Endothelial barrier antigen-immunoreactivity is conversely associated with blood-brain barrier dysfunction after embolic stroke in rats

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Abstract

While the concept of the Neurovascular Unit (NVU) is increasingly considered for exploring mechanisms of tissue damage in ischemic stroke, immunohistochemical analyses are of interest to specifically visualize constituents like the endothelium. Changes in immunoreactivity have also been discussed to reflect functional aspects, e.g., the integrity of the blood-brain barrier (BBB). This study aimed to characterize the endothelial barrier antigen (EBA) as addressed by the antibody SMI-71 in a rat model of embolic stroke, considering FITC-albumin as BBB leakage marker and serum levels of BBB-associated matrix metalloproteinases (MMPs) to explore its functional significance. Five and 25 h after ischemia onset, regions with decreased BBB integrity exhibited a reduction in number and area of EBA-immunopositive vessels, while the stained area per vessel was not affected. Surprisingly, EBA content of remaining vessels tended to be increased in areas of BBB dysfunction. Analyses addressing this interrelation resulted in a significant and inverse correlation between the vessels’ EBA content and degree of BBB permeability. In conclusion, these data provide evidence for a functional relationship between EBA-immunoreactivity and BBB dysfunction in experimental ischemic stroke. Further studies are required to explore the underlying mechanisms of altered EBA-immunoreactivity, which might help to identify novel neuroprotective strategies.

Introduction

As consequence of numerous neuroprotective failures in acute ischemic stroke,1 the perspective of tissue damage shifted from the initial neurocentric to an extended view considering the vasculature as an essential component of the so-called Neurovascular Unit (NVU).2-4 The NVU including endothelial and associated cells like neurons and glia also forms the blood-brain barrier (BBB),5,6 which critically impacts on local homeostasis and ischemia-related complications like edema generation and hemorrhagic transformation.7 However, to gain further insight into the functional role of NVU constituents, their reliable visualization under experimental conditions is required and largely based on specific markers allowing the simultaneous detection of multiple vascular compartments. In this context, antibodies directed against the endothelial barrier antigen (EBA) have frequently been applied to address the cerebral endothelium in rodents also under pathological conditions like ischemic stroke,9-12 intracerebral hemorrhage,13 experimental allergic encephalomyelitis,14 traumatic brain injury,15 and spinal cord trauma.16 Initially described in 1987 by Sternberger and Sternberger,17 anti-EBA is a monoclonal mouse antibody of the immunoglobulin M (IgM) class, that binds to the luminal side of cerebral endothelial cells. The name EBA was chosen due to the assumption that antigen reactivity is associated with parts of the cerebral vasculature exhibiting a selective permeability barrier,6 since EBA-immunoreactivity in the rat brain was found to be absent or diminished in regions with naturally restricted BBB integrity like the area postrema and the choroid plexus.17 Subsequently, EBA was also detected in endothelial cells in the reproductive tract of male rats, probably as a part of the blood-testis barrier.18 However, the distribution of EBA varies strongly in the CNS with a mixture of EBA-immunopositive and -immunonegative cells in pial vessels,19 a complete lack of EBA in cerebral arterioles and a mosaic pattern in capillaries and venules.20 Notably, previous studies demonstrated that cerebral areas with reduced EBA-positive vessels provide increased BBB-permeability for endogenous albumin and IgG19,21,22 leading to the hypothesis that EBA-immunoreactivity might represent a potential tool to explore functional characteristics of the NVU with a special focus on BBB integrity. However, this perspective has so far not been investigated under ischemic conditions, although such a feature might notably complement further research while covering ischemia-related consequences in more detail.

The present study therefore aimed to characterize EBA in a descriptive and functional fashion using an embolic model of focal cerebral ischemia in rats. For this purpose, EBA-immunoreactivity was analyzed to identify the spatio-temporal pattern in ischemia-affected areas exhibiting increased BBB permeability as identified by leakage of intravenously applied fluorescein isothiocyanate (FITC)-albumin. Moreover, relationships between different aspects of EBA-immunoreactivity and BBB permeability as well as matrix metalloproteinases (MMPs), known to promote disintegration of the BBB during stroke,23 were analyzed by correlation coefficients to further explore the functional significance of EBA under ischemic conditions.
Materials and Methods

Study design and tissue preparation

Seventeen male Wistar rats (mean body weight 324 g), provided by Charles River (Sulzfeld, Germany), underwent embolic middle cerebral artery occlusion as described below and were consecutively assigned to an observation period of 5 or 25 h, respectively. Premature death occurred in one animal, resulting in a total of 16 animals (8 in each group) that finished the study addressing spatio-temporal characteristics of EBA-immunoreactivity. In control experiments, 4 rats (mean body weight 358 g) underwent sham-operation, while additional 2 rats suffering from embolic middle cerebral artery occlusion (mean body weight 322 g) were used for triple fluorescence labeling of vascular constituents. For visualizing ischemia-associated changes in BBB integrity, 4 or 24 h after ischemia onset FITC-albumin (molecular weight about 70 kDa; 20 mg dissolved in 1 mL saline; Sigma, Taufkirchen, Germany), usually not crossing the BBB under physiological conditions,21 was applied intravenously via a femoral catheter. After an additional 1-hour circulation period animals were sacrificed and blood was sampled for measuring serum levels of MMPs, followed by transcardial perfusion with a fixative containing 4% paraformaldehyde in phosphate-buffered saline (PBS). Next, brains were removed from the skulls and immersed in the same fixative for 24 hours before their equilibration in 30% phosphate-buffered sucrose. Free-floating 30 µm thick coronal sections from the entire forebrain, stored in 0.1 M Tris-buffered saline, pH 7.4 (Sigma, Taufkirchen, Germany), usually not crossing the BBB under physiological conditions,21 was applied intravenously via a femoral catheter. After an additional 1-hour circulation period animals were sacrificed and blood was sampled for measuring serum levels of MMPs, followed by transcardial perfusion with a fixative containing 4% paraformaldehyde in phosphate-buffered saline (PBS). Next, brains were removed from the skulls and immersed in the same fixative for 24 hours before their equilibration in 30% phosphate-buffered sucrose. Free-floating 30 µm thick coronal sections from the entire forebrain, stored in 0.1 M Tris-buffered saline, pH 7.4 (PBS), containing sodium azide, served for subsequent histological analyses. The experimental protocol involving animals was approved by local authorities (Regierungsspräsidium Leipzig, reference numbers TVV-02/09 and TVV-34/11). Care and treatment of animals conformed to the standards of the European Communities Council Directive (86/609/EEC).

Experimental induction of focal cerebral ischemia and sham procedure

Focal cerebral ischemia was induced by an embolic model originally described by Zhang et al.,22 with some minor modifications as previously reported.24 Briefly, animals were anesthetized breathing 2.0-2.5% isoflurane in a mixture of 70% N2O and 30% O2. During the surgical procedure the body temperature was adjusted to 37°C using a warming pad with a rectal probe (Fine Science Tools, Heidelberg, Germany). After preparation of left femoral vessels and catheter insertion, a tissue-conserving preparation of right cervical vessels was performed. Afterwards, a pointed PE-50 catheter was inserted into the right external carotid artery and advanced into the distal internal carotid artery where a blood clot with a length of approximately 45 mm was released. Finally, the catheter was removed and the stump of the external carotid artery ligated. The sham-procedure was performed in analogous manner, except of catheter insertion into cervical vessels and thus without clot release.

Immunofluorescence labeling of EBA and enhancement of FITC-albumin

A first series of brain sections was washed 3 times for 10 min with TBS before being rinsed with distilled water, mounted on fluorescence-free glass slides, air-dried and coverslipped with Entellan in toluene (Merck, Darmstadt, Germany). Afterwards, sections were screened using a fluorescence microscope (Axioplan; Zeiss, Germany) to identify regions with increased BBB permeability indicated by green fluorescence of extravasated FITC-albumin. Preceding brain sections were then applied to indirect red fluorescent Cy (carbocyanine) 3 staining of EBA-immunoreactivity and concomitant enhancement of FITC-albumin fluorescence. In detail, brain sections were washed 3 times for 10 min each with TBS before being blocked for 1 hour with TBS containing 5% normal donkey serum and 0.3% Triton X-100 (TBS-NDS-T) followed by an overnight incubation with SMI-71, an IgM mouse monoclonal antibody directed against EBA (1-400 in TBS-NDS-T; Covance, Emeryville, CA, USA). Further additional washings with TBS for 10 min each, a cocktail containing Cy2-anti-FITC-IgG (20 µg/mL TBS-RSA; Dianova, Hamburg, Germany), for enhancement and long-term stabilization of the green fluorescent signal of FITC-albumin, and Cy3-donkey-anti-mouse-IgM (20 µg/mL TBS-RSA; Dianova) visualizing SMI-71.

In additional experiments, selected brain sections from animals that underwent middle cerebral artery occlusion and injection of FITC-albumin were blocked with TBS-NDS-T for 1 h and then incubated overnight with a mixture of two endothelial markers, biotinylated Solanum tuberosum agglutinin (20 µg/mL TBS-NDS-T; Vector, Burlingame, CA, USA) and SMI-71 (1:200). Next, the sections were rinsed with TBS and processed with a cocktail composed of Cy5-streptavidin (Dianova), Cy2-anti-FITC-IgG and Cy3-donkey-anti-mouse-IgM (all at 20 µg/mL TBS-RSA) for 1 h. Finally, brain sections were washed, mounted, air-dried and coverslipped as described above.

Quantification of EBA-immunoreactivity

Fluorescence-based measurements were done by a blinded investigator using an Axioplan 2 microscope (Zeiss) with a structured-light confocal system (OptiGrid; Qioptiq, Fairport, NY, USA) and a digital camera (ORCA-ER; Hamamatsu, Hamamatsu City, Japan). Applying a 10x magnification, brain sections were first scanned for regions with clear extravascular green fluorescence signal in terms of FITC-albumin leakage arising from the inner part of the vessels indicating increased BBB permeability. Within those regions, 2 microscopic fields (each covering an area of about 0.55 mm2) were digitized and the total number as well as the stained area of EBA-immunopositive vessels were assessed automatically using the imaging software Velocity 4.3.2 (PerkinElmer, Waltham, MA, USA). Criteria to identify a vessel were a length of at least 12 µm and an area of at least 100 µm2. Thereby, the following settings were applied: Find objects by intensity and exclude objects by size (<100 µm²), while the overall technical standard was a 10x magnification with numerical aperture 0.3. Further settings were: Immersion medium air, exposure time 400 ms, gain 0, offset 0, binning 1, grid wide field, Cy2: excitation 480 nm, emission 527 nm, Cy3: excitation 545 nm and emission 610 nm. The stained area per vessel was obtained by dividing the area of vessels by their total number. To obtain intensity of the EBA-immunofluorescence signal per µm² vessel area, representing the vessels' EBA content, the total fluorescence intensity of the microscopic field was measured by Velocity 4.3.2 and manually corrected for the surprisingly low background staining. Finally, the background-corrected intensity of the EBA-immunofluorescence signal was divided by the previously obtained vessel area. For control issues on the ischemia-affected hemisphere, number and area of EBA-immunopositive vessels, area per vessel and the intensity of the EBA-immunofluorescence signal per µm² vessel area were also measured in the hippocampus. Due to the nature of embolic stroke with varying infarct degrees and locations between animals,26 we also analyzed corresponding regions of the contralateral (non-ischemic) hemisphere, allowing the calculation of a ratio between the ischemic and non-ischemic hemispheres for vessels' EBA content, finally enabling intergroup comparison. In sections from sham-processed animals devoid of FITC-albumin leakage, 4 microscopic fields in both hemispheres (each 0.55 mm²), covering hypothalamus, striatum, parietal cortex and hippocampus, were analyzed as described above. For illustration of data, high-resolution images were taken from selected sections using the confocal laser-scanning microscopes 510
cerebral ischemia, anti-EBA reliably detected

Quantification of BBB integrity and measurement of BBB-related serum markers

Quantification of BBB permeability in ischemia-affected brain regions using FITC-albumin as leakage marker as well as acquisition of MMP-2 and MMP-9 serum levels were described previously. The resulting ratio of extra-/intravascular Cy2-enhanced FITC-albumin as surrogate for BBB integrity as well as MMP serum levels were utilized for calculations in the present study.

Statistical analysis

Calculations were performed with SPSS version 20.0 (IBM Corp.; New York, NY, USA). The Wilcoxon test was utilized to check for statistical significance between values of the ischemic vs the non-ischemic hemisphere at a single time point. For inter-group comparisons at different time points the t-test was used. To explore interrelations between assessed variables, partial correlations, controlled for observation time, were applied. In general, a P<0.05 was considered as statistically significant.

Results

EBA-immunoreactivity after experimental ischemic stroke

In sham-proceded animals devoid of focal cerebral ischemia, anti-EBA reliably detected cerebral vessels in neocortical and subcortical regions except of the circumventricular organs, e.g., the choroid plexus and some pial vessels showing an incomplete staining. Compared to the neocortex and striatum, the number of EBA-immunopositive vessels appeared to be symmetrically reduced in the hippocampus.

To ensure the specificity of endothelial EBA-immunolabeling in ischemia-affected brain regions, triple fluorescence labeling was applied resulting in nearly complete congruence of EBA-immunoreactivity and the endothelium as visualized by Solanum tuberosum lectin (STL), confirmed by FITC-albumin that remained intravascularly (Figure 1A). In contrast to the surrounding tissue with regular BBB integrity and to the non-affected contralateral hemisphere, vessel-associated EBA-immunoreactivity was found to be rarefied in areas with ischemia-related BBB dysfunction, indicated by FITC-albumin leakage (Figure 1B). Apart from the striatum as primary region of ischemia, this feature was also seen in the neocortex if affected by ischemia. A remarkably weak background staining in all animals, facilitated quantitative analyses of the red fluorescence signal.

Diverse aspects of EBA-immunoreactivity in regions with decreased BBB integrity

Sham-proceded animals showed no differences in number of EBA-immunopositive vessels, stained area of vessels and area per vessel between both hemispheres (data not shown). Compared to corresponding regions in the non-affected hemisphere, 5 h after ischemia onset the number of EBA-immunopositive vessels was significantly reduced in the area of decreased BBB integrity as indicated by FITC-albumin leakage (Wilcoxon, P=0.02; overview in Table 1), while stained vessel area provided the same tendency, scarcely missing statistical significance (Wilcoxon, P=0.09). Similar findings were obtained at 25 h after ischemia onset with a significant decrease in the stained area of vessels (Wilcoxon, P=0.04) and a reduction in number of EBA-immunopositive vessels (Wilcoxon, P=0.05), simultaneously occurring with FITC-albumin leakage (Figure 2). Surprisingly, the stained area per vessel did not change at both time points (Wilcoxon, 5 h: P=0.12; 25 h: P=0.33).

Additionally, at both time points no significant changes in number of EBA-immunopositive vessels, stained vessel area or area per vessel were found between the ipsi- and contralateral hippocampus (Wilcoxon, P>0.05 each).

Vessels’ EBA content in regions with BBB dysfunction

Five and 25 h after ischemia onset, intensity of EBA-immunofluorescence signal per µm² vessel area as surrogate for vessels’ EBA content tended to be increased in areas with FITC-albumin leakage compared to corresponding regions on the non-affected hemisphere, however, failing statistical significance (Wilcoxon, P=0.12 and P=0.17). Even clearer in the hippocampus, no significant change in vessels’ EBA content was found at both time points (Wilcoxon, P>0.05 each). Concerning the general time course, regions with decreased BBB integrity exhibited a slightly increased EBA-immunofluorescence signal per µm² vessel area towards the 25 h time point, which, however, failed statistically significance when compared to 5 h (t-test, P=0.58).

Interrelations among number of EBA-immunopositive vessels, stained vessel area, vessels’ EBA content and degree of BBB dysfunction

As previously described, in the applied model of focal cerebral ischemia affected brain regions displayed a decreased tendency of the extra-/intravascular FITC-albumin ratio as surrogate for BBB dysfunction between 5 and 25 h after ischemia onset, indicating an increase in BBB integrity in the early phase of stroke. However, in areas with increased BBB permeability.
ability no significant correlation (corrected for time points) was found between the number or area of EBA-immunopositive vessels and the ratio of extra-/intravascular FITC-albumin (r=0.22, P=0.47; r=0.05, P=0.87).

When further corrected for serum levels of MMP-2 and -9, the ratio of vessels’ EBA content and the intensity of EBA-immunofluorescence signal per µm² vessel area correlated in an inverse manner with the ratio of extra-/intravascular FITC-albumin in regions of increased BBB permeability (r= –0.68, P=0.02; r=0.60 P=0.05; Figure 3). No correlation was found between intensity of either EBA-immunosignal per µm² vessel area or ratio of vessels’ EBA content and serum levels of both MMP-2 and -9 (MMP-2: r= –0.03, P=0.91; r= –0.2, P=0.52; MMP-9: r= –0.02, P=0.95; r= –0.11, P=0.72).

Discussion

Following previous investigations, regular establishment of EBA has been considered to be associated with an intact cerebral vasculature as well as unaltered BBB integrity, although clear evidence for such a functional perspective is still missing, especially under...
ischemic conditions. To the best of our knowledge, the present study for the first time provides descriptive details and spatio-temporal characteristics of EBA-immunoreactivity in brain regions with decreased BBB integrity due to focal cerebral ischemia using an embolic model of stroke in rats.

Five and 25 h after ischemia induction, both number of EBA-immunopositive vessels and stained area of vessels were reduced in regions with increased BBB permeability. At first glance, these findings confirm previous reports about a decrease of EBA-immunopositive vessels under ischemic conditions, but contrary to Lin and Ginsberg as well as Lu et al., the stained area per vessel did not change significantly at both time points investigated, probably indicating a lack of vessel fragmentation. This discrepancy might be attributed to several reasons, e.g., differing stroke models (embolic vs filament-based) and localization of measurement (associated with impaired BBB vs infarcted area itself). Indeed, Michalski et al. could show that the applied model of embolic stroke in rats leads to FITC-albumin leakage that is restricted to ischemia-affected areas, but *vice versa* loss of neurons could naturally exceed the area of increased BBB-permeability. Mismatch between previous immunohistochemical studies might also result from considerably differing staining methods, e.g., SMI-71 was utilized in dilutions ranging from 1:100 to 1:2000 for immunoperoxidase labeling with diaminobenzidine as chromogen.

So far, a loss of EBA-immunopositive vessels has predominantly been interpreted as a loss of EBA-immunoreactivity linked to an increase in BBB permeability, but specific data on the functional role in focal cerebral ischemia are lacking, especially whether reduced EBA-immunolabeling is the cause or the consequence of BBB dysfunction. Bhattacharjee et al. showed that papaverine, when applied *in vivo*, resulted in an opening of the BBB as
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