Association of stroke lesion shape with newly detected atrial fibrillation – Results from the MonDAFIS study

Bernardo Crespo Pimentel1,2*, Thies Ingwersen1*, Karl Georg Haeusler3,4, Eckhard Schlemm1, Nils D Forkert5, Deepthi Rajashekar5, Pauline Mouches5, Alina Königsberg1, Paulus Kirchhof1,4,6,7,8, Claudia Kunze9, Serdar Tütüncü9, Manuel C Olma9, Michael Krämer9, Dominik Michalski10, Andrea Kraft11, Timolaos Rizos12, Torsten Helberg13, Sven Ehrlich14, Darius G Nabavi15, Joachim Röther16, Ulrich Laufs17, Roland Veltkamp18,19, Peter U Heuschmann20,21, Bastian Cheng1, Matthias Endres8,22,23,24,25,26 and Götz Thomalla1

Abstract
Paroxysmal Atrial fibrillation (AF) is often clinically silent and may be missed by the usual diagnostic workup after ischemic stroke. We aimed to determine whether shape characteristics of ischemic stroke lesions can be used to predict AF in stroke patients without known AF at baseline. Lesion shape quantification on brain MRI was performed in selected patients from the intervention arm of the Impact of standardized MONitoring for Detection of Atrial Fibrillation in Ischemic Stroke (MonDAFIS) study, which included patients with ischemic stroke or TIA without prior AF. Multiple

1Department of Neurology, Medical Center Hamburg-Eppendorf, Hamburg, Germany
2Department of Neurology, Christian Doppler Medical Center, Paracelsus Medical University, Salzburg, Austria
3Department of Neurology, Universitätssklinikum Würzburg, Würzburg, Germany
4German Atrial Fibrillation Network (AFNET), Münster, Germany
5Department of Radiology, University of Calgary, Calgary, AB, Canada
6Department of Cardiovascular Sciences, College of Medical and Dental Sciences, Medical School, University of Birmingham, UK
7Departments of Cardiology, UHB and SWBH NHS Trusts, Birmingham, UK
8University Heart and Vascular Center Hamburg, Hamburg, Germany
9Center for Stroke Research Berlin, Charité-Universitätsmedizin Berlin, Berlin, Germany
10Department of Neurology, Universitätssklinikum Leipzig, Leipzig, Germany
11Department of Neurology, Maritha Maria Hospital, Halle Dölau, Germany
12Department of Neurology, University of Heidelberg, Heidelberg, Germany
13Department of Neurology, Clinical Center of Hubertusburg, Wermsdorf, Germany
14Clinical Center of Hubertusburg, Wermsdorf, Germany
15Department of Neurology, Vivantes Klinikum Neukölln, Berlin, Germany
16Department of Neurology, Asklepion Klinik Altona, Hamburg, Germany
17Department of Cardiology, Universitätssklinikum Leipzig, Leipzig, Germany
18Department of Neurology, Alfried Krupp Krankenhaus, Essen, Germany
19Department of Brain Sciences, Imperial College London, UK
20Comprehensive Heart Failure Center & Clinical Trial Centre Würzburg, University Hospital Würzburg, Germany
21Institute of Clinical Epidemiology and Biometry, University Würzburg, Würzburg, Germany
22Klinik und Hochschulambulanz für Neurologie mit Abteilung für Experimentelle Neurologie, Charité-Universitätsmedizin Berlin, Berlin, Germany
23German Center for Neurodegenerative Diseases, Partner Site Berlin, Germany
24German Center for Cardiovascular Diseases, Partner Site Berlin, Germany
25ExcellenceCluster NeuroCure, Berlin, Germany
26Berlin Institute of Health (BiH), Berlin, Germany

*Bernardo Crespo Pimentel & Thies Ingwersen contributed equally to this work.

Corresponding author:
Götz Thomalla, Department of Neurology, University Hospital Hamburg-Eppendorf, Martinistrasse 52, Hamburg 20246, Germany.
Email: thomalla@uke.de
morphologic parameters were calculated based on lesion segmentation in acute brain MRI data. Multivariate logistic models were used to test the association of lesion morphology, clinical parameters, and AF. A stepwise elimination regression was conducted to identify the most important variables. A total of 755 patients were included. Patients with AF detected within 2 years after stroke ($n = 86$) had a larger overall oriented bounding box (OBB) volume ($p = 0.003$) and a higher number of brain lesion components ($p = 0.008$) than patients without AF. In the multivariate model, OBB volume (OR 1.72, 95%CI 1.29–2.35, $p < 0.001$), age (OR 2.13, 95%CI 1.52–3.06, $p < 0.001$), and female sex (OR 2.45, 95%CI 1.41–4.31, $p = 0.002$) were independently associated with detected AF. Ischemic lesions in patients with detected AF after stroke presented with a more dispersed infarct pattern and a higher number of lesion components. Together with clinical characteristics, these lesion shape characteristics may help in guiding prolonged cardiac monitoring after stroke.

Keywords
Ischemic stroke, cardioembolism, atrial fibrillation, lesion shape

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Introduction
Cardioembolism accounts for approximately 20% of all ischemic strokes$^{1,2}$ and is mostly caused by atrial fibrillation (AF). Identification of AF following acute ischemic stroke (AIS) or transient ischemic attack (TIA) is crucial as oral anticoagulation is more effective for secondary prevention in this subgroup of patients than antiplatelet agents.$^3$ As AF is often clinically silent, it may go undetected during the in-hospital diagnostic work-up, thus depriving a significant proportion of patients of the most efficient preventive treatment.$^4$

Different approaches can be used to detect non-permanent AF after stroke. Repeated Holter-ECG is frequently suggested but is not universally available. Invasive cardiac monitoring (ICM) is an ancillary procedure that increases the detection rate of AF$^5$ and is suggested for patients without detected AF but high suspicion of a cardioembolic source of stroke.$^5$ However, there are no established criteria to guide ICM implantation in stroke patients, as there is no randomized controlled trial demonstrating a subsequent reduction of stroke recurrence or death.

Diffusion-weighted imaging (DWI) and fluid-attenuated inversion recovery (FLAIR) have shown high sensitivity in detecting even small ischemic lesions and allow the assessment of complex lesion characteristics such as shape and distribution patterns.$^7,9$ The analysis of stroke morphologic features, such as size, lesion dispersion, and sphericity, has been explored in several clinical studies.$^7,8,10,11$ Limited evidence from MRI-based studies suggests a potential role of location patterns, such as scattered stroke lesions, cortical involvement, or involvement of multiple arterial territories in anticipating a cardio-embolic source of stroke.$^{12–17}$

Thus, it remains unknown if specific morphologic lesion patterns can be used to identify patients with a high risk of AF. The aim of this pre-defined sub-study of the Impact of standardized MONitoring for Detection of Atrial Fibrillation in Ischemic Stroke (MonDAFIS) study$^{18,19}$ was to characterize ischemic stroke lesion shape and location patterns and use these markers to predict a first episode of AF.

Material and methods
Study design
In this exploratory, predefined analysis, we included data of patients randomized in the intervention arm of the MonDAFIS study who underwent brain MRI during in-hospital diagnostic work-up. MonDAFIS was an investigator-initiated, randomized, multi-center study conducted at 38 certified stroke units in Germany that assessed the effect of prolonged inpatient Holter ECG monitoring on anticoagulation rates 12 months after AIS/TIA.

Patients with AIS or TIA without known AF before hospital admission were included and randomized 1:1 to standard diagnostic procedures or standard diagnostic procedures plus in-hospital Holter ECG monitoring for up to 7 days. Inclusion criteria included age $\geq 18$ years and stroke unit admission within 72 h after onset of stroke-related symptoms. Patients with pre-stroke life expectancy less than a year and post-stroke life expectancy less than a month, participation in an interventional study, pregnancy or breastfeeding, independent indication for long-term oral anticoagulation as well as patients with implanted devices with the ability to record an ECG were excluded.$^{18}$

For the current analysis, we further excluded subjects without acute ischemic lesions on DWI, as well as patients with MRI brain imaging acquired later than 7 days following the index stroke or insufficient image quality. In the MonDAFIS study, presence of AF in hospital was defined in accordance with the current European Society of Cardiology guidelines as an absolute arrhythmia lasting $\geq 30$ s.$^5$ During the follow-up period of 2 years, detection of AF was assessed through a telephone interview conducted by the study team at the Center for Stroke Research Berlin.

For descriptive and analysis purposes, we divided patients into two groups. In the first group, we included
patients who had AF detected in-hospital after the index stroke or TIA as well as patients self-reported to have documented AF within 24 months after the index stroke or TIA. All other study patients were assigned to the second group. All study procedures were carried out in accordance with the Declaration of Helsinki. The study was approved by the ethics committees of Charité – Universitätsmedizin Berlin, Germany, and all participating sites. All patients provided written informed consent.

Lesion segmentation

Subject datasets were analyzed using a segmentation algorithm (SONIA) based on the in-house developed software tool ANTONIA.20 SONIA performs DWI image co-registration to FLAIR images, semi-automated stroke lesion segmentation based on an apparent diffusion coefficient (ADC) threshold, and registration to standard MNI (Montreal Institute of Neurology) space. Ischemic stroke lesions were semi-automatically segmented using an ADC maximum threshold, and registration to standard MNI (Montreal Institute of Neurology) space. Ischemic stroke lesions were semi-automatically segmented using an ADC maximum threshold of 620 × 10^{-6} \text{mm}^2/\text{s}. Segmented voxels were then manually corrected by two trained neurology residents (BCP, TI). Each individual lesion mask was registered to the corresponding FLAIR image, being consequently transformed to MNI standard space using an affine registration process. Registration results were visually evaluated and compared with the original lesion in native space as a quality assurance step. Native space masks were further post-processed with removal of implausible lesion components smaller than 10 \text{mm}^3. MRI data was acquired on 1.5 or 3 T scanners in 20 participating centers. Pixel spacing ranged from 0.71 to 1.88 mm with a slice thickness of 3–6 mm.

Lesion shape parameters

Morphological descriptors were calculated in lesion masks in native space using customized in-house developed tools. The following brain lesion shape parameters were calculated: (1) number of lesion components; (2) volume of the stroke lesion (mL); (3) surface area of the ischemic lesion (mm2); (4) the oriented minimum bounding-box of the lesion (mL, OBB), that is, the smallest rectangular volume comprising all lesion components (whereas the orientation is independent of the coordinate axes); (5) the ratio between lesion volume and OBB volume, that is, the proportion of the OBB filled by the actual stroke lesion; and (6) the normalized shape factor S of the ischemic lesion, which describes how closely the lesion resembles the shape of a sphere, defined as:

\[ S = \frac{\sqrt[3]{A}}{2.199085233} \]

where \( A \) is the surface area and \( V \) the volume of the lesion.

Lesion location characterization

Inferences about lesion location were determined using the mindboggle-101 brain atlas21 and a previously published vascular territory map.22 Location parameters included the affected hemisphere (left, right, both), vascular territory (left/right internal carotid artery, basilar artery, multiple affected territories), and any cortical region affected. A minimum of 100 lesion voxels overlapping the respective location was chosen as a threshold.

Statistical analysis

Descriptive statistics are reported in absolute counts and percentages or medians and interquartile range, as applicable. Group differences in frequencies were analyzed using the chi-squared test and \( p \)-values were calculated by Monte Carlo simulation. Groups were compared using the Mann–Whitney test. To determine a potential relation between lesion shape parameters and AF, logistic regression models were fitted in the following way: first, logarithmized parameters were analyzed individually in univariate models. In order to compare the individual impact of different variables, all values were scaled to standardized values (\( z \)-scores) beforehand. Second, a multivariate model was fitted containing the parameters “number of components,” “lesion volume,” “sphericity,” “OBB volume,” and “ratio between lesion and OBB volume.” Due to collinearity with “lesion volume,” “surface area,” was excluded from the model. This regression model was further refined by a simple stepwise elimination algorithm. During every step, variables that did not reach the significance level of 0.05 were eliminated from the model starting with the least significant variable.

In a second analysis, the multivariate models were complemented by clinical variables (age, sex, NIHSS score at admission) and imaging location parameters (multiple vascular territories affected, cortical involvement). All tests were performed two-sided and statistical significance was set to an alpha of 0.05. Statistical analysis was performed using R Studio (R software package, v. 3.6.2; R Foundation for 159 Statistical Computing, Vienna, Austria).

Results

We analyzed brain MRI data from 1,054 patients of the MonDAFIS intervention arm (see Figure 1). We excluded patients with missing clinical information (\( n = 146 \)), brain MRI performed >7 days following the index stroke (\( n = 30 \)) and patients without an acute ischemic brain lesion in DWI (\( n = 59 \)). From the remaining subjects, we excluded datasets with insufficient image quality or missing MRI sequences (\( n = 64 \)). Thus, we performed semi-automated segmentation of acute ischemic brain lesions in 755 patients. Information on AF at 24 months after index stroke was missing in 54
patients, which were also excluded. Individual space lesion masks were used for the lesion shape analysis \((n = 701)\) and lesion location analysis was carried out in standard MNI space. Due to unsuccessful image normalization to MNI space, 57 datasets were excluded, resulting in 644 subjects available for lesion location analysis.

Summary descriptive statistics are displayed in Table 1. Median age was 66 years (IQR 57–76) and 261 patients (37.5%) were female in the shape analysis group \((n = 701)\). MRI was performed in a median of 1 day (IQR 1–3 days) after hospital admission. AF was detected in a total of 86 patients (12.3%) within the 24 months follow-up period. Compared to patients without AF, female sex (52% vs 35%, \(p = 0.003\)) and advanced age (74 years (IQR 67–78) vs 65 years (IQR 56–74), \(p < 0.001\)) was more often observed in patients with detected AF. No difference regarding the NIHSS score at hospital admission was observed. In the AF group, lesions were more often located in cortical regions (44% vs 28%, \(p = 0.008\)) (Table 2). Figure 2 shows overlap lesion maps for patients with and without detected AF. The same analysis was done for the cases of in-hospital diagnosed AF (comprising cases detected through monitoring on the stroke unit and normal neurological ward). Due to the low case number of subjects \((n = 25)\), these results are displayed as supplemental material (see Supplemental Table 1).

Comprehensive lesion shape analysis was carried out in native lesion space. Infarcts in patients with AF had a significantly higher number of separate lesion components (3.0 (IQR 1.0–7.0) vs 2.0 (IQR 1.0–4.0), \(p = 0.008\)). Additionally, the OBB volume was significantly larger (58 mL (IQR 6.1–180.4) vs 10 mL (IQR 3.2–78.2), \(p = 0.003\)) in the AF group (see Figure 4).

In univariate generalized logistic regression models, AF detection within 24 months after the index stroke/TIA was associated with a larger OBB volume (OR 1.42, 95%CI 1.09–1.87, \(p = 0.010\)) and higher number of lesion components (OR 1.33, 95%CI 1.06–1.70, \(p = 0.013\)), while no association with lesion volume, surface area, and sphericity was found. In the multivariate analysis including lesion shape characteristics, clinical variables (age, sex, NIHSS at admission), as well as cortical involvement and involvement of multiple vascular territories, only female sex and old age were significantly associated with AF detection (see Table 3). The stepwise elimination algorithm identified age (OR 2.13, 95%CI 1.52–3.06, \(p < 0.001\)), female sex (OR 2.45, 95%CI 1.41–4.31, \(p = 0.002\)), and OBB volume (OR 1.72, 95%CI 1.29–2.35, \(p < 0.001\)) as statistically associated with AF (see Table 3). Figure 3 displays an example of two acute stroke lesions with different OBB volumes.

**Discussion**

In this pre-defined analysis of the prospective MonDAFIS study, patients with AF detected within 24 months after AIS showed a higher number of ischemic lesions and a wider

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**Table 1.** Group difference in clinical and shape parameters between patients with and without detected atrial fibrillation (AF) within 24 months after ischemic stroke or TIA.

| Clinical variables              | Total \((n = 701)\) | No AF \((n = 615)\) | Detected AF \((n = 86)\) | \(p\)  |
|--------------------------------|---------------------|---------------------|--------------------------|-------|
| Female sex, \(n\) (%)          | 261 (37)            | 216 (35)            | 45 (52)                  | 0.003 |
| Age, years (IQR)               | 66 (57–76)          | 65 (56–74)          | 74 (67–78)               | <0.001|
| Admission NIHSS score, points (IQR) | 2 (1–4)             | 2 (1–4)             | 2.5 (1–4)                | 0.532 |
| Shape parameters               |                     |                     |                          |       |
| Lesion components, \(n\) (IQR) | 2 (1–4)             | 2 (1–4)             | 3 (1–7)                  | 0.008 |
| Lesion volume, mL (IQR)        | 0.39 (0.13–1.13)    | 0.39 (0.14–1.00)    | 0.45 (0.11–3.03)         | 0.336 |
| Surface area, mm\(^2\) (IQR)   | 39.35 (17.77–79.49) | 38.91 (17.86–74.15) | 40.34 (16.99–141.5)      | 0.336 |
| Sphericity, value (IQR)        | 1.26 (1.21–1.32)    | 1.26 (1.21–1.318)   | 1.28 (1.22–1.33)         | 0.319 |
| OBB Volume, mL (IQR)           | 13 (3.37–95.8)      | 9.87 (3.15–78.3)    | 58.14 (6.10–180.4)       | 0.003 |
| Lesion/OBB Volume, ratio (IQR)| 0.05 (0.01–0.13)    | 0.05 (0.01–0.14)    | 0.02 (0.00–0.13)         | 0.108 |
Table 2. Group difference in location parameters between patients with and without detected atrial fibrillation (AF) within 24 months after ischemic stroke or TIA.

| Location parameters                  | Total (n = 644) | No AF (n = 565) | Detected AF (n = 79) | p    |
|--------------------------------------|-----------------|-----------------|----------------------|------|
| Cortical involvement, n (%)          | 192 (30)        | 158 (28)        | 34 (44)              | 0.008|
| Multiple territories, n (%)          | 42 (6)          | 35 (6)          | 7 (9)                | 0.491|
| Bilateral involvement, n (%)         | 50 (8)          | 41 (7)          | 9 (12)               | 0.249|

Figure 2. Overlay map of lesion density in patients without detected atrial fibrillation (AF) and with newly detected AF. Lesion density is displayed in the color bar on the right side.

Figure 3. Representative example of two differing right-sided stroke lesions. The large lesion consists of eight components and measures 9.8 mL in total. The resulting overall bounding box (OBB) is 1204 mL in volume (A and B). In contrast, the OBB of a smaller lesion of 0.36 mL consisting of one component is only 7.2 mL in volume (C and D). Three-dimensional reconstructions are depicted from a fronto-lateral (left column) and dorsolateral (right column) view.
Older age, female sex, and spatial lesion distribution were independently associated with AF in the analyzed subcohort of study patients. The clinical benefit of imaging parameters should be evaluated in further studies.

In our cohort, stroke patients with detected AF displayed a higher number of lesion components as compared to patients without AF during follow-up. This corroborates results from previous observational studies using MRI-DWI or CT imaging.\textsuperscript{11,14,15,23} In the context of AF, repetitive embolism or large emboli generated in the left atrium that fragment on their way and end up at multiple distinct sites within the cerebral vascular bed likely leads to a multi-site infarct pattern.

Additionally, detection of AF was associated with a significantly larger minimum OBB volume, consistent with a manifold infarct pattern with extensive spatial distribution. In contrast to distal arterial sources of embolic material, cardiac emboli can simultaneously travel through any of the major arteries supplying the brain and may induce a multiple vascular pattern. Indeed, multiple lesion patterns affecting various cerebral vascular territories are associated with a cardio-embolic stroke, a proximal embolic source or the presence of tumors (e.g., in the thoracic aorta).\textsuperscript{9,11,24,25}

Moreover, lesions in patients with AF detected during follow-up were of similar size as in patients in whom no AF was detected. This finding contradicts some previous studies demonstrating that ischemic lesions related to AF tend to be larger compared to other etiologies,\textsuperscript{24,26} probably owing to the larger size of cardiac emboli as compared to other sources. The possibility that AF was detected as an incidental finding not etiologically related to the index stroke could have also contributed to the volume similarities.

Cortical involvement in strokes related to AF and other cardiac sources is a matter of conflicting evidence. Supporting the findings of previous studies,\textsuperscript{27–29} cortical involvement was more frequent in patients with detected

**Table 3.** Multivariate logistic regression model containing lesion shape parameters and clinical variables and final stepwise elimination model.

| OR (95% CI) | p     |
|------------|-------|
| Multivariate logistic regression |       |
| Age        | 2.15 (1.53–3.12) | <0.001 |
| Female Sex | 2.42 (1.38–4.28)  | 0.002  |
| NIHSS score on admission | 1.15 (0.88–1.49) | 0.297  |
| Cortical involvement | 1.41 (0.65–3.05) | 0.383  |
| Number of components | 1.19 (0.66–2.16) | 0.554  |
| Multiple territories | 0.71 (0.21–2.08) | 0.556  |
| Lesion volume | 1.16 (0.71–1.90) | 0.562  |
| OBB volume   | 1.24 (0.58–2.60) | 0.579  |
| Sphericity   | 1.03 (0.73–1.44) | 0.855  |
| Stepwise elimination model |       |
| Age        | 2.13 (1.53–3.06) | <0.001 |
| Female sex | 2.45 (1.41–4.31)  | 0.002  |
| OBB volume | 1.72 (1.29–2.35)  | <0.001 |

Spatial lesion distribution in the acute imaging work-up. Older age, female sex, and spatial lesion distribution were independently associated with AF in the analyzed subcohort of study patients. The clinical benefit of imaging parameters should be evaluated in further studies.

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AF in our cohort. Nevertheless, it was not independently associated with AF in the multivariate regression analysis.

Apart from lesion OBB volume, old age and female sex were identified as significant factors associated with AF in the multivariate regression analysis. This high predictive value in detection of AF is in line with previous findings.\textsuperscript{30,31}

To our knowledge, the HAVOC and ASSF scores\textsuperscript{32,33} are the only clinical scores developed to stratify patients with cryptogenic stroke/TIA into risk categories of AF detection. Both scores are however solely based on clinical and/or ECG data. With this study, we contribute to a growing body of evidence supporting imaging-based scores to identify stroke patients at high-risk of a first episode of AF.

Our study has several limitations. First, the MonDAFIS study was not powered to identify significant predictors for AF in subgroups of patients and a relevant number of designated patients for this subgroup analysis had to be excluded. Despite the relatively large patient cohort, the pragmatic inclusion criteria of the MonDAFIS study constitute a limiting factor, as a substantial number of patients was aged <65 years, and thus, at comparably low risk of AF. Of note, there was no systematic ECG monitoring after hospital discharge. The use of a stepwise regression model may also, despite the small number of variables, favor nuisance variables and therefore affect the robustness of our data.

Conclusion

Ischemic stroke in patients with detected AF during a 24 months follow-up tend to present with a widespread multiple infarct pattern and a higher number of brain lesions. In addition to old age and female sex, spatial lesion dispersion was independently associated with AF detection. Our results therefore support visual rating of ischemic lesion patterns for AF risk stratification. Whether prolonged ECG monitoring in these patients improves stroke prevention has to further investigated.

Declarations of conflict of interest

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Informed consent

All patients provided written informed consent.

Ethical approval

All study procedures were carried out in accordance with the Declaration of Helsinki. The study was approved by the ethics committees of Charité – Universitätsmedizin Berlin, Germany, and all participating sites.

Guarantor

Prof. Götz Thomalla, MD

Contributorship

GT, KGH, ME, BCP, BC and TI researched literature and conceived the study. ME, KGH and GT were involved in protocol development of the PreDAFIS substudy, gaining ethical approval. KGH, CK, ST, MO, DM, AK, TR, TH, SE, DGN, JR, ME and GT were involved in patient recruitment. BCP, TI, DR, PM, NF, AK, GT conducted the data analysis. BCP and TI wrote the first draft of the manuscript. All authors reviewed and edited the manuscript and approved the final version of the manuscript.

ORCID iD

Bernardo Crespo Pimentel \url{https://orcid.org/0000-0002-8965-8081}

Supplemental material

Supplemental material for this article is available online.
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