18/689 (2.6%)                      | 94/706 (13%)             |          |
| None  | 72/1395 (5.2%)                 | 10/689 (1.5%)                      | 62/706 (8.8%)            |          |

ACE, angiotensin-converting enzyme; AF, atrial fibrillation; BL, baseline; CABG, coronary artery bypass graft; IQR, interquartile range; NOAC, novel oral anticoagulants; MDRD, modification of diet in renal disease; MI, myocardial infarction; PCI, percutaneous coronary intervention; SD, standard deviation; VKA, vitamin K antagonists.

<sup>a</sup>Mean (SD) or frequency with no./total no. (%)

\*P-values resulting from mixed linear regression models for metric variables and mixed (multinomial or ordinal) logistic regression models for categorical variables. For categorical variables with more than two categories (not ordinal), random effect is not included.

**Figure 1** Details of Sodium channel blocker intake for patients with first intake later or at baseline and stratified for randomized groups.

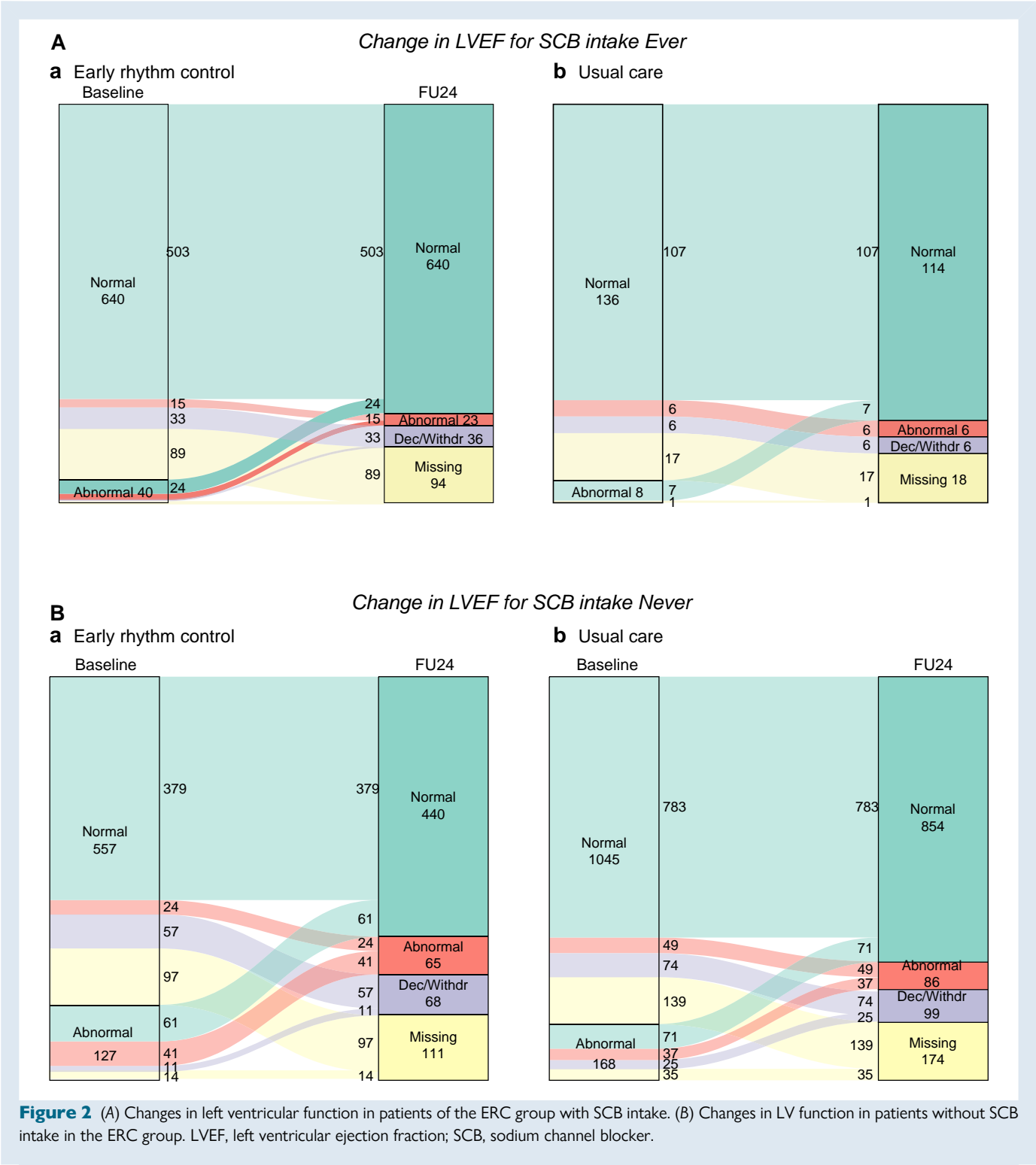
Cycle - Cycle of intake. Example: Cycle = 2 → patient took SCB for a while, then stopped and later started again. End of cycle can be end of follow-up, death, withdrawal or end of intake during follow-up on specific date.

[patients with SCB intake: 640/680 (94%) patients with normal LVEF; patients SCB<sup>never</sup>: 557/684 (81%) patients with normal LVEF; Table 1].

Of the 177 patients with SCB intake and HF, 3/177 (1.7%) patients had heart failure with reduced ejection fraction (HFrEF), 37/177 (21%) patients had heart failure with mildly reduced ejection fraction (HFmrEF), and 136/177 (77%) had heart failure with preserved ejection fraction (HFpEF).

Within the follow-up period, no relevant changes in LV function were observed in patients with or without SCB intake (Figure 2). Similar findings were found for the NYHA class with no worsening of NYHA class in any group (Figure 3). The group of patients with SCB intake comprised a lower number of patients with stable HF [i.e. SCB intake: 177/689 (26%); SCB<sup>never</sup> 219/706 (31%),  $P$ -value < 0.001], and changes in LV function or NYHA class were of similarity to those without SCB intake (Table 1, Figures 2 and 3).

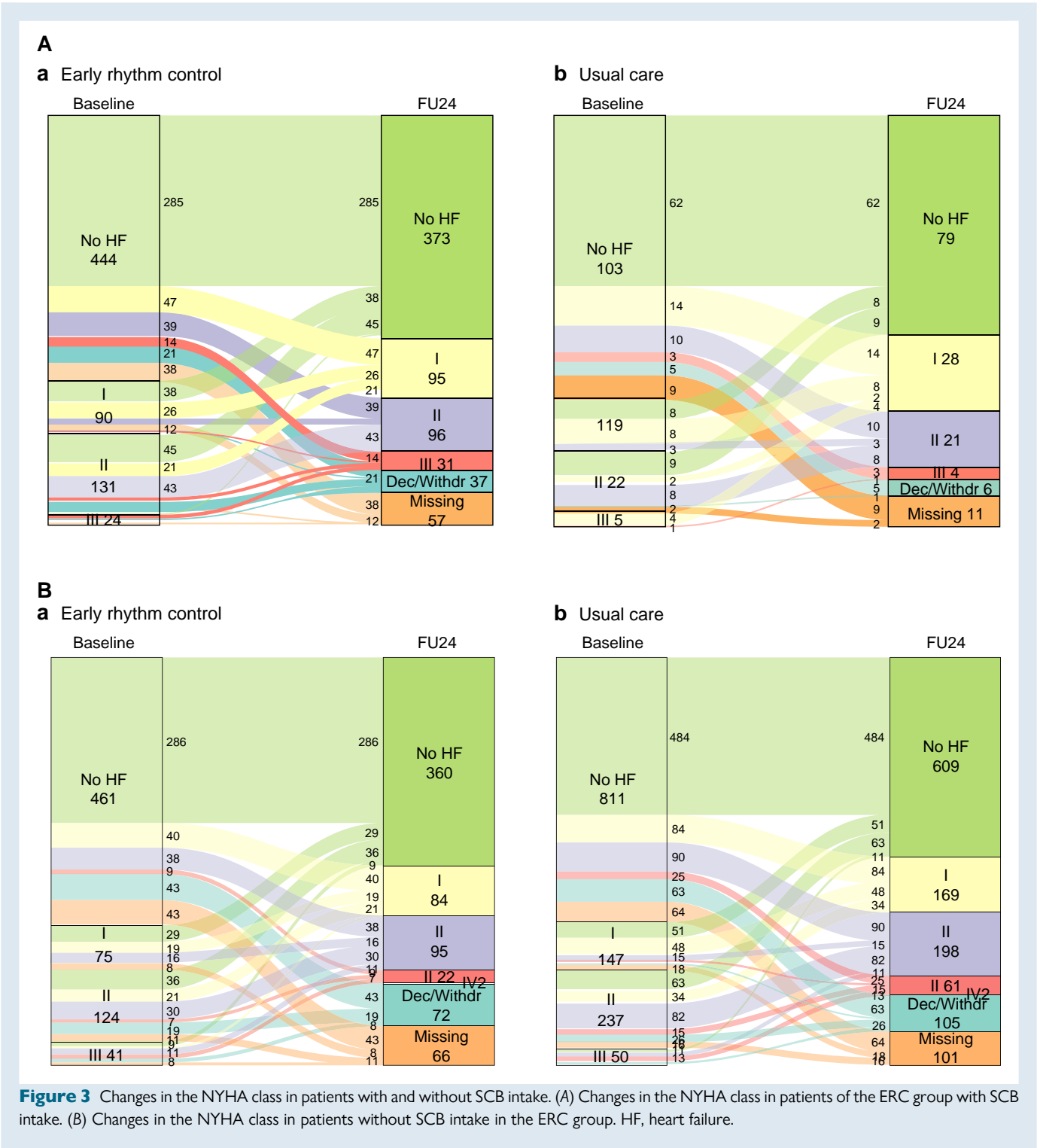




Efficacy and safety outcomes in patients with sodium channel blocker intake

The effect on the primary efficacy endpoint differed in patients with and without SCB intake. Patients with ERC on SCB had less outcomes of cardiovascular death, stroke, or hospitalization with worsening of HF or acute coronary syndrome [HR 0.55 (0.39–0.77); SCB intake: 3/100 (99/3316) patient-years; SCB<sup>never</sup> (4.9/100 (150/3083) patient-years,

multivariable Cox model  $P < 0.001$ , Table 2, Supplementary material online, Tables S6A and S6B, Supplementary material online, Figure S2) as well as for the secondary endpoints (see Supplementary material online, Table S6B). Incidence rate ratios for the second primary outcome parameter (nights spent in hospital) were lower in patients with SCB intake as compared to patients without SCB intake (see Supplementary material online, Table S7 and Supplementary material online, Figure S2).



The primary safety endpoint was numerically less often observed in patients with SCB intake as compared to SCB<sup>never</sup> patients [SCB 2.9/100 (96/3359) patient-years vs. SCB<sup>never</sup> 4.2/100 (135/3220) patient-years,  $P = 0.027$ , adjusted  $P = 0.11$ ] Table 3, Figure 4A]. When in multivariable Cox models, treatments were adjusted for age, male gender, CAD, LVH on ECG, and stable HF the primary safety endpoint and its components were observed less frequently in patients with ERC [HR 0.62 (0.45–0.86),  $P = 0.004$ ; Table 4]. Serious adverse events related to rhythm control therapy in the ERC group were observed

with similar frequency in SCB and SCB<sup>never</sup> patients [HR 0.89 (0.52–1.53),  $P = 0.685$ ].

**Changes in ECG parameters during sodium channel blocker intake**

Resting ECGs at baseline were compared to resting ECGs at 12 and 24 months, and compared between patients with SCB intake and SCB<sup>never</sup> patients (baseline ECG characteristics of patients with or without SCB

**Table 2** Cox models with time-dependent SCB intake for patients with ERC—first primary outcome and its components

| Predictors                | First primary outcome |                  |  | Death from cv causes |                  |  | Stroke           |              |  | Hospitalization worsening HF |                  |  | Hospitalization acute coronary syndrome |                  |  |
|---------------------------|-----------------------|------------------|--|----------------------|------------------|--|------------------|--------------|--|------------------------------|------------------|--|---|------------------|--|
|                           | HR (CI)               | P-value          |  | HR (CI)              | P-value          |  | HR (CI)          | P-value      |  | HR (CI)                      | P-value          |  | HR (CI)                                 | P-value          |  |
| Time-dependent SCB intake | 0.55 (0.39–0.77)      | <b>&lt;0.001</b> |  | 0.37 (0.18–0.79)     | <b>0.010</b>     |  | 0.70 (0.33–1.50) | 0.346        |  | 0.34 (0.21–0.58)             | <b>&lt;0.001</b> |  | 0.95 (0.48–1.88)                        | 0.885            |  |
| Age                       | 1.05 (1.03–1.07)      | <b>&lt;0.001</b> |  | 1.08 (1.05–1.12)     | <b>&lt;0.001</b> |  | 1.06 (1.02–1.11) | <b>0.003</b> |  | 1.06 (1.03–1.08)             | <b>&lt;0.001</b> |  | 1.01 (0.97–1.04)                        | 0.586            |  |
| Male gender               | 1.18 (0.91–1.53)      | 0.218            |  | 1.10 (0.67–1.83)     | 0.707            |  | 1.36 (0.71–2.61) | 0.362        |  | 0.98 (0.69–1.38)             | 0.890            |  | 1.27 (0.70–2.30)                        | 0.421            |  |
| CAD                       | 1.61 (1.20–2.15)      | <b>0.001</b>     |  | 1.15 (0.64–2.05)     | 0.620            |  | 1.01 (0.45–2.27) | 0.983        |  | 1.27 (0.85–1.88)             | 0.265            |  | 3.74 (2.07–6.76)                        | <b>&lt;0.001</b> |  |
| LVH on ECG                | 1.43 (0.81–2.52)      | 0.237            |  | 2.33 (0.91–5.93)     | 0.078            |  | 1.23 (0.29–5.21) | 0.799        |  | 1.02 (0.41–2.53)             | 0.960            |  | 1.89 (0.66–5.37)                        | 0.244            |  |
| Stable HF                 | 1.74 (1.35–2.26)      | <b>&lt;0.001</b> |  | 1.80 (1.10–2.96)     | <b>0.017</b>     |  | 0.71 (0.33–1.55) | 0.392        |  | 2.65 (1.89–3.71)             | <b>&lt;0.001</b> |  | 0.99 (0.54–1.81)                        | 0.974            |  |

CAD, coronary artery disease; CI, confidence interval; ECG, electrocardiogram; HF, heart failure; HR, hazard ratio; LV, left ventricular; SCB, sodium channel blocker. Values reaching statistical significance are shown in bold characters.

intake at baseline are shown in [Supplementary material online, Table S8](#)). QRS duration in baseline ECGs was slightly shorter in patients with SCB intake [SCB: 95 (17) ms, SCB<sup>never</sup>: 97 (21) ms;  $P < 0.001$ ]. No clinically relevant changes in baseline ECG characteristics at 12 and 24 months were observed (see [Supplementary material online, Table S9](#)).

**Safety of sodium channel blocker intake in patients with coronary heart disease, stable heart failure, and left ventricular hypertrophy**

Stable HF, prior myocardial infarction, PCI, or CABG, and LVH > 15 mm were observed in 596 patients of the ERC group (SCB:  $n = 224$ ; SCB<sup>never</sup>:  $n = 372$ ; [Table 1](#)). In those 224 patients with SCB intake, stable HF was observed in 177 patients, prior myocardial infarction, PCI, or CABG in 41 patients, and LVH >15 mm in 26 patients ([Table 1](#)). There were numerically similar primary safety outcomes in patients receiving SCB with previous myocardial infarction, CABG, or PCI and stable HF or LVH [34 (15.2%)] than in patients not receiving SCB [74 (19.9%), [Table 4](#)]. However, as outlined above, when assessed in multivariable Cox models the primary safety endpoint and its components were observed in fewer frequency in patients with ERC [HR 0.62 (0.45–0.86),  $P = 0.004$ ; [Table 5](#)]. To substantiate the safety of SCB therapy, we performed a separate safety analysis including all patients who received SCB including those who received SCB as part of UC. The overall safety was comparable (see [Supplementary material online, Tables S10 and S11](#)).

**Discussion**

This analysis provides information on the long-term safety and effectiveness of the SCBs flecainide and propafenone as part of ERC therapy in patients with AF and stroke risk factors. These findings include safety information in selected patients with HFpEF and with stable or revascularizedCAD. The study provides an increase in information on the safety of flecainide and propafenone, substances that have mainly been used in patients with no or only a few cardiovascular diseases.<sup>9,16,20</sup> The results might encourage the use of flecainide and propafenone in similar patients when safety precautions are followed, including assessment of QRS duration with swift action to halt drug therapy in the case of extensive QRS prolongation upon therapy.

**Long-term sodium channel blocker treatment in clinical practice**

Although SCB has shown high efficacy in reducing AF burden and maintaining sinus rhythm, precautions still exist to prescribe antiarrhythmic drugs (AADs), especially in patients with higher age and higher comorbidity burden.<sup>12,18</sup> The reservations against using SCB mainly originate from the CAST and CAST II, where SCB intake (flecainide, moricizine, and encainide) was associated with a 2.5-fold excess mortality in patients with previous myocardial infarction and a high burden of premature ventricular contractions. Mortality was significantly higher in patients with non-Q-wave infarction as compared to patients with Q-wave infarction with a 5-time higher relative risk of mortality. Further analysis in CAST revealed that acute ischaemia served as one of the main triggers for lethal tachyarrhythmias.<sup>15,21</sup> The findings of CAST have led to an Food and Drug Administration (FDA) recommendation that labels flecainide use to be contraindicated in all patients with structural heart disease of any aetiology.<sup>16</sup> However, patients with (untreated or treated) stable CAD or HF with preserved ejection fraction or mildly reduced ejection fraction without prior myocardial infarction were not studied in CAST.<sup>15,21</sup> There are also few data on the safety of



**Table 3** Primary safety endpoint of patients with (ever) or without (never) SCB intake in patients with ERC or UC

|   | ERC       |            |          |               | UC        |            |
|---|-----------|------------|----------|---------------|-----------|------------|
|   | Ever      | Never      | P-value* | P-value adj** | Ever      | Never      |
| n   | 689       | 706        |          |               | 149       | 1245       |
| Primary composite safety outcome  | 96 (13.9) | 135 (19.1) | 0.027    | 0.11          | 20 (13.4) | 203 (16.3) |
| Stroke  | 17 (2.5)  | 23 (3.3)   | 0.438    | 0.496         | 7 (4.7)   | 55 (4.4)   |
| Death   | 45 (6.5)  | 93 (13.2)  | < 0.001  | 0.001         | 9 (6.0)   | 155 (12.4) |
| Serious adverse event of special interest related to rhythm control therapy | 34 (4.9)  | 34 (4.8)   | 0.783    | 0.587         | 6 (4.0)   | 13 (1.0)   |
| Serious adverse event related to antiarrhythmic drug therapy                |           |            |          |               |           |            |
| Non-fatal cardiac arrest  | 1 (0.1)   | 0 (0.0)    | 0.851    | 1             | 0 (0.0)   | 1 (0.1)    |
| Drug toxicity of AF-related drug therapy                                    | 5 (0.7)   | 5 (0.7)    | 0.969    | 0.835         | 2 (1.3)   | 1 (0.1)    |
| Drug-induced bradycardia  | 8 (1.2)   | 6 (0.8)    | 0.561    | 0.525         | 1 (0.7)   | 4 (0.3)    |
| Atrioventricular block  | 1 (0.1)   | 1 (0.1)    | 0.968    | 0.477         | 0 (0.0)   | 0 (0.0)    |
| Torsade de pointes tachycardia  | 1 (0.1)   | 0 (0.0)    | 1        | 1             | 0 (0.0)   | 0 (0.0)    |
| Serious adverse event related to AF ablation                                |           |            |          |               |           |            |
| Pericardial tamponade   | 1 (0.1)   | 2 (0.3)    | 0.585    | 0.36          | 0 (0.0)   | 0 (0.0)    |
| Major bleeding related to AF ablation                                       | 1 (0.1)   | 5 (0.7)    | < 0.001  | 0.88          | 0 (0.0)   | 0 (0.0)    |
| Non-major bleeding related to AF ablation                                   | 1 (0.1)   | 0 (0.0)    | 0.9      | 1             | 1 (0.7)   | 1 (0.1)    |
| Serious adverse event of special interest related to RC therapy             |           |            |          |               |           |            |
| Blood pressure-related event  | 0 (0.0)   | 1 (0.1)    | 1        | 0.95          | 0 (0.0)   | 0 (0.0)    |
| Hospitalization for AF  | 4 (0.6)   | 7 (1.0)    | 0.432    | 0.896         | 1 (0.7)   | 2 (0.2)    |
| Other cardiovascular event  | 1 (0.1)   | 4 (0.6)    | 0.222    | 0.349         | 1 (0.7)   | 0 (0.0)    |
| Other event   | 1 (0.1)   | 0 (0.0)    | 0.831    | 0.993         | 1 (0.7)   | 2 (0.2)    |
| Syncope   | 3 (0.4)   | 1 (0.1)    | 0.23     | 0.264         | 0 (0.0)   | 1 (0.1)    |
| Hospitalization for worsening of HF with Decomp HF                          | 2 (0.3)   | 1 (0.1)    | 0.22     |               | 0 (0.0)   | 0 (0.0)    |
| Implantation of a pacemaker, defibrillator, or other                        | 5 (0.7)   | 3 (0.4)    | 0.614    | 0.789         | 0 (0.0)   | 4 (0.3)    |

AF, atrial fibrillation; HF, heart failure; RC, rhythm control; UC, usual care.

\*Mixed logistic regression models with a random effect for site were used for comparison of intake at ever vs. never for patients with ERC treatment.

\*\*Mixed logistic regression models with a random effect for site were used for comparison of intake at ever vs. never for patients with ERC treatment adjusted for age, stable heart failure, CAD, and type of heart failure by LVEF (cut-off 35).

SCBs in patients with LVH or in those with HF with preserved ejection fraction.<sup>10,16,22,23</sup> The recommendations of the current ESC guidelines for the management of patients with ventricular arrhythmias and the prevention of sudden cardiac death provide more flexibility for SCB treatment also in patients with structural heart disease, when no myocardial infarction has been reported.<sup>24,25</sup>

## Considerations for the safety of long-term sodium channel blocker intake in patients with structural heart disease

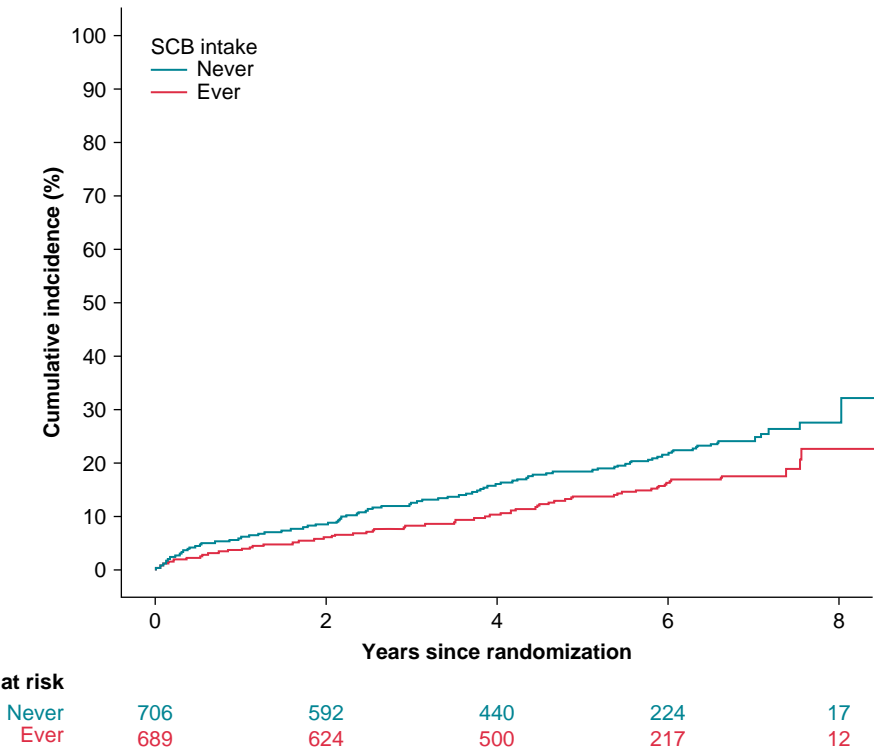
The Flec-SL trial has shown that long-term use of flecainide is more effective as compared to short-term use after electrical cardioversion with a comparable safety profile.<sup>9</sup> However, long-term SCB use in the Flec-SL trial was defined as an intake of no longer than 6 months and patients with a reduced LV function <40% were excluded.<sup>9</sup> This underlines the need for additional data from large prospective patient cohorts for long-term safety of SCB use in patients with and without stable cardiac comorbidities. Recent analyses, obtained from non-randomized cohorts, have shown that flecainide does not show an increased rate of proarrhythmia or HF events in patients with stable or revascularized CAD when compared to the treatment with class III AADs.<sup>26</sup> In addition,

experimental data have demonstrated only limited impact of flecainide and propafenone on voltage-gated potassium channels.<sup>27</sup>

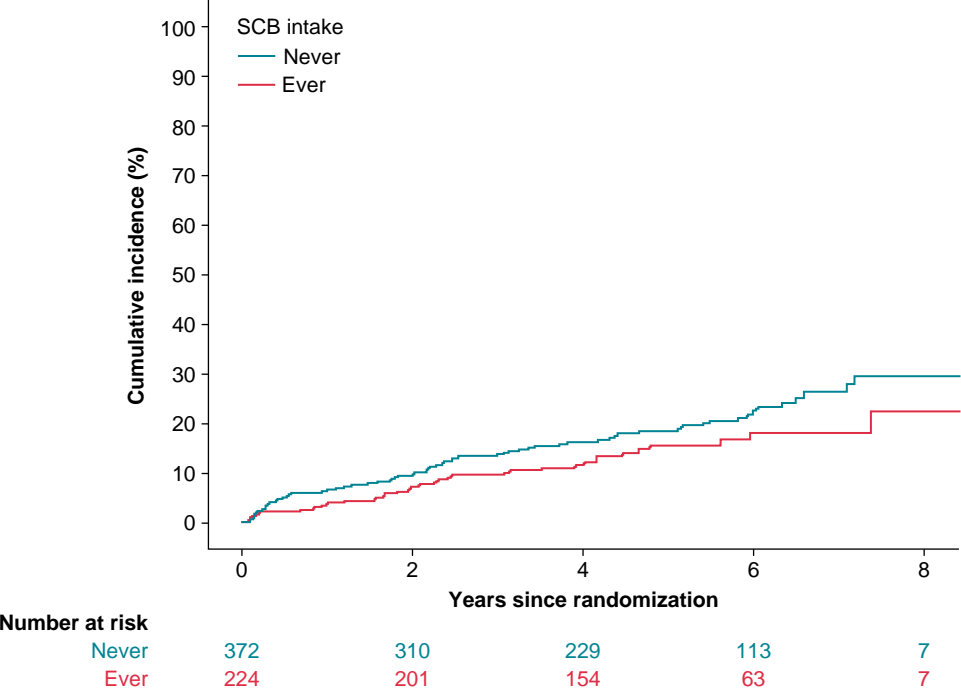
Specific trials have shown that antiarrhythmic drugs remain effective after AF ablation.<sup>28</sup> The original trials of propafenone and flecainide tested their use in patients not undergoing AF ablation. Of note, in the POWDER-AF trial patients treated with antiarrhythmic drugs, mainly based on SCBs, after catheter ablation did not show a higher number of adverse events related to antiarrhythmic drug therapy during a 1-year follow-up period.<sup>28</sup>

In the EAST-AFNET 4 trial, rhythm control was obtained using AADs in the majority of patients (>85%), although SCB therapy considered as the primary initial treatment in patients randomized to ERC in the EAST-AFNET 4 trial was higher (>40%)<sup>1</sup> than the final treatment with SCB (21% of patients at baseline, Table 1). The present subanalyses provide detailed insights into the safety and efficacy of long-term SCB intake in the EAST-AFNET 4 population. Several primary safety events were reported in patients treated with SCB in the present subanalyses, but events potentially related to AAD treatment such as bradycardia, torsade de pointes tachycardia or sudden cardiac death as well as life-threatening events were rarely seen in both groups (Table 2). Remarkably, similar event rates of the primary safety endpoint were observed in patients with and without stable structural heart disease, which suggests that patients with stable heart disease including stable

A Early rhythm control group — primary safety outcome



B Early rhythm control group — primary safety outcome for patients with stable heart failure, CAD, or LV hypertrophy



**Figure 4** Cumulative incidence of the primary safety outcome in all patients with SCB intake (A) and patients with stable cardiovascular disease (severe CAD, HF, and LV hypertrophy) (B) in the ERC group. CAD, coronary artery disease; LV, left ventricular; SCB, sodium channel blocker.

**Table 4** Cox models with time-dependent SCB intake for patients with ERC—safety outcomes

| Predictors                | Primary composite safety outcome |                  | Death            |                  | SAE of special interest related to RC therapy |       |
|---------------------------|----------------------------------|------------------|------------------|------------------|---|-------|
|                           | HR (CI)                          | P                | HR (CI)          | P                | HR (CI)                                       | P     |
| Time-dependent SCB intake | 0.62 (0.45–0.86)                 | <b>0.004</b>     | 0.40 (0.24–0.68) | <b>0.001</b>     | 0.89 (0.52–1.53)                              | 0.685 |
| Age                       | 1.07 (1.05–1.09)                 | <b>&lt;0.001</b> | 1.09 (1.07–1.12) | <b>&lt;0.001</b> | 1.03 (1.00–1.06)                              | 0.055 |
| Male gender               | 1.10 (0.84–1.44)                 | 0.483            | 1.39 (0.97–1.98) | 0.074            | 0.74 (0.45–1.22)                              | 0.243 |
| CAD                       | 1.05 (0.76–1.46)                 | 0.760            | 0.99 (0.65–1.50) | 0.961            | 1.14 (0.60–2.17)                              | 0.683 |
| LVH on ECG                | 1.85 (1.08–3.16)                 | <b>0.022</b>     | 2.20 (1.13–4.25) | <b>0.017</b>     | 1.56 (0.56–4.36)                              | 0.401 |
| Stable HF                 | 1.26 (0.95–1.66)                 | 0.112            | 1.52 (1.06–2.16) | <b>0.022</b>     | 1.15 (0.68–1.95)                              | 0.595 |

CAD, coronary artery disease; CI, confidence interval; HF, heart failure; HR, hazard ratio; LVH, left ventricular hypertrophy; RC, rhythm control; SAE, serious adverse event; SCB, sodium channel blocker.

Values reaching statistical significance are shown in bold characters.

**Table 5** Primary safety outcomes in patients with stable cardiovascular comorbidities (stable CAD, stable HF, LVH >15 mm) stratified for SCB intake at baseline, later SCB intake, and no SCB intake

|   | Early rhythm control |           |          |               | Usual care |            |
|---|----------------------|-----------|----------|---------------|------------|------------|
|   | Ever                 | Never     | P-value* | P-value adj** | Ever       | Never      |
| n   | 224                  | 372       |          |               | 42         | 550        |
| Primary composite safety outcome  | 34 (15.2)            | 74 (19.9) | 0.557    | 0.622         | 6 (14.3)   | 109 (19.8) |
| Stroke  | 4 (1.8)              | 13 (3.5)  | 0.233    | 0.401         | 4 (9.5)    | 22 (4.0)   |
| Death   | 18 (8.0)             | 51 (13.7) | 0.121    | 0.166         | 1 (2.4)    | 86 (15.6)  |
| Serious adverse event of special interest related to rhythm control therapy | 12 (5.4)             | 18 (4.8)  | 0.604    | < 0.001       | 1 (2.4)    | 10 (1.8)   |
| Serious adverse event related to antiarrhythmic drug therapy                |                      |           |          |               |            |            |
| Non-fatal cardiac arrest  | 0 (0.0)              | 0 (0.0)   |          |               | 0 (0.0)    | 1 (0.2)    |
| Drug toxicity of AF-related drug therapy                                    | 1 (0.4)              | 3 (0.8)   | 0.607    | 0.348         | 0 (0.0)    | 1 (0.2)    |
| Drug-induced bradycardia  | 4 (1.8)              | 3 (0.8)   | 0.295    | 0.342         | 0 (0.0)    | 3 (0.5)    |
| Atrioventricular block  | 1 (0.4)              | 0 (0.0)   | 0.996    | 0.996         | 0 (0.0)    | 0 (0.0)    |
| Torsade de pointes tachycardia  | 1 (0.4)              | 0 (0.0)   | < 0.001  | 1             | 0 (0.0)    | 0 (0.0)    |
| Serious adverse event related to AF ablation                                |                      |           |          |               |            |            |
| Pericardial tamponade   | 1 (0.4)              | 0 (0.0)   | 0.865    |               | 0 (0.0)    | 0 (0.0)    |
| Major bleeding related to AF ablation                                       | 1 (0.4)              | 3 (0.8)   | 0.607    | 0.927         | 0 (0.0)    | 0 (0.0)    |
| Non-major bleeding related to AF ablation                                   | 1 (0.4)              | 0 (0.0)   | 0.926    | 1             | 1 (2.4)    | 0 (0.0)    |
| Serious adverse event of special interest related to RC therapy             |                      |           |          |               |            |            |
| Blood pressure-related event  | 0 (0)                | 0 (0)     |          |               | 0 (0)      | 0 (0)      |
| Hospitalization for AF  | 1 (0.4)              | 5 (1.3)   | 0.312    |               | 0 (0.0)    | 2 (0.4)    |
| Other cardiovascular event  | 1 (0.4)              | 2 (0.5)   | 0.45     | 0.588         | 0 (0.0)    | 0 (0.0)    |
| Other event   | 0 (0.0)              | 0 (0.0)   |          |               | 0 (0.0)    | 2 (0.4)    |
| Syncope   | 0 (0.0)              | 1 (0.3)   | 1        | 1             | 0 (0.0)    | 1 (0.2)    |
| Hospitalization for worsening of HF with Decomp HF                          | 0 (0.0)              | 1 (0.3)   | 1        |               | 0 (0.0)    | 0 (0.0)    |
| Implantation of a pacemaker, defibrillator, or other                        | 1 (0.4)              | 2 (0.5)   | 0.268    | 0.198         | 0 (0.0)    | 3 (0.5)    |

AF, atrial fibrillation; HF, heart failure.

\*Mixed logistic regression models with a random effect for site were used for comparison of intake at BL vs. never for patients with ERC treatment.

\*\*Mixed logistic regression models with a random effect for site were used for comparison of intake at BL vs. never for patients with ERC treatment adjusted for age, stable HF, CAD, and type of HF by LVEF (cut-off 35).

or revascularized CAD were safely treated with SCB therapy in the EAST-AFNET 4 trial unless otherwise contraindicated. Sinus rhythm at the 12- and 24-month follow-up was similar in patients with or

without SCB use in the ERC group. However, patients not treated with SCB were often treated with other effective antiarrhythmic drugs such as amiodarone or dronedarone.

## Safety of long-term sodium channel blocker intake in patients with coronary artery disease, left ventricular hypertrophy, and heart failure

In the EAST-AFNET 4 trial, patients with unstable angina, untreated CAD, or unstable HF were excluded, but a relevant number of patients with stable CAD were randomized. According to the findings of these subanalyses, SCBs were safely applied in this patient population of the EAST-AFNET 4 trial as safety events were observed only in a minority of these patients and lethal complications such as cardiovascular death and life-threatening arrhythmias were rare (Table 5).

Apparently, in our subanalyses, primary safety events were not more often observed in patients with stable HF as compared to patients without. Furthermore, LV function and NYHA class remained stable in the majority of patients and did less often worsen during follow-up when compared with patients without structural heart disease (Figures 2 and 3); neither relevant impairment of systolic LV function nor an increase of the NYHA class was observed in any of the subgroups with SCB intake. The observations mainly apply to patients with preserved LV function. These findings show that patients with stable cardiac comorbidities receiving SCB therapy did not have more safety events than patients treated with other AADs in the EAST-AFNET 4 trial supporting early medical rhythm control in these patients with high efficacy and a low risk of harm. Of note, patients in the EAST-AFNET 4 trial were treated with the recommended SCB dose (200 mg flecainide/day, 600 mg propafenone/day), whereas clinical practice tends to prescribe lower doses.<sup>29</sup>

## Strengths and limitations

This is a post hoc subgroup analysis of the prospective randomized EAST-AFNET 4 trial, and therefore, although obtained from a large international randomized multicentre cohort, the results remain hypothesis-generating. Sodium channel blocker intake varied during study participation resulting in some patients with continuous SCB intake and others with on/off SCB therapy. The term severe CAD was defined as previous myocardial infarction, CABG, or PCI; however, detailed information about the severity of the disease (single-/multivessel disease as well as presence of untreated stenoses of the coronary arteries) was not available for analysis. Although the available information, especially the normal global LV function, suggests that only patients with small myocardial infarctions were treated with SCBs in the EAST-AFNET 4, no information on exercise testing and no information on the type, size, or location of previous myocardial infarction were available. The suitability for SCB therapy was assessed by the local study team. The main outcome of this analysis is the safety of SCB therapy in the trial without mandated exercise testing or routine angiography. A majority of patients with HF had HFpEF; the definition of HF in patients with ejection fraction <50% was based on symptoms and therefore provides limited granularity. Similarly, the definition that the authors use for LVH does not consider the underlying aetiology.

As flecainide therapy alone might accelerate ventricular conduction during AF and could result in 1:1 flutter with high ventricular rates, concomitant  $\beta$ -blocker therapy is recommended due to its AV node slowing effects. In the EAST-AFNET 4 trial, 1:1 atrial flutter was rarely observed. The high use of concomitant  $\beta$ -blocker therapy in the SCB group (flecainide-only-treated patients 78% and propafenone-only-treated patients 80%) might have contributed to the encouraging results for a safe and effective long-term use of flecainide in the present subanalyses. The low overall number of safety events precluded a meaningful analysis of specific patient features that may be associated with safety events with and without SCB therapy. Much larger data bases, e.g. stemming from merged electronic health records and prescribing information, may address this topic.

No information to the actual dosage of the medications can be provided. However, recommended dosing of SCBs was defined in the study protocol according to the AF guidelines (flecainide daily dose 200–300 mg, propafenone daily dose 450–600 mg).<sup>1,10</sup>

Of note, the results have to be interpreted with caution due to differences in age and cardiovascular comorbidities of the SCB therapy group with other patients, making comparison more difficult. The main finding of this analysis is the long-term safety of therapy with flecainide and propafenone, including in selected patients deemed unsuitable for these drugs. In addition, patients in the SCB group were less often treated with digoxin which may have contributed to the observed safety profile.<sup>30–32</sup>

Nonetheless, patients in this analysis were treated for a long time period with a median SCB intake of 2105 patient-years (median therapy duration 1153 [237, 1828] days), providing robust information on the long-term effectiveness and safety of SCB in ERC therapy in patients with AF with and without stable structural heart disease so far.

Although sensitivity analyses were performed considering age, stable HF, CAD, and type of HF as stratified by LVEF, we cannot exclude other confounders in the cohort of non-SCB intake, as patients in the SCB group had a higher comorbidity burden. This might at least in part explain why the primary safety endpoint in patients with SCB intake was less often observed than in patients not treated with SCB. Some patients initiated SCB later in the trial, but the overall findings mainly apply to patients with relatively recently diagnosed AF.

## Conclusion

The findings of this subanalysis in selected patients of the EAST-AFNET 4 trial show that no safety signals were observed during SCB therapy for ERC therapy in patients with AF with or without stable cardiovascular disease such as CAD, LVH, or stable HF (mainly patients with HFpEF) in the EAST-AFNET 4 trial.

## Supplementary material

Supplementary material is available at *Europace* online.

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## Data availability

Data are available on reasonable request (contact: info@kompetenznetz-vorhofflimmern.de).

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