Microbleed Prevalence and Burden in Anticoagulant-Associated Intracerebral Bleed

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Microbleed prevalence and burden in anticoagulant-associated intracerebral bleed

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Abstract

Prior studies suggest an association between Vitamin K antagonists (VKA) and cerebral microbleeds (CMBs); less is known about nonvitamin K oral anticoagulants (NOACs). In this observational study we describe CMB profiles in a multicenter cohort of 89 anticoagulation-related intracerebral hemorrhage (ICH) patients. CMB prevalence was 51% (52% in VKA-ICH, 48% in NOAC-ICH). NOAC-ICH patients had lower median CMB count [2(IQR:1–3) vs. 7(4–11); P < 0.001]; ≥5 CMBs were less prevalent in NOAC-ICH (4% vs. 31%, P = 0.006). This inverse association between NOAC exposure and high CMB count persisted in multivariable logistic regression models adjusting for potential confounders (OR 0.10, 95%CI: 0.01–0.83; P = 0.034).

Introduction

Cerebral microbleeds (CMBs) are cerebral small vessel disease markers identified in blood-sensitive brain MRI sequences (gradient-recalled echo [GRE] or susceptibility-weighted imaging [SWI]).¹ They correspond to diverse underlying pathological processes. While hypertensive² or cerebral amyloid angioopathy-related microhemorrhages are the most commonly associated conditions, ischemic processes such hemorrhagic transformation of ischemic microinfarcts and extravasation of red thrombi might also have similar imaging appearance CMBs have been linked
to increased incident risk of both ischemic and hemorrhagic stroke. Although ischemic strokes are more frequent in terms of absolute event rates, the association of CMB presence with incident intracerebral hemorrhage (ICH) for the individual patient is more potent, especially in those with higher CMB burden. Thus, CMBs are recognized as a potentially useful prognostic of ICH especially in high-risk patients exposed to therapeutic oral anticoagulation.

Less is known regarding the relationship between oral anticoagulation and the pathogenesis of CMBs. Data regarding Vitamin K antagonists (VKAs) and CMBs generally suggest higher risk, especially among those with highly variable international normalized ratio (INR). Data regarding nonvitamin K oral antiocoagulants (NOACs) are scarce and suggest no overall association between NOAC exposure and CMB incidence or burden. Notably, a recent multicenter study reported higher CMB prevalence in patients experiencing ICH than ischemic stroke while taking NOACs. We undertook this observational, cross-sectional study to characterize the prevalence and burden of CMBs in patients with symptomatic anticoagulation-associated ICH and examine potential differences in VKA versus NOAC-exposed patients.

Methods

Patient population

This is a retrospective analysis of two prospectively enrolled consecutive patient cohorts of nontraumatic ICH and positive history of oral anticoagulant intake in 15 participating tertiary-care stroke centers during a 2-year period (August 2015–July 2016 and August 2016–July 2017). Patient characteristics and outcome have been described in detail previously and are available in the online supplement (Table S1). Briefly, the definition of VKA-related ICH required effective use of a VKA with an international normalized value of >1.5 on hospital admission, while the definition of NOAC-related ICH required confirmed ingestion of the relevant NOAC during the last 24 h before the index event. Patients with major head trauma or known underlying structural or vascular cause of ICH were excluded from further evaluation. We also excluded patients with hemorrhagic transformation of ischemic infarcts and patients with pure intraventricular hemorrhage. Lastly, we excluded patients without baseline MRI with available blood-sensitive sequences allowing CMB detection.

Imaging characteristics and criteria

We defined CMBs according to the Standards for Reporting Vascular changes on neuroimaging (STRIVE) consensus criteria. We recorded magnet strength (1.5 vs. 3T) and type of sequence used. Further details on CMB reading are available in the online supplement. MRI was performed in the acute phase (during the admission for the index ICH). Images were reviewed by experienced vascular neurologists at each center, blinded to anticoagulation exposure and official Neuroradiologist read for both CMB presence and ICH location. Formal Neuroradiological interpretation was used to corroborate ICH location.

Statistical analyses

Continuous variables are presented as median with interquartile range, whereas categorical variables are presented as percentages. Statistical comparisons between different subgroups were performed using the Pearson’s χ² test and Mann–Whitney U test, where appropriate.

Multivariable logistic regression analyses were performed on the association of baseline characteristics with the presence of five or more cerebral microbleeds in baseline neuroimaging. In univariable models of all baseline characteristics a threshold of P < 0.1 was used to identify candidate variables for inclusion in the multivariate regression models that tested statistical significance hypothesis using the likelihood ratio test with an alpha value of 0.05. We also performed sensitivity univariable/multivariable logistic regression analyses after excluding patients with no presence of cerebral microbleeds on baseline neuroimaging. Finally, we also performed adjusted for baseline ICH volume analyses on the probabilities of hematoma expansion and ICH volume more than 30 cm³ at the follow-up neuroimaging in 24 h according to baseline CMB burden (<5 CMBs or ≥5 CMBs).

We used Stata Statistical Software Release 13 for Windows (College Station, TX, StataCorp LP) for all analyses.

Results

Our two previous cohorts comprised a total of 109 NOAC-related and 248 VKA-related ICHs. Of these, a total of 89 patients (25 NOAC-, 64 VKA-associated) received brain MRI with GRE or SWI sequences allowing CMB detection. Compared to those who did not receive MRI (Table S1), included patients had higher median BMI (30 vs. 27 kg/m², P = 0.002), were less likely to have hypertension (89% vs. 97%; P = 0.002), hyperlipidemia (52% vs. 65%; P = 0.03), and coronary artery disease (28% vs. 40%; P = 0.04); otherwise there were no significant differences. Patient characteristics according to anticoagulant type are summarized in Table 1. NOAC-ICH patients were older (median age 78 [70–81] years vs. 70
[60–77] years; \( P = 0.005 \), had higher CHA2DS2-VASc score (4[3–5] vs. 4[3–5]); \( P = 0.017 \), and were less likely to have lobar ICH location (28% vs. 58%; \( P = 0.001 \)).

A total of 45 patients (51%) had \( \geq 1 \) CMB. There was no difference between VKA- and NOAC- ICH (52% vs. 48%; \( P = 0.763 \); Table 1). However, the median CMB number was significantly lower in NOAC-ICH patients (2 [1–3] vs. 7[4–11]; \( P < 0.001 \) (Table 1 and Fig. 1)). A total of 21 patients had high CMB burden (\( \geq 5 \) CMBs). Their characteristics are summarized in Table S2. High CMB burden (\( \geq 5 \) CMBs) was less prevalent in NOAC-ICH patients (31% vs. 4%; \( P = 0.006 \), Table 1). In contrast, high CMB burden was more common in younger patients and in patients who underwent 3 Tesla MRI (Table 2). This inverse association between NOAC (vs. VKA) exposure and high CMB count persisted in multivariable logistic regression models adjusting for potential confounders including demographics, risk factors, laboratory and brain imaging parameters: (OR 0.10, 95%CI:0.01–0.83; \( P = 0.034 \); Table 2). In sensitivity analyses, four factors were associated with high CMB burden in this sensitivity analysis: age, NOAC pretreatment, antiplatelet pretreatment and MRI strength (Table S3). Pretreatment with NOACs was independently related to lower odds of high CMB burden (OR 0.02, 95%CI: 0.01–0.25; \( P = 0.006 \)) in the sensitivity analysis (Table S3).

### Table 1. Baseline characteristics and outcomes according to the type of oral anticoagulant treatment

| Variable | VKA (n = 64) | NOAC (n = 25) | \( P \)-value |
|----------|-------------|--------------|--------------|
| Baseline clinical characteristics | | | |
| Age (years, median, IQR) | 70 (60–77) | 78 (70–81) | 0.005 |
| Males (%) | 65.6% | 52.0% | 0.234 |
| BMI (median, IQR) | 30 (25–33) | 27 (17–34) | 0.133 |
| CHA2DS2-VASc score (median, IQR) | 4 [3–5] | 4 [4–6] | 0.017 |
| HAS-BLED score (median, IQR) | 3 [2–3] | 2 [2–4] | 0.917 |
| Hypertension (%) | 92.1% | 96.0% | 0.519 |
| Diabetes (%) | 42.2% | 32.0% | 0.376 |
| Hyperlipidemia (%) | 50.0% | 48.0% | 0.865 |
| Heart failure (%) | 21.9% | 12.0% | 0.287 |
| Current smoking (%) | 10.9% | 4.0% | 0.304 |
| Coronary artery disease (%) | 31.2% | 32.0% | 0.945 |
| Chronic kidney disease (%) | 17.2% | 16.0% | 0.893 |
| Prior history of ischemic stroke (%) | 29.7% | 24.0% | 0.592 |
| Baseline Laboratory values | | | |
| INR admission (median, IQR) | 2.4 [1.8–3.6] | 1.2 (1.1–1.6) | <0.001 |
| aPTT admission (sec, median, IQR) | 39 (33–42) | 30 (28–32) | <0.001 |
| Platelet count \( \times 10^9/\mu L \) (median, IQR) | 192 (159–259) | 218 (184–270) | 0.217 |
| CrCl on admission (ml/min, median, IQR) | 60 (44–70) | 60 (45–75) | 0.291 |
| Baseline neuroimaging findings | | | |
| Lobar hemorrhage (%) | 57.8% | 28.0% | 0.001 |
| Intraventricular hemorrhage (%) | 35.9% | 32.0% | 0.726 |
| Baseline ICH volume (cm\(^3\), median, IQR) | 11.3 (5.1–26.3) | 4.9 (2.1–22.1) | 0.051 |
| ICH score (median, IQR) | 1 (1–2) | 1 (1–2) | 0.635 |
| Severe ICH (%)\(^1\) | 14.5% | 16.0% | 0.861 |
| CMB presence (%) | 51.6% | 48.0% | 0.763 |
| CMB number (median, IQR)\(^2\) | 7 (4–11) | 2 (1–3) | <0.001 |
| CMB \( \geq 5 \) (%) | 31.2% | 4.0% | 0.006 |
| CMB \( \geq 10 \) (%) | 17.1% | 4.0% | 0.102 |
| 3T MRI | 39.1% | 36.0% | 0.789 |
| SWI sequence | 6.2% | 0% | 0.201 |

\(^1\)Defined as ICH score \( \geq 2\).

\(^2\)After excluding patients without CMB presence.
Discussion

In this cross-sectional observational study of anticoagulation-associated ICH, we documented an overall prevalence of CMBs of 51%. Although CMB prevalence was well-balanced between VKA and NOAC-ICH, NOAC-ICH patients had significantly lower CMB burden, both in median number and when dichotomized as <5 versus ≥5 CMBs. These associations remained significant after adjusting for potential confounders.

The observed CMB prevalence of ~50% is significantly higher than the frequency reported in population-based studies\textsuperscript{12–14} and studies in ischemic stroke patients\textsuperscript{3} which range from 8.8 to 23%, with tendency to increase with age. However, this prevalence is comparable or lower than CMB frequency reported in ICH cohorts which often

Table 2. Univariable and multivariable logistic regression analyses on the association of baseline characteristics with the presence of five or more cerebral microbleeds in baseline neuroimaging

|                          | Univariable analysis | Multivariable analysis |
|--------------------------|----------------------|------------------------|
|                          | OR (95%CI)           | P                      | OR (95%CI) | P   |
| Age (years)              | 0.94 (0.89, 0.98)    | 0.011                  | 0.94 (0.89, 0.99) | 0.031 |
| Males (%)                | 1.28 (0.48, 3.40)    | 0.616                  | –          | –   |
| BMI                      | 1.03 (0.97, 1.09)    | 0.338                  | –          | –   |
| Hypertension             | 1.59 (0.17, 14.40)   | 0.681                  | –          | –   |
| Diabetes                 | 1.21 (0.45, 3.27)    | 0.705                  | –          | –   |
| Hyperlipidemia           | 1.50 (0.56, 4.02)    | 0.421                  | –          | –   |
| Heart failure            | 0.99 (0.29, 3.46)    | 0.994                  | –          | –   |
| Current smoking          | 2.10 (0.46, 9.64)    | 0.340                  | –          | –   |
| Coronary artery disease  | 0.61 (0.20, 1.88)    | 0.390                  | –          | –   |
| Kidney failure           | 1.81 (0.54, 6.06)    | 0.335                  | –          | –   |
| Prior history of ischemic stroke | 1.39 (0.48, 3.99)   | 0.542                  | –          | –   |
| Prior history of intracerebral hemorrhage | 3.47 (0.46, 26.32) | 0.228                  | –          | –   |
| Statin pretreatment      | 1.32 (0.47, 3.69)    | 0.600                  | –          | –   |
| Antiplatelet pretreatment| 0.48 (0.16, 1.38)    | 0.172                  | –          | –   |
| NOAC pretreatment        | 0.09 (0.01, 0.72)    | 0.024                  | 0.10 (0.01, 0.83) | 0.034 |
| Admission SBP            | 0.99 (0.98, 1.01)    | 0.575                  | –          | –   |
| Admission DBP            | 1.02 (0.99, 1.05)    | 0.191                  | –          | –   |
| Lobar hemorrhage         | 1.17 (0.44, 3.11)    | 0.758                  | –          | –   |
| 3T MRI                   | 4.80 (1.68, 13.67)   | 0.003                  | 6.42 (1.96, 21.03) | 0.002 |
| SWI sequence             | 3.47 (0.46, 26.32)   | 0.228                  | –          | –   |
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External validation in a larger cohort is necessary. Given the cross-sectional nature of this analysis, it is not possible to ascertain whether there is a causative link between anticoagulation exposure and CMB formation or whether CMB presence and anticoagulation exposure act synergistically to increase the risk of hemorrhage. Approximately 25% of the entire cohort received MRI and was available for analysis. Despite small imbalances in baseline cardiovascular risk factor prevalence, we found no evidence of a systematic selection bias with regard to parameters of interest, including anticoagulation type, concomitant medications, ICH severity and overall CHA2DS2-VASc and HAS-BLED score (Table S1). MRI protocols and strengths which are well-known to affect the sensitivity of CMB detection were heterogeneous among participating centers. However, both 3T MRI and SWI sequence use were evenly distributed between anticoagulation allocation groups (Table 1) and the association between NOAC-ICH and lower CMB burden remained significant after adjusting for magnet strength. We had no information regarding duration of exposure to anticoagulation which might be an important factor determining development of CMBs. However, we note that NOAC-ICH patients were older and with higher CHA2DS2-VASc score; it is plausible that they might have been exposed to VKA and subsequently changed to NOAC, which would further highlight lower risk of CMB formation associated with NOAC. We do not have information regarding CMB anatomical location, which is a shortcoming subtracting granularity from our findings, given that different CMB location implies differential underlying pathology.13,14

In conclusion, we documented similar prevalence but significantly higher burden of CMBs in VKA compared to NOAC exposure, in a cohort of 89 patients with anticoagulation-related ICH. Results of additional ongoing prospective studies (Intracerebral Hemorrhage Due to Oral Anticoagulants: Prediction of the Risk by Magnetic Resonance (HERO) https://clinicaltrials.gov/ct2/show/NCT02238470) and Cerebral Microbleeds During NOACs or Warfarin Therapy in NVAF Patients With Acute Ischemic Stroke (CMB-NOW) https://clinicaltrials.gov/ct2/show/NCT02356432) are expected to further refine our understanding of the interaction between CMBs and anticoagulant therapy.

Author Contributions

Vasileios Lioutas participated in conception and design of study and first manuscript draft. Aristeidis Katsonas involved in drafting of manuscript and figures/tables. Georgios Tsivgoulis involved in study conception and design, manuscript, and figures drafting. All authors involved in acquisition and analysis of the data.
Conflict of Interest

GT reports advisory board and speaker honoraria from Boehringer Ingelheim, Bayer, Daichii Sankyo, Medtronic, Shire, CSL Behring, Allergan, and Biogen; and an unrestricted research grant from Medtronic.

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Supporting Information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Table S1. Univariable comparisons of baseline characteristics between patients who received MRI vs those who did not.

Table S2. Baseline characteristics and outcomes according to high cerebral microbleed burden (≥5) on baseline neuroimaging.

Table S3. Univariable and multivariable logistic regression analyses on the association of baseline characteristics with the presence of five or more cerebral microbleeds on baseline neuroimaging, after excluding patients with no cerebral microbleeds on baseline neuroimaging.