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Recurrent ischemic stroke and bleeding in patients with atrial fibrillation who suffered an acute stroke while on treatment with NOACs: the RENO-EXTEND study.

Short title: RENO Extend Study

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Abstract

Background: In patients with atrial fibrillation (AF) who suffered an ischemic stroke while on treatment with NOACs, rates and determinants of recurrent ischaemic events and major bleedings remain uncertain.

Methods: This prospective multicenter observational study aimed to estimate the rates of ischaemic and bleeding events and their determinants in the follow-up of consecutive patients with AF who suffered an acute cerebrovascular ischaemic event while on NOAC treatment. Afterwards, we compared the estimated risks of ischaemic and bleeding events between the patients in whom anticoagulant therapy was changed to those who continued the original treatment.

Results: After a mean follow-up time of 15.0±10.9 months, 192 out of 1240 patients (15.5%) had 207 ischemic or bleeding events corresponding to an annual rate of 13.4%. Among the events, 111 were ischaemic strokes, 15 systemic embolisms, 24 intracranial bleedings and 57 major extracranial bleedings. Predictive factors of recurrent ischemic events (strokes and systemic embolisms) included CHA_2DS_2-VASc score after the index event [OR 1.2 (95% CI 1.0-1.3) for each point increase, p=0.05] and hypertension [OR 2.3 (95% CI 1.0-5.1), p=0.04]. Predictive factors of bleeding events (intracranial and major extracranial bleedings) included age [OR 1.1 (95% CI 1.0-1.2) for each year increase, p=0.002], history of major bleeding [OR 6.9 (95% CI 3.4-14.2), p=0.0001] and the concomitant administration of an antiplatelet agent [OR 2.8 (95% CI 1.4-5.5), p=0.003]. Rates of ischemic and bleeding events were no different in patients who changed or not changed the original NOACs treatment [OR 1.2 (95% CI 0.8-1.7)].

Conclusions: Patients suffering a stroke despite being on NOAC therapy are at high risk of recurrent ischemic stroke and bleeding. In these patients, further research is needed to improve secondary prevention by investigating the mechanisms of recurrent ischemic stroke and bleeding.

Non-standard Abbreviations and Acronyms

NOAC   Non-vitamin K antagonist oral anticoagulant

AF   Atrial fibrillation

OR   Odds ratio

HR   Hazard ratio

PSM   Propensity Score Matching

Introduction

In patients with AF, oral anticoagulation reduces the risks of ischemic stroke and systemic embolism in patients with AF by 60–70%. (1) and Non–vitamin K antagonist oral anticoagulants (NOACs) are currently the gold standard for this clinical indication (2,3). In a meta-analysis of randomized trials (RCT), the residual risk of ischemic stroke in patients treated with NOACs was estimated at 1.4% per year (4); observational studies have reported an annual risk of 8.9% (7.3–10.8) in patients with AF on anticoagulant treatment (5-7). Several possible mechanisms for this residual risk of stroke have been suggested, including an absent or reduced adherence to anticoagulation, non-cardioembolic stroke etiologies, or cardioembolic mechanisms different from AF. Identifying one or more of these mechanisms could allow for a more targeted and individualized approach for secondary prevention. (8,9). To this regard, the results of the recent "Causes and Risk Factors of Cerebral Ischemic Events in Patients With Nonvalvular AF Treated With NOACs for Stroke Prevention (RENO)" study suggested that the prescription of reduced off-label dose of NOACs, atrial enlargement, hyperlipidemia and CHA_2DS_2-VASc were independently associated with high ischemic stroke rates. (10). Currently, guidelines on managing stroke recurrences under NOACs are lacking as no specific RCTs have addressed the issue of changing or not the NOAC administered at the time of stroke.

In light of the aforementioned, the RENO-EXTEND study sought to estimate the rates of ischemic and bleeding events after an acute ischemic stroke in patients with AF while on treatment with NOACs and investigate for predictive factors of these events. Subsequently, we compared the estimated risks for ischemic and bleeding events between those who had or not their anticoagulant therapy changed.

Methods

The data that supported the findings of this study are available from the corresponding author upon reasonable request.

In this multicentre observational cohort study performed between January 2018 and December 2020, consecutive acute cerebrovascular ischemic patients with AF who were taking NOAC treatment at the time of the event were prospectively collected. Patients were enrolled from 43 Stroke Unit across Europe and United States (25 from academic hospitals and 18 from non-academic hospitals) all with high expertise regarding the management of patients with stroke. Patients who had suspended anticoagulant therapy at least 24 hours before the cerebrovascular event for any reason and patients who did not guarantee adherence to treatment were excluded. To verify compliance, the patients and family members were asked how the prescribed anticoagulant was taken.

Each research center was free to change or not the NOAC treatments.

The study was approved by the pertinent institutional review boards if required. Informed consent was obtained according to local requirements.

This study was designed following recommendation of the STROBE statement (see online-only Data Supplement).

Clinical, radiological and laboratory risk factors

Data on known risk factors for stroke were collected as previously described (11,12): see online-only Data Supplement.

The doses of NOACs were recorded, and any reasons for prescribing reduced doses were also collected. Reduced doses of NOACs were considered off label in the absence of the recommended clinical and laboratory criteria for dose reduction (13-16).

Both the CHA_2DS_2-VASc score and the HAS-BLED score before and after the index event were also calculated (17) (see online-only Data Supplement).

Lead investigators were required to report on the type and dosage of prescribed NOACs prescribed and the vascular management. Moreover, physicians were also required to perform neuroradiological examinations: cerebral computed tomography (CT) or cerebral magnetic

resonance imaging (MRI). Any recurrent ischemic stroke, symptomatic systemic embolism, intracranial bleeding, and death were recorded during this period.

Etiology of the index stroke

For the causes of stroke, the A-S-C-O-D (A, atherosclerosis; S, small vessel disease; C, cardiac pathology; O, other causes; D, dissection) classification was adopted (18). A-S-C-O-D phenotyping assigns a degree of likelihood of causal relationship to every potential disease (1 for potentially causal, 2 for causality is uncertain, 3 for unlikely causal but the disease is present, 0 for the absence of disease, and 9 for an insufficient workup to rule out the disease). The cause of the index event was considered cardioembolic in the case of [A(0,2,3)-S(0,2,3)-C1-O(0,2,3)] according to the A-S-C-O-D classification.

Outcome evaluation

Patients were followed up prospectively by face-to-face or telephone interviews. The duration of follow-up was at least 12 months. Patient follow-ups started at the moment of index events. The initiation of oral anticoagulation followed international guideline recommendations (2).

The primary outcome measure was the composite of ischemic stroke, systemic embolism, intracranial bleeding, and major extracranial bleeding. Recurrent stroke was defined as the sudden onset of a new focal neurological deficit of vascular origin in a site consistent with the territory of a major cerebral artery and was categorized as ischemic or hemorrhagic. Intracranial bleeding was defined as a spontaneous hemorrhagic stroke (intraparenchymal bleeding), subdural or subarachnoid hemorrhage. Traumatic intracranial bleeding was not considered as an outcome event. Systemic embolism was defined as an acute vascular occlusion of an extremity or organ confirmed by imaging at either surgery or autopsy.

Death data were recorded; functional recovery was assessed by the modified Rankin Scale (mRS), dichotomizing between functional independence (mRS 0-2) and disability (mRS≥3).

Major extracranial bleeding was defined as a reduction in the hemoglobin level of 2g per deciliter or more, the requirement of a blood transfusion of at least 2 units, or symptomatic bleeding in either a critical area or organ.

Follow-up visits and outcome adjudication were performed by local investigators, not in a blinded fashion.

Statistical analysis

For patients with or without outcome events, differences in clinical characteristics and risk factors were calculated using the chi-square test of proportions (with a two-sided alpha level of 5%). Ninety-five percent confidence intervals were calculated for odds ratios (OR). A second step analysis aimed to identify predictors of outcome events among baseline characteristics. Univariate tests were performed to compare clinical characteristics and risk factors associated with stroke on admission in patients with and without outcome events. Multivariable logistic regression analysis was performed to identify independent predictors for outcome events. The variables included in this analysis were the following: CHA_2DS_2-VASc score after admission for the index stroke (separately as a continuous variable or including the risk factors within the score excluding the CHA_2DS_2-VASc score), hyperlipidemia, current alcohol abuse, current smoking habit, paroxysmal AF, presence of malignancy, antiplatelet therapy in addition to NOACs after the index stroke, cardioembolic index stroke, history of previous major bleeding and changes in the type of anticoagulant therapy after the index stroke.

To compare the risk of outcome events in those patients who did not have their anticoagulant therapy changed after the index event and those who changed their anticoagulant therapy, the relation between the survival function and the set of explanatory variables were calculated by Cox proportional hazard models. These models provide an estimate of the treatment effect on survival after an adjustment for other explanatory variables. The same Cox proportional hazard analyses were performed in patients diagnosed with a cardioembolic ischemic index stroke to compare the

risk of outcome events in those patients who had had their anticoagulant therapy changed after the index event and those who had not had their anticoagulant therapy modified. The results of these analyses were reported as hazard ratios (HRs).

Using Propensity Score Matching (PSM), a further analysis was performed to compare the overall risk of outcome events in those patients who did not have their anticoagulant therapy modified after their index events versus those who underwent modifications of anticoagulant therapy. In this PSM, survival function and empirical cumulative hazard function were estimated via the Kaplan–Meier estimator for the two groups; any differences between survival functions were tested using the Log-rank statistic (or Mantel–Haenszel test), that in the case of large samples has an asymptotic Chi-square distribution (19). Patients were censored at the time of an outcome event, death, or if they had been lost to follow-up. Data were analyzed utilizing the SPSS/PC Win package 25.0.

Sample size calculation

To perform a logistic regression analysis, we needed at least 10 patients with outcome for each variable included in the model (20). The expected outcome event rate at 12 months was estimated to be 11% (9). In light of the above, to evaluate the predictors of the primary outcome events, it was calculated that 910 patients would have been needed; meaning that at least 10 variables were required for inclusion in the model to address a sufficient level of confounding

Results

Characteristics of the patients

Overall, 1,300 consecutive patients who suffered an acute ischemic stroke while on treatment with NOACs were included in the study. Of these, 1,240 patients were included in the analysis (10 patients were excluded due to incomplete data, and 50 were lost to follow-up, and there were no data for them) (Figure S1 in the online-only Data Supplement). Of these, 16 patients had their NOACs changed, while 15 did not and 19 did not receive NOACs. The characteristics of the

10

included patients are reported in Table I in the online-only Data Supplement. Out of 1,240 patients, 491 (39.6%) were appropriately treated with low-dose NOACs before the index event, and 149 of these (30.3%) were treated with a non-label low dose.

Concerning the causes of the index events, 920 (74.2%) were diagnosed as cardioembolic strokes, according to the A-S-C-O-D classification.

After the index event, 490 patients treated with an appropriate dose of NOAC (39.5%) had their NOACs changed (68 switched from dabigatran to factor Xa inhibitor, 139 from factor Xa inhibitor to dabigatran and 283 switched to another Xa inhibitor). In contrast, 527 treated with appropriate dose (42.5%) continued with the same NOAC at the same dose, 83 (6.7%) continued with the same NOAC but had the dose increased, 58 (4.7%) were shifted to warfarin and 82 (6.6%) were shifted to low molecular weight heparin and never prescribed oral anticoagulants for the following reasons: early ischemic stroke recurrence, early hemorrhagic transformation, early death or severe index stroke. The characteristics of the patients subdivided by the types of treatment received after the index event are summarized in Table II in the online-only Data Supplement while patterns of switch are revealed by Sankey diagram in Figure S2 in the online-only Data Supplement.

On multivariable analysis, predictive factors associated with change of NOACs included CHA_2DS_2-VASc score [OR 1.4 (95% CI 1.1-1.8) for each point increase, p=0.001] and index event of cardioembolic origin [OR 2.4 (95% CI 1.4-4.1), p=0.001], while adding an antiplatelet to a NOAC was associated with retaining the preexisting NOAC regimen [OR 0.5 (95% CI 0.3-1.0), p=0.05]. When including in the model the different research centres, similar results were obtained for CHA_2DS_2-VASc score [OR 1.3 (95% CI 1.1-1.6) for each point increase, p=0.001] and for index event of cardioembolic origin [OR 2.9 (95% CI 1.8-4.7), p=0.001].

Rates and predictive factors of ischemic and bleeding events

After a mean follow-up time of 15.0±10.9 months (1,550 patient-years) (median 12 months, Interquartile Range 12-15), 192 patients (15.5%) had 207 outcome events corresponding to an annual rate of 13.4%. The following events were observed: 111 ischemic strokes, 15 systemic embolisms, 24 intracranial bleedings and 57 were major extracranial bleedings. The ischemic and bleeding events according to the type of treatment are summarized in Table III in the online-only Data Supplement.

On multivariable analysis, predictive factors for thromboembolic and bleeding events were history of major bleeding [OR 4.1 (95% CI 2.2-7.6), p=0.0001], the addition of an antiplatelet to NOACs [OR 1.7 (95% CI 1.1-2.9), p=0.03], age [OR 1.0 (95% CI 1.0-1.0) for each year increase or OR 1.2 (1.0-1.4) for each decade increase p=0.045], male sex [OR 1.5 (95% CI 1.0-2.2), p=0.04] and hypertension [OR 2.2 (95% CI 1.1-4.2), p=0.02].

Predictive factors of ischemic events (ischemic stroke and systemic embolism) included CHA_2DS_2-VASc score after the index event [OR 1.2 (95% CI 1.0-1.3) for each point increase, p=0.05] and hypertension [OR 2.3 (95% CI 1.0-5.1), p=0.04].

Predictive factors of haemorrhagic events (intracranial and major extracranial bleedings) included age [OR 1.1 (95% CI 1.0-1.2) for each year increase, p=0.002], history of major bleeding [OR 6.9 (95% CI 3.4-14.2), p=0.0001] and the addition of an antiplatelet to a NOAC [OR 2.8 (95% CI 1.4-5.5), p=0.003]. Figure 1 and Table IV in the online only Data Supplement describe the events in those patients who had or not, antiplatelet therapy associated with NOAC after index event.

In patients with a history of stroke or TIA, the rate of the combination of thromboembolic and bleeding events was 16.8% (79/471) compared to 14.7% (113/769) of the patients without a history of stroke or TIA (p=0.1).

Outcome events in patients who continued anticoagulant therapy unchanged after the index event compared to those who changed the type of NOAC.

The Cox regression curve analyses comparing the overall outcome events in patients who did not have their NOAC changed after the index event (n=527) with those who changed the type of their NOACs (n=490) are reported in Figure 2. There was no difference in the rate of the primary outcome [HR of 1.1 (95% CI 0.8-1.4)], or of ischemic outcome events [HR of 1.1 (95% CI 0.7-1.4)] or bleeding outcome events [HR of 1.4 (95% 0.7-2.5)]. Comparing strategies of anticoagulation after the index event, it resulted that, all the strategies were significantly less associated with the primary outcome, when comparing the results from a NOAC to the low molecular weight heparin switch (Log Rank – Mantel-Cox, p=0.01 vs NOAC-to-NOAC switch and NOAC dosage change, p=0.003 vs NOAC-to-warfarin switch, and p=0.005 vs no NOAC change) (Figure S3 in the online only Data Supplement). No observed differences emerged between NOAC-to-NOAC change strategies. The Cox regression curve analyses that compared the outcome events in patients with cardioembolic stroke as index event who did not have their type of NOAC changed after the index event, with those who underwent NOAC type changes are reported in Figure S4 in the online only Data Supplement. Out of the 920 patients who had a cardioembolic stroke as an index event, 401 changed their type of NOAC after the index event, whereas 351 did not: 58 (14.5%) from the former group had a combined outcome event, compared to 44 (12.5%) from the latter group (p=0.4). Furthermore, 34 (8.6%) of the patients who had their NOAC type changed had an ischemic outcome event, compared to 27 (7.7%) of the patients who did not undergo such a change (p=0.7). Concerning the bleeding outcome events, no differences were observed between the groups: 24 (5.9%) and 17 (4.8%), respectively (p=0.1).

After PSM, 421 patients who did not have their type of NOAC changed were compared with 421 patients who did it. In Table V in the online-only Data Supplement, the characteristics of the patients after PSM are reported.

Concerning outcome events, 67 (15.9%) of the patients who had their NOAC type changed had a combined outcome event, compared to 59 (14.1%) of those who did not (Log Rank – Mantel-Cox, p=0.4). Furthermore, 41 (9.7%) of the patients who had their NOAC type changed had an ischemic outcome event, compared to 37 (8.8%) of the patients who did not (Log Rank – Mantel-Cox, p=0.6). Concerning bleeding outcome events, a difference between the groups: 26 (6.2%) and 22 (5.2%), respectively (Log Rank – Mantel-Cox, p=0.5), was not observed.

The Kaplan-Meier curve that compared the combined outcome events (ischemic stroke, systemic embolism, intracranial haemorrhage and major extracranial) in patients who did not have their type NOAC changed with those who did not after PSM is reported in Figure S5 in the online-only Data Supplement.

Discussion

The results of this study in patients with the previous stroke while on treatment with NOACs, showed a combined rate of thromboembolic and bleeding events of 16.7%, corresponding to an annual rate of 13.4%. These rates are higher than those observed in previous RCTs: 2.83% in the ENGAGE AF-TIMI 48 for edoxaban (21); 2.32% in RELY for Dabigatran 110 mg BID (RR 0.84; [95% CI: 0.58, 1.20]) and 2.07% for 150 mg BID (22); 1.01% in ARISTOTLE for apixaban (15) and 2.79% in ROCKET AF for rivaroxaban (23). An individual patient data pooled analysis of 7 prospective real-world cohort studies on AF patients with recent cerebral ischemia, who had been on anticoagulation treatment, reported an annual risk for recurrent ischemic stroke and death of 4.7% and 10.2%, respectively, which is in line with our study (5). This high residual stroke risk needs to be quantified for targeted treatment in these patients. The currently used CHA_2DS_2-VASc score may not be adequate to quantify the residual stroke risk as it weighs the history of previous strokes with a score of only two. Furthermore, the CHA_2DS_2-VASc score does not acknowledge an added risk associated with every stroke recurrence in patients already under anticoagulation treatment.

The residual risk for stroke is often considered a treatment failure or related to etiologies other than cardioembolic. So, the first step in these patients is excluding non-cardioembolic causes that would lead the physician to adjust to the original treatment strategy. In clinical practice, one of the options for patients with non-cardioembolic recurrences and intra-or-extracranial atherosclerosis and/or small vessel disease may be to add antiplatelet therapy. However, this regimen was associated with an increase in the risk of major bleeding, with only a small stroke prevention benefit (24). Likewise, our study found that add-on antiplatelets significantly increased the risk of bleeding in line with the AUGUSTUS trial and by Capodanno et al. results on patients undergoing percutaneous coronary intervention, leading us to suggest the avoidance of prescribing the combination therapy including dual antiplatelet therapy and NOACs (25,26).

In this study, we observed that changing the type of anticoagulant after the index event was not associated with a decreased risk of ischemic stroke recurrence, as reported by two observational studies (5,6). These no responders to anticoagulant stroke therapy have an increased risk of ischemic events for each point increase of CHA_2DS_2-VASc score. Such high-risk patients might benefit from alternative preventive options, including left atrial appendage occlusion (27) even if there is a lack of data regarding the benefit of this procedure in patients on NOACs with a prior stroke, but only in those patients undergoing cardiac surgery.

The RENO-EXTEND study has several limitations: (1) it was observational, and neither individual NOAC type nor their doses were randomized; (2) other pharmacological treatments besides NOACs were not investigated. However, interactions between NOACs and other drugs are reported to be much lower than those of warfarin; (3) we have no information about the plasma level of NOACs; (4) the sample size was calculated to investigate for predictive factors associated with ischemic and bleeding events, but not to compare the risks of these events in patients who had and those did not have their anticoagulation molecule changed; (5) 50 patients were lost at follow-up; (6) as the reduced dose criteria are different in the label in Europe and USA, we adopted a single definition of the appropriate dose.

The strengths of our study include its large sample size and its prospective design. Additionally, our analyses reflect real-life experiences and thus may provide valuable information that could significantly reduce the incidence of ischemic events in patients with AF and stroke during NOAC therapy.

In conclusion, patients having a stroke despite being on therapy with NOACs are at high risk of recurrent ischemic stroke and bleeding. Further research is needed to investigate mechanisms of recurrent stroke and improve secondary prevention in these patients.

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Supplemental Materials:

Expanded Methods

Tables I-V

Figures S1-S2

STROBE checklist

References:

1. EAF Trial Study Group. Secondary prevention in non-rheumatic AF after TIA or minor stroke. Lancet 1993;342:1255–1262.

2. Klijn CJ, Paciaroni M, Berge E, Korompoki E, Korv J, Lal A, Putaala J, Werring DJ. Antithrombotic treatment for secondary prevention of stroke and other thromboembolic events in patients with stroke or TIA and non-valvular AF. Eur Stroke J 2019;4:198-223.

3. Hindricks G, Potpara T, Dagres N, Arbelo E, Bax JJ, Blomstrom-Lundqvist C, Boriani G, Castella M, Dan GA, Dilaveris PE et al. 2020 ESC Guidelines for the diagnosis and management of AF developed in collaboration with the European Association for Cardio-Thoracic Surgery: The Task Force for the diagnosis and management of AF of the ESC developed with the special contribution of the EHRA of the ESC. Eur Heart J 2021;42:373-498.

4. Ruff CT, Giugliano RP, Braunwald E, Hoffman EB, Deenadayalu N, Ezekowitz MD, Camm AJ, Weitz JI, Lewis BS, Parkhomenko A et al. Comparison of the efficacy and safety of new oral anticoagulants with warfarin in patients with AF: a meta-analysis of randomized trials. Lancet. 2014;383:955-962.

5. Seiffge DJ, De Marchis GM, Koga M, Paciaroni M, Wilson D, Cappellari M, Macha Md K, Tsivgoulis G, Ambler G, Arihiro S et al. Ischemic Stroke despite Oral Anticoagulant Therapy in Patients with AF. Ann Neurol. 2020;87:677-87.

6. Yaghi S, Henninger N, Giles JA, Leon Guerrero C, Mistry E, Liberman AL, Asad D, Liu A, Nagy M, Kaushal A et al. Ischemic stroke on anticoagulation therapy and early recurrence in acute cardioembolic stroke. J Neurol Neurosurg Psychiatry 2021;92:1062-1067.

7. Tanaka K, Koga M, Lee KJ, Kim BJ, Park EL, Lee J, Mizoguchi T, Yoshimura S, Cha JK, Lee BC et al. AF associated ischemic stroke patients with prior anticoagulation have higher risk of recurrent stroke. Stroke 2020;51:1150-1157.

8. Rota E, Testa L, Di Brigida G, Agosti S, Rovere ME, Risso R, Morelli N. The management of patients with acute ischemic stroke while on DOACs: data from an Italian cohort and a proposed algorithm. J Thromb Thrombolysis 2020;50: 732-738.

9. Stretz C, Wu TY, Wilson D, Seiffge DJ, Smith EE, Gurol ME, Yaghi S. Ischemic stroke in anticoagulated patients with AF. J Neurol Neurosurg Psychiatry 2021; 92:1164-1172.

10. Paciaroni M, Agnelli G, Caso V, Silvestrelli G, Seiffge DJ, Engelter S, De Marchis GM, Polymeris A, Zedde ML, Yaghi S et al. Causes and Risk Factors of Cerebral Ischemic Events in Patients With AF Treated With NOACs for Stroke Prevention. Stroke. 2019;50:2168-2174.

11. Paciaroni M, Agnelli G, Falocci N, Caso V, Becattini C, Marcheselli S, Rueckert C, Pezzini A, Poli L, Padovani A et al. Early Recurrence and Cerebral Bleeding in Patients With Acute Ischemic Stroke and AF: Effect of Anticoagulation and Its Timing. Stroke 2015;46:2175-2182.

12. Paciaroni M, Agnelli G, Falocci N, Tsivgoulis G, Vadikolias K, Liantinioti C, Chondrogianni M, Bovi P, Carletti M, Cappellari M et al. Early Recurrence and Major Bleeding in Patients With Acute Ischemic Stroke and AF Treated With NOACs. J Am Heart Assoc 2017;6e007034.

13. Connolly SJ, Ezekowitz MD, Yusuf S, Eikelboom J, Oldgren J, Parekh A, Pogue J, Reilly PA, Themeles E, Varrone J et al. Dabigatran versus warfarin in patients with AF. N Engl J Med. 2009;361:1139–1151

14. Patel MR, Mahaffey KW, Garg J, Pan G, Singer DE, Hacke W, Breithardt G, Halperin JL, Hankey GJ, Piccini JP et al. Rivaroxaban versus warfarin in nonvalvular AF. N Engl J Med. 2011;365:883–891.

15. Granger CB, Alexander JH, McMurray JJ, Lopes RD, Hylek EM, Hanna M, Al-Khalidi HR, Ansell J, Atar D, Avezum A et al. Apixaban versus warfarin in patients with AF. N Engl J Med. 2011;365:981–992.

16. Giugliano RP, Ruff CT, Braunwald E, Murphy SA, Wiviott SD, Halperin JL, Waldo AL, Ezekowitz MD, Weitz JI, Spinar J et al. Edoxaban versus warfarin in patients with AF. N Engl J Med 2013;369:2093–2104.

17. Pisters R, Lane DA, Nieuwlaat R, De Vos CB, Crijns HJ, Lip GY. A Novel User-Friendly Score to Assess 1-Year Risk of Major Bleeding in Patients with AF. Chest 2010;138:1093–100.

18. Amarenco P, Bogousslavsky J, Caplan LR, Donnan GA, Wolf ME, Hennerici MG. The ASCOD phenotyping of ischemic stroke (Updated ASCO Phenotyping). Cerebrovasc Dis. 2013;36:1–5.

19. Mantel N. Evaluation of survival data and two new rank order statistics arising in its consideration. Cancer Chemother Rep. 1966;50:163–170.

20. Reboldi G, Angeli F, Verdecchia P. Multivariable analysis in cerebrovascular research: practical notes for the clinicians. Cerebrovasc Dis 2013;35:187-193.

21. Rost N, Giugliano RP, Ruff, CT, Murphy SA, Crompton AE, Norden AD, Silverman S, Singhal AB, Nicolau JC, Somaraju B et al. Outcomes with edoxaban versus warfarin in patients with previous cerebrovascular events: findings from ENGAGE AF-TIMI 48. Stroke 2016;47:2075-2082.

22. Diener HC, Connolly SJ, Ezekowitz MD, Wallentin L, Reilly PA, Yang S, Xavier D, Di Pasquale G, Yusuf S. Dabigatran compared with warfarin in patients with AF and previous TIA or stroke: a subgroup analysis of the RE-LY trial. Lancet Neurol 2010;9:1157-63.

23. Hankey GJ, Patel MR, Stevens SR, Becker RC, Breithardt G, Carolei A, Diener HC, Donnan GA, Halperin JL, Mahaffey KW et al. Rivaroxaban compared with warfarin in patients with AF and previous stroke or TIA: a subgroup analysis of ROCKET AF. Lancet Neurol 2012;11:315-322.

24. So, C.H., Eckman, M.H. Combined aspirin and anticoagulant therapy in patients with AF. J Thromb Thrombolysis 2017;43:7–17.

25. Lopes RD, Heizer G, Aronson R, Vora AN, Massaro T, Mehran R, Goodman SG, Windecker S, Darius H, Li J et al. Antithrombotic therapy after acute coronary syndrome or PCI in AF. N Engl J Med 2019;380:1509-1524.

26. Capodanno D, Huber K, Mehran R, Lip GYH, Faxon DP, Granger CB, Vranckx P, Lopes RD, Mantalescot G, Cannon CP et al. Management of antithrombotic therapy in AF patients undergoing PCI. J Am Coll Cardiol 2019;74:83-99.

27. Whitlock RP, Belley-Cote EP, Paparella D, Healey JS, Brady K, Sharma M, Reents W, Budera P, Baddour AJ, Fila P et al. Left atrial appendage occlusion during cardiac surgery to prevent stroke. N Engl J Med 2021;384:2081-2091.
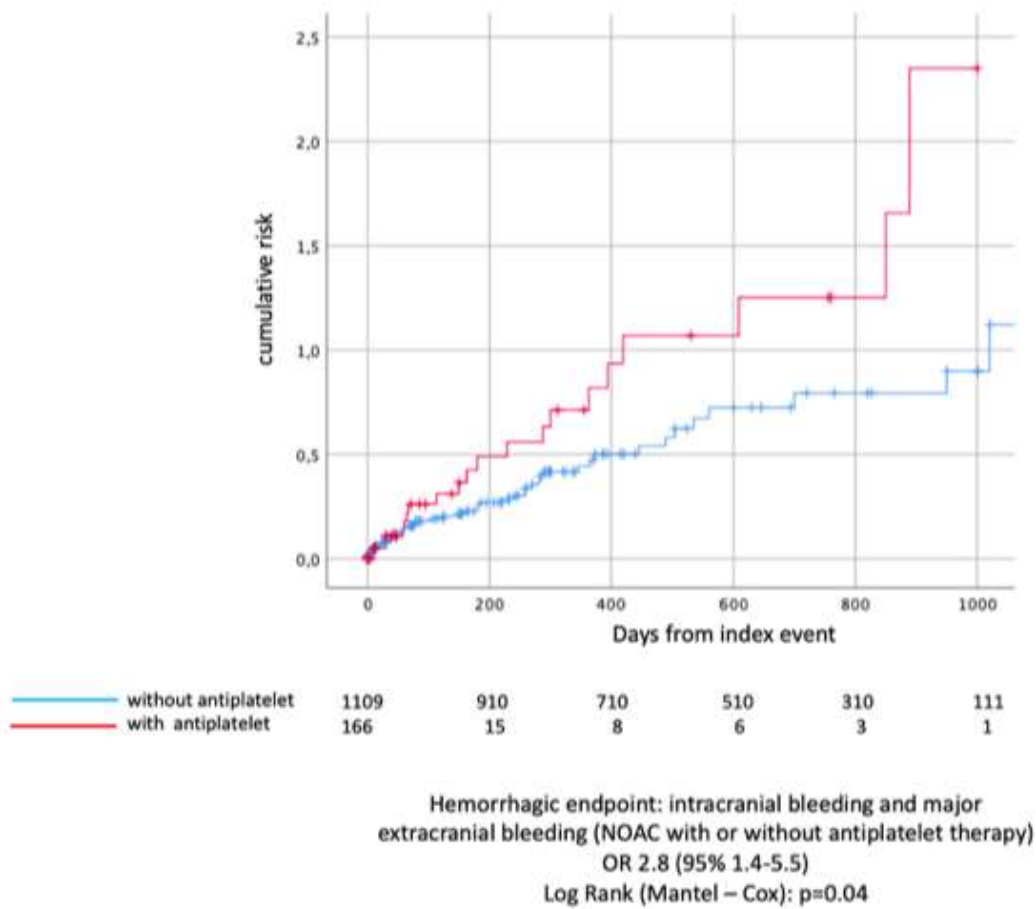
Figure legend

Hemorrhagic endpoint: intracranial bleeding and major
extracranial bleeding (NOAC with or without antiplatelet therapy)
OR 2.8 (95% 1.4-5.5)
Log Rank (Mantel – Cox): p=0.04

Figure 1: Hemorrhagic endpoint events (intracranial hemorrhage and major extracranial

hemorrhage) in patients who had or not, associated antiplatelet therapy to NOAC in the overall
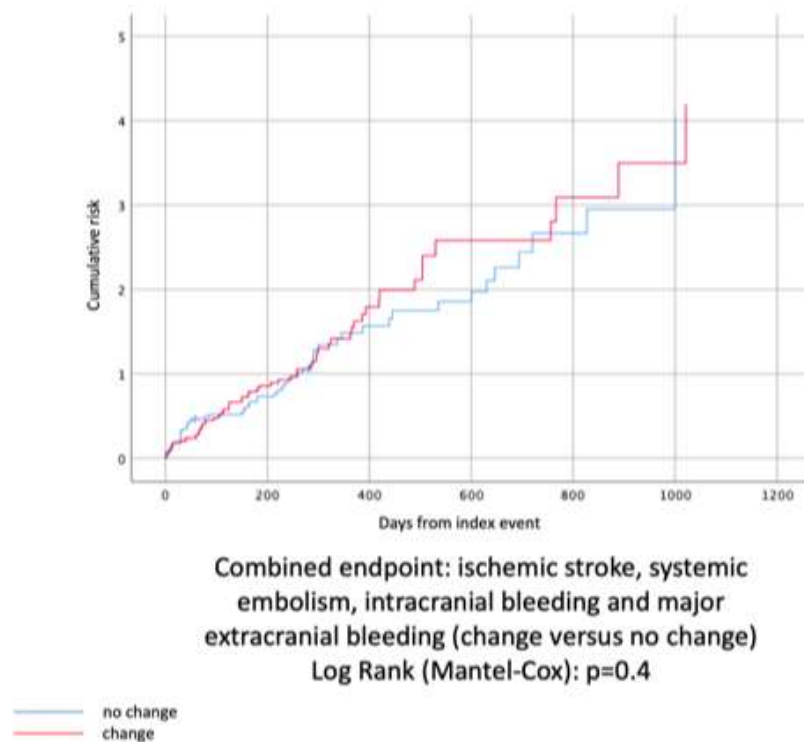
population

Figure 2

Combined endpoint: ischemic stroke, systemic
embolism, intracranial bleeding and major
extracranial bleeding (change versus no change)
Log Rank (Mantel-Cox): p=0.4

Figure 2: Cox regression curve analyses comparing the combined outcome events (ischemic stroke, systemic embolism, intracranial hemorrhage and major extracranial hemorrhage) in patients who did not have their NOAC changed after the index event with those who changed the type of their NOACs in the overall population.