CD84 Links T Cell and Platelet Activity in Cerebral Thrombo-Inflammation in Acute Stroke

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RATIONALE: Ischemic stroke is a leading cause of morbidity and mortality worldwide. Recanalization of the occluded vessel is essential but not sufficient to guarantee brain salvage. Experimental and clinical data suggest that infarcts often develop further due to a thromboinflammatory process critically involving platelets and T cells, but the underlying mechanisms are unknown.

OBJECTIVE: We aimed to determine the role of CD (cluster of differentiation)-84 in acute ischemic stroke after recanalization and to dissect the underlying molecular thromboinflammatory mechanisms.

METHODS AND RESULTS: Here, we show that mice lacking CD84—a homophilic immunoreceptor of the SLAM (signaling lymphocyte activation molecule) family—on either platelets or T cells displayed reduced cerebral CD4+ T-cell infiltration and thrombotic activity following experimental stroke resulting in reduced neurological damage. In vitro, platelet-derived soluble CD84 enhanced motility of wild-type but not of \( \text{Cd84}^{-/-} \) CD4+ T cells suggesting homophilic CD84 interactions to drive this process. Clinically, human arterial blood directly sampled from the ischemic cerebral circulation indicated local shedding of platelet CD84. Moreover, high platelet CD84 expression levels were associated with poor outcome in patients with stroke.

CONCLUSIONS: These results establish CD84 as a critical pathogenic effector and thus a potential pharmacological target in ischemic stroke.

Key Words: blood platelets ■ brain ischemia ■ infarction ■ stroke ■ T lymphocytes

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stroke is one of the major causes of death and disability worldwide with limited treatment options. In acute ischemic stroke (IS), the primary therapeutic goal is the rapid reconstitution of cerebral blood flow, achievable either by treatment with recombinant tPA (tissue-type plasminogen activator) or mechanical thrombectomy (MTE). MTE, which is experimentally mimicked by the most widely used animal stroke model, the transient middle cerebral artery occlusion (IMCAO) model—is particularly effective for severe clinical syndromes caused by occlusion of large cerebral arteries as the internal carotid artery (ICA) or the middle cerebral artery. However, despite recent advances in MTE with high recanalization rates of up to \( \approx 80\% \), the therapeutic efficacy of vessel reopening remains limited. Some studies indicate that infarction may progress significantly during reperfusion, and for many patients, infarction has already progressed unfavorably before hospital arrival preventing...
Novelty and Significance

What Is Known?
- Ischemia/reperfusion (I/R) injury in cerebral ischemia depends on the concerted action of platelets and T cells, a process referred to as thromboinflammation, but the molecular basis of the interplay of these 2 cell types is only partially understood.
- CD (cluster of differentiation) 84 is highly expressed on different immune cell populations and platelets. It acts as a homophilic cell adhesion molecule and serves as a functional coreceptor in lymphocytes.
- In activated platelets, CD84 has been shown to undergo ectodomain shedding, resulting in the release of soluble CD84. However, the (patho)physiological significance of soluble CD84 remains elusive.

What New Information Does This Article Contribute?
- Mouse models of experimental stroke identified CD84 as critical pathogenic effector in ischemic stroke.
- Platelet surface abundance of CD84 is reduced locally and directly within the ischemic circulation during hyperacute total vascular occlusion in stroke patients.
- High platelet CD84 expression levels are associated with poor outcome in stroke patients.
- CD84 links platelet and T-cell activities during cerebral I/R injury and aggravates neuronal damage.

In acute ischemic stroke, the primary therapeutic goal is reconstitution of cerebral blood flow. However, in the majority of patients, infarcts worsen despite successful recanalization due to I/R injury. Here, we demonstrate that CD84 promotes infarct progression following cerebral I/R injury by linking T-cell and platelet activities in the acutely ischemic brain, as mice lacking CD84 in platelets or T cells displayed smaller infarct sizes. In vitro studies revealed that platelet-derived soluble CD84 enhances motility of wild-type but not of Cd84-deficient T cells, suggesting that homophilic soluble CD84:CD84 interactions trigger this response. A critical role for CD84 in stroke progression was further supported by direct human observation during the hyperacute stroke phase of total vascular occlusion. Local probing of occlusive ischemic blood showed that platelet expression of CD84 decreases in the ischemic arterial compartment during persisting vessel occlusion, indicating shedding of platelet CD84. In addition, high systemic platelet CD84 expression levels were associated with poor outcome in stroke patients. Collectively, our data establish CD84 as key molecule linking detrimental T-cell and platelet activities in acute stroke and indicating that this receptor might be a promising pharmacological target to limit I/R injury.
key orchestrators of experimental cerebral I/R injury that promote inflammation and subsequent thrombotic activity in the microcirculation. Microvascular thromboinflammation can be efficiently prevented by blocking early steps of platelet adhesion and activation via GP (glycoprotein) Ib and GPVI receptors, respectively.11 Importantly, the pathogenic effect of platelets is linked to that of T cells as depletion of platelets in Rag1−/− mice abolished the detrimental effect of adoptively transferred CD4+ T cells.12 These findings led to the recent concept of thromboinflammation as the driving force underlying I/R injury in the ischemic brain.13–15 Thereby platelets act in concert with T cells to cause further brain injury, but the underlying mechanisms and the molecular link between the two cell types have remained elusive.

CD84 is a member of the SLAM (signaling lymphocyte activation molecule) family and acts as a homophilic cell adhesion molecule highly expressed on different immune cell populations and platelets.16–18 The N-terminal ectodomain of CD84 mediates the homophilic interaction between CD84 proteins,17,18 whereas the C-terminal intracellular portion of CD84 bears 2 immunoreceptor tyrosine-based switch motifs, which can bind the intracellular adapters SAP (SLAM-associated protein; also termed SH2D1A) and EAT-2 (Ewing sarcoma activated transcript 2).16,19 CD84 ligation triggers immunoreceptor tyrosine-based switch motif phosphorylation and SAP recruitment, resulting in enhanced IFN (interferon)-γ production and proliferation in T cells stimulated with low doses of anti-CD3 antibodies.20 CD84 has been established as a functional coreceptor in lymphocytes that facilitates prolonged B:T-cell interactions required for germinal center formation.20 Thus, while CD84 appears to have a role in immune cell activation, its function in platelets is not well understood. Upon platelet activation, CD84 is shed from the platelet surface,21 but the (patho)physiological significance of soluble CD84 (sCD84) is unknown. Notably, CD84 deficiency had no effect on the hemostatic or thrombotic function of platelets in mice in vitro and in vivo.22

Here, we show that platelet-derived sCD84 acts on CD4+ T-cell CD84 leading to enhanced CD4+ T-cell motility in vitro and aggravated infarct growth following cerebral I/R. Finally, we corroborated the clinical relevance of these findings by directly demonstrating reduced platelet CD84 levels in the ischemic circulation of stroke patients indicating CD84 shedding during the hyperacute stage of stroke. Furthermore, human platelet CD84 expression levels were associated with poor outcome in acute IS.

**METHODS**

The data that support the findings of this study are available from the corresponding author upon reasonable request. All animal experiments and the analysis of the corresponding data were performed blinded.

**Experimental Animals**

Mice used in this study were matched for age (7–14 weeks), sex, and genetic background. Experiments were conducted in accordance with the regulations of the local authorities (Regierung von Unterfranken) and performed in accordance with the current Animal Research: Reporting of In Vivo Experiments guidelines (https://www.nc3rs.org.uk/arrive-guidelines). Exclusion criteria and dropout rates are provided in Table I in the Data Supplement.

**Sample Size Calculation for tMCAO Experiments**

Assuming a reduction of infarct volume of 30% as functionally relevant and an SD of 20% to the respective mean values, a group size of 8 to 10 was necessary to show this effect with a power of 0.8 and a probability of a type I error of <0.05 (calculated with GraphPad StatMate 2.00).

**Animal Treatment**

Heart rate and blood pressure were determined in isoflurane anesthetized mice using a Pressure Meter LE5001 (Harvard Apparatus, Boston, MA) according to the manufacturer’s instructions. For T-cell transfer experiments into Rag1−/− and Cd84−/− mice, splenic CD4+ T cells were isolated by negative selection (Miltenyi Biotech). Cells were injected intravenously (750 000 cells/mouse) 1 day before tMCAO, and successful T-cell transfer was controlled 1 hour after iv injection by flow cytometry of blood immune cells.10

**Focal Ischemia Model**

Focal cerebral ischemia was induced in 10- to 14-week-old C57BL/6, Cd84−/−,21 Rag1−/−,23 Cd84fl/fl,PF4-Cre−/−, or Cd84fl/fl,PF4-Cre mice by tMCAO as described previously.24 Inhalation anesthesia was induced by 2% isoflurane. The duration of the surgical procedure per animal was kept below 10 minutes. A silicon rubber-coated 6.0 nylon monofilament (6021PK10; Doccol, Redlands, CA) was advanced through the carotid artery up to the origin of the middle cerebral artery to the origin of the middle cerebral artery infarction. After an occlusion time of 60 minutes, the filament was removed allowing reperfusion. Animals were sacrificed 23 hours after reperfusion, and brains were checked for intracerebral hemorrhages. Neurological function was analyzed calculating a neuroscore (score, 0–10) based on the direct sum of the Grip test (score, 0–5) and the inverted Bederson score (score, 0–5).25–27 For assessment of the long-term neurological outcome, the occlusion time was reduced to 30 minutes, and Bederson score, modified neurological severity score, corner test, body swing test, grip test, latency to move, and adhesive tape removal test were performed daily as described previously.

**Infarction Size Measurement**

The extent of infarction was quantitatively assessed 24 hours after occlusion. Animals were sacrificed, and brains were cut in three 2-mm-thick coronal sections. The slices were stained for 20 minutes at 37°C with 2% 2,3,5triphenyltetrazolium chloride (Sigma-Aldrich; 2% [w/v] solution) to visualize the infarctions.29 Edema-corrected infarct volumes were calculated by planimetry.
Histology

Histology and immunohistochecmistry were performed according to standard procedures, and all antibodies were validated using isotype controls (not shown).24 Cryo-embedded coronal brain sections (2 mm) were cut into 10-µm-thick slices. Every tenth slice was used for evaluation. The following antibodies were used: monoclonal antibodies: anti-CD11b (MCA711; Serotec), anti-Ly6B (MCA771G; Bio-Rad), and anti-CD31 (MCA2398; Bio-Rad) and polyclonal antibodies: anti-CD4 (100506; BioLegend) and anti-albumin (ab207327; Abcam). Apoptotic neurons were visualized by double immunolabeling of mouse anti-NeuN (neuronal nuclei) antibody (MAB377; Merck, Darmstadt, Germany) and TUNEL (Terminal deoxynucleotidyl transferase dUTP nick end labeling) in situ cell death detection kit TMR red (12156792910; Merck). For quantification of occluded microvessels, brain slices were stained with hema-topoxylin and eosin. Afterward, the numbers of occluded and opened vessels per hemisphere were counted to determine the percentage of occlusions as described previously.24 All immunohistologic stainings were analyzed and acquired using a Nikon Eclipse 50i microscope or a Leica SP8 confocal microscope. Representative images were chosen among the images of one genotype to best reflect the mean values of this genotype.

Cell Separation and Flow Cytometry

For platelet analysis, peripheral blood was collected in heparin (20 U/mL in TBS [Tris-buffered saline]) and diluted in Tyrode-HEPES buffer and incubated with the indicated antibodies. CD84 expression (in-house generated JER1-FITC21 [fluorescein isothiocyanate]) of GPⅠb/Ⅲa–positive platelets (in-house generated JON6-PE30) and expression of major platelet receptors was analyzed using FACS Celesta or FACS Calibur (Becton Dickinson, Heidelberg, Germany). In the remaining blood, RBCs (red blood cells) were lysed using ACK (ammonium-chloride-potassium) buffer and immune cells were stained with the indicated antibodies for 30 minutes on ice in FACS buffer. For the isolation of brain-infiltrating mononuclear cells, a Percoll (GE Healthcare) density gradient (50%/30%) was used as a control. Tissue homogenization, RNA isolation, and real-time polymerase chain reaction procedures using antibodies against albumin (ab106582; Abcam), CD84 (in-house generated JER121), GAPDH (G9545; Sigma), and anti-β-actin (A5441; Sigma-Aldrich).31

Real-Time Polymerase Chain Reaction

Tissue homogenization, RNA isolation, and real-time polymerase chain reaction were performed as described.31 Relative gene expression levels of TNFa (tumor necrosis factor-alpha; assay ID: Mm 00443258_m1; Applied Biosystems), IL (interleukin)-β (assay ID: Mm 00434228_m1; Applied Biosystems), MMP (matrix metalloproteinase)-2 (assay ID: Mm 00439498_m1; Applied Biosystems), and MMP-9 (assay ID: Mm 00442991_m1; Applied Biosystems) were analyzed with a fluorescent TaqMan technology. As an endogenous control Gapdh (TaqMan Predeveloped Assay Reagent for gene expression, part number 4352339E; Applied Biosystems) was used. Polymerase chain reaction was performed using the StepOnePlus Real-Time PCR System (Applied Biosystems).

T-Cell Migration Assays

The migratory capacity of CD4+ T cells was measured using ibidi µ-Slides VI coated with poly-D-lysine (10 µg/mL; Merck) and laminin (20 µg/mL; Merck) or ibidi Glass Bottom µ-Slides 8 well cultured with primary primary murine brain microvascular endothelial cells as described previously. The migration assays were either performed in DMEM high glucose (31053-028; Thermo Fisher) with penicillin/streptomycin (1%, P4333; Merck) and B27 supplement (2%, 17504044; Thermo Fisher) or Roswell Park Memorial Institute (RPMI) (12633020; Sigma) with glucose (1%, 59202C; Merck) and penicillin/streptomycin (1%, P4333; Merck). Each condition (phorbol-12-myristate-13-acetate [8 µg/mL; Merck] and CCL20 [C-C chemokine cysteine motif chemokine ligand 20; 24.5 µg/µL; PeproTech]), Fc-proteins (0.2 µg/mL), washed WT or Cd84-/- platelets (diluted 50:1 with CD4+ T cells), platelet-releasate (PLT-R; 1:1 diluted with migration media) was performed in duplicates or triplicates. PLT-R was obtained by stimulating platelets (500000/µL) with 10 µg/mL of the GPⅠb agonist CRP (collagen-related peptide) for 15 minutes; subsequently, the platelet supernatant was harvested (5 minutes with 800 g, followed by 5 minutes with 22000 g, 4°C). The GPⅠb-Fc fusion protein (control-Fc) that did not affect CD4+ T-cell migration compared with vehicle (not shown) served as control for CD84-Fc (recombinant sCD84 fused to the Fc part of human IgG1); both Fc fusion proteins were purified in-house from transfected HEK 293 (human embryonic kidney) cells using standard techniques. For time-lapse video microscopy, MACS (magnetic-activated cell sorting) isolated mouse CD4+ T cells (130-104-454; Miltenyi; 100 cells/µL) were added to the chamber. Images were collected every 30 seconds for 30 minutes on a Leica DMIB inverted microscope with 20× objective. Cells were tracked with LAS X Imaging software and analyzed with ImageJ 1.51 software.
Analysis of Human Blood Samples

Both the local ischemic and systemic blood samples were drawn into citrate phosphate dextrose adenine monovettes (S-Monovette; Sarstedt, Nümbrecht, Germany). Samples were diluted after blood collection in PBS, and platelets were stained using anti-CD42b-FITC (SZ2; Immunotech SAS, Marseille, France) and anti-CD84-PE antibodies (CD84.1.2.1; BioLegend, San Diego). After an incubation period of 30 minutes, the samples were analyzed by flow cytometry instrument FACS Calibur (Becton Dickinson, Franklin Lakes). CD84 expression was defined as mean fluorescence intensity of PE (phycoerythrin) in platelets gated by CD42b positivity.

Clinical Observation of Local Human Arterial Blood Samples

The protocol of the prospective study of local ischemic and systemic arterial blood samples during hyperacute stroke was approved by the local ethics committee (approval number 135/17) and performed as described previously. During first-ever IS, it was determined by CT angiography that the embolic occlusion location matched the ICA-T/M1 segment. MTE was then indicated based on clinical and radiological criteria recommended by current guidelines. All procedures were performed by board-certified neurointerventional experts or by supervised neurointerventional fellows. A detailed description of the MTE technical procedural steps used in this study is given elsewhere. Briefly, microcatheter navigation was performed into the occluded vascular field via a 0.014 microinch microwire over a coaxial guide catheter system positioned in the cervical common carotid arteries and ICAs. Having reached this position, one local arterial blood sample was drawn by microcatheter aspiration. Two further samples were drawn after recanalization at the systemic levels of the proximal internal carotid and common femoral arteries.

Clinical Observation of the SICFAIL Cohort

To assess the transferability of the findings from mouse model into humans, the association of CD84 expression on platelets and stroke severity at day 3 after hospital admission was analyzed in a substudy within the SICFAIL study (Stroke-Induced Cardiac Failure in Mice and Men). SICFAIL is an ongoing, prospective cohort study, aiming to describe the natural course of cardiac function in 750 unselected acute IS patients (DRKS00011615). The subsample of the SICFAIL study population, analyzed in this study, included 98 patients with the diagnosis of acute IS according to World Health Organization definition aged ≥18 years, who were hospitalized at the stroke unit of the University Hospital Würzburg between June 2016 and January 2017. Written informed consent was obtained prior study enrollment. The Ethics Committee of the Medical Faculty of the University of Würzburg (vote 176/13) approved the study. All patients or their legal representatives provided written informed consent to participate.

All patients underwent standard treatment and stroke unit monitoring. Information on stroke severity and acute treatment (eg, intravenous thrombolysis) was retrieved by review of patient records. Stroke severity was assessed by a neurologist according to the National Institutes of Health Stroke Scale (NIHSS). Additionally, all study participants underwent a personal standardized interview to collect information on demographic characteristics and comorbidities. Based on previously published data, poor outcome was defined before analyses as NIHSS ≥5 on day 3 of hospitalization, representing moderate or severe stroke.

For CD84 analysis, blood was drawn the morning after study enrollment in tubes containing citrate phosphate dextrose adenine and analyzed as described above.

Statistical Analyses

All data from animal experiments are given as box plots including median with the 25th percentile, the 75th percentile, minimum deviation, and maximum deviation except for the migration assays, which are depicted as scatter plots including the mean and SEM. Human data are presented as median and interquartile range (IQR; 25th and 75th percentile) or, in case of categorical data, frequencies and percentages, respectively. Cell migration data were analyzed by 1-way ANOVA with Bonferroni post hoc tests. All other data were analyzed by nonparametric tests (specified in the respective figure and table legends) due to small sample sizes. Corrections for multiple tests within one experimental assay in animal studies were done by appropriate post hoc tests (eg, Dunn test for Kruskal-Wallis tests) or the Holm-Sidak method. No corrections for multiple testing were made across assays. For nonparametric repeated measures of human data, we used Friedman rank test followed by Dunn multiple comparison test. In case of frequencies, χ² test or Fisher exact test was used to compare groups. Multivariable logistic regression analysis, adjusted for age and NIHSS at baseline, was used to identify the association of CD84 expression and poor outcome on day 3 of hospitalization (NIHSS score ≥5). For sensitivity analyses, the model was recalculated by further adjusting for comorbidities including previous stroke or transient ischemic attack, coronary heart disease, hypertension, type 2 diabetes mellitus, and atrial fibrillation. The α-level was set to 0.05 (2 sided). P values displayed in figures are corrected for multiple testing, as described above; P > 0.05 is not displayed in the figures (albeit all data were analyzed with the statistical tests provided in the figure or table legends). For statistical analysis, the GraphPad Prism 5.0 and 8.2.1 software package (GraphPad Software) and SAS 9.4 (SAS Institute, Inc, Cary, NC) were used.

RESULTS

Cd84<sup>−/−</sup> Mice Have Less CD4<sup>+</sup> T-Cell Infiltration Into the Ischemic Hemisphere and Smaller Infarcts Following IS

We have previously shown that Cd84<sup>−/−</sup> mice display no overt phenotype and exhibit no defects in platelet function in hemostasis or thrombosis. Of note, brain morphology was unaltered in Cd84<sup>−/−</sup> mice as assessed by ink staining of the vasculature and Nissl staining of brain slices (Figure I in the Data Supplement). To test for a possible role of CD84 in thromboinflammation, Cd84<sup>−/−</sup> mice were subjected to 1 hour of tMCAO and 23 hours of reperfusion. Strikingly, median infarct...
volumes in the mutant mice were significantly reduced compared with WT littermates at 24 hours after tMCAO as measured by triphenyltetrazolium chloride staining (97.8 [IQR, 75.7–124.2] versus 83.8 [IQR, 45.1–92.2] mm3; \( P = 0.0198 \); Figure 1A and 1B), while heart rate (median, 532.8 bpm [IQR, 520.2–544.4] versus 533.5 bpm [IQR, 512.6–588.2]) and blood pressure in naive mice were indistinguishable (not shown) and local cerebral blood flow during and after middle cerebral artery occlusion was comparable (Figure IIA in the Data Supplement). The reduction in infarct volumes corresponded with improved neurological outcome in \( Cd84^{-/-} \) mice as reflected by a higher mean neuroscore (4.3±1.0 versus 5.2±1.5, respectively; \( P = 0.0269 \); Figure 1C). Sex can have a significant impact on stroke outcome.40 Therefore, we subjected male and female mice to 60 minutes of tMCAO. Gender-specific evaluation revealed significantly smaller infarctions in both male and female \( Cd84^{-/-} \) mice (Figure IIB and IIC in the Data Supplement). To investigate the role of CD84 in long-term functional outcomes following stroke, we analyzed sensorimotor deficits of male \( Cd84^{-/-} \) mice and control littermates over 7 days after 30 minutes of tMCAO (Figure 2). In line with the results following 60-minute tMCAO, \( Cd84^{-/-} \) mice displayed less severe functional defects on day 1 following 30-minute tMCAO as compared with WT mice (modified neurological severity score, 3 [IQR, 2.5–4.0] versus 6 [IQR, 4.8–8.3]; \( P = 0.0111 \); Figure 2A). This difference between the 2 genotypes persisted for the 7-day observation period as confirmed by the modified neurological

Figure 1. \( Cd84^{-/-} \) mice are protected after transient middle cerebral artery occlusion (tMCAO).
A, Representative triphenyltetrazolium chloride–stained 2-mm brain slices from 1 WT and 1 \( Cd84^{-/-} \) mouse 24 h after tMCAO. Bar=1 cm. B, Infarct volumes 24 h after tMCAO of WT (n=16) and \( Cd84^{-/-} \) mice (n=18; \( P =0.0198 \)). C, WT (n=18) and \( Cd84^{-/-} \) (n=21) neuroscore at 24 h after tMCAO (\( P =0.0269 \)). Statistical significances analyzed by Mann-Whitney U test. *\( P <0.05 \).

Figure 2. Role of CD (cluster of differentiation) 84 in long-term functional outcome following stroke.
Neurological outcome over time in WT (gray boxes) and \( Cd84^{-/-} \) mice (green boxes) as assessed by modified neurological severity score (mNSS; A), Corner test (B), and Bederson score (C) after 30-min transient middle cerebral artery occlusion (tMCAO; n=7–10 mice per group). Each symbol represents one individual. Statistical significances between WT and \( Cd84^{-/-} \) mice were determined using the Holm-Sidak correction on \( P \) obtained by Mann-Whitney U tests. *\( P <0.05 \), **\( P <0.01 \), and ***\( P <0.001 \); exact n-numbers and \( P \) are listed in Table V in the Data Supplement.
severity score or different additional tests (Figure 2B and 2C; Figure III in the Data Supplement). Three WT but no Cd84−/− mice had to be euthanized due to severe impairments (Figure IIIA in the Data Supplement).

Next, we assessed accumulation of lymphocytes and monocytes in the brain after tMCAO. Cd84−/− mice showed significantly reduced numbers of CD4+ T cells in the infarcted brain (reduction by ≈50%) compared with

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**Figure 3.** CD (cluster of differentiation) 4+ T-cell recruitment to the postischemic brain is diminished in CD84-deficient mice. 
A, Representative immunocytologic stainings and quantification of brain-infiltrating CD4+ T lymphocytes in the ipsilateral hemisphere on day 1 after transient middle cerebral artery occlusion (tMCAO) of WT (wild type) and Cd84−/− mice (n=5 mice per group; P=0.0328). Bar=100 µm. B, Representative CD11b immunoreactivity and quantification of CD11b+ cells in the ipsilateral hemisphere 24 h after tMCAO of WT and Cd84−/− mice (n=5 mice per group; P=0.5476). Bar=100 µm. C, Representative images and quantification of Ly6B.2+ cells in the ipsilateral hemisphere 24 h after tMCAO of WT and Cd84−/− mice (n=5 mice per group; P=0.5794). Bar=100 µm. D, Representative hematoxylin and eosin (H&E) staining and quantification of the percentage of occluded vessels in the infarcted hemispheres of WT and Cd84−/− mice (n=5 mice per group; P=0.0079). White arrows indicate occluded vessels; blue arrows indicate patent vessels. Bar=100 µm. Statistical significances analyzed by Mann-Whitney U test. *P<0.05 and **P<0.01. DAPI indicates 4′,6-diamidino-2-phenylindole.
WT mice (Figure 3A), whereas the amount of CD11b+ cells (Figure 3B) and the accumulation of neutrophils (Ly6B.2+ cells; Figure 3C) was comparable between the 2 groups. These data were confirmed by flow cytometry (Figure IV in the Data Supplement). Furthermore, the percentages of occluded vessels in the ipsilateral hemisphere was reduced by 35% in Cd84−/− mice 24 hours after stroke induction (Figure 3D), and less platelet deposition in the ipsilateral hemisphere of Cd84−/− mice was observed (not shown). These data demonstrated that Cd84 deficiency is protective after tMCAO and that Cd84 contributes to CD4+ T-cell recruitment into the infarcted brain territory. We analyzed neuronal apoptosis and blood-brain barrier damage in WT and Cd84−/− mice 24 hours after stroke induction (Figure VA through VC in the Data Supplement). In line with smaller infarct volumes, neuronal survival and blood–brain barrier (BBB) function in the mutant mice were significantly improved, whereas the expression levels of TNFα, IL-1β, MMP-2, and MMP-9 did not differ significantly between WT and Cd84−/− mice (Figure VD through VG in the Data Supplement).

### CD84 Expression on CD4+ T Cells and Platelets Is Required to Promote Infarct Growth After tMCAO

Then, we assessed the relevance of CD4+ T cell–expressed CD84 for infarct progression by performing adoptive T-cell transfer experiments. Immunodeficient Rag1−/− mice, which lack B and T cells, or Cd84−/− mice received CD4+ T cells from WT or Cd84−/− mice 24 hours before subjecting them to tMCAO. While Rag1−/− mice displayed small infarcts 24 hours after tMCAO,5,10,41 fully evolved infarctions were found in Rag1−/− mice after adoptive transfer of WT CD4+ T cells (Figure 4A and 4B). In sharp contrast, Rag1−/− mice reconstituted with Cd84−/− CD4+ T cells had significantly smaller infarcts (median, 100.9 [IQR, 74.8–126.0] versus 28.6 [IQR, 22.2–111.4] mm3; P=0.0050; Figure 4A and 4B) and better neurological outcome (Figure VI in the Data Supplement) at day 1 after stroke than those transplanted with WT CD4+ T cells. Of note, lymphocyte and thymocyte subsets were indistinguishable between WT and Cd84−/− mice (not shown), nor did the secretion of typical cytokines from stimulated CD4+ T cells differ (Table II in the Data Supplement).

In contrast to adoptive CD4+ T-cell transfer in Rag1−/−, infarct volumes and functional outcome did not differ when adoptively transferring WT or Cd84−/− CD4+ T cells into Cd84−/− mice (Figure 4A and 4B) indicating that an interaction of CD84+ T cells with another CD84-expressing cell type is required to induce I/R injury. To address a possible role of platelets in this setting, we generated Cd84fl/fl,PF4-Cre mice. Knockout efficacy was confirmed using flow cytometry and Western blot (Table III in the Data Supplement; Figure VIIA through VIIE in the Data Supplement). As reported previously,22 Cd84 deficiency did not affect classical platelet activation (Figure VIIID and VIIE in the Data Supplement). Cd84−/− mice developed significantly (>30%) smaller infarcts at day 1 after tMCAO when compared with littermate control mice (Cd84fl/fl,Cre- neg; median, 92.3 [IQR, 61.0–105.9] versus 63.4 [IQR, 26.2–79.4] mm3; P=0.041; Figure 4A and 4B) suggesting that platelet CD84 is required for the development of CD4+ T cell–dependent cerebral I/R injury. This is in agreement with the previous observation that platelet depletion abolished detrimental CD4+ T-cell effects during I/R in stroke after adoptive transfer into immune-deficient Rag1−/− mice.12 Of note, also in mice lacking Cd84 only on megakaryocytes and platelets, less CD4+ T cells were detected in the ischemic hemisphere 24 hours after tMCAO (Figure 4C), while the expression of Cre recombinase under the Pf4 promoter did not affect infarct sizes (Figure VIII in the Data Supplement).

### sCD84 Promotes CD4+ T-Cell Migration

To assess whether Cd84 is required for T-cell migration, we measured the motility of WT and Cd84−/− CD4+ T cells in a laminin/PDL (poly-D-lysine)-coated 2-dimensional in vitro system and did not observe differences between the 2 genotypes in response to phorbol-12-myristate-13-acetate or the CCL20 (Figure 5A; Figure IXA in the Data Supplement). We have previously shown that the activatory platelet receptor GPVI plays a critical role in cerebral ischemia,11,42 we tested the effect of the releasate from GPVI–stimulated WT platelets on CD4+ T-cell motility in this assay. Treatment of WT CD4+ T cells with PLT-R induced an increase of >25% in distance and speed of migrating CD4+ T cells, whereas this effect was absent in Cd84−/− T cells (Figure 5B; Figure IXB in the Data Supplement). Given the established homophilic binding of CD84 and the efficient ectodomain shedding of CD84 in activated platelets,12 we suspected platelet-derived sCD84 to mediate this effect. Indeed, the releasate of Cd84−/− platelets failed to increase the migratory capacity of WT T cells, strongly suggesting that platelet-derived sCD84 acts as a costimulatory factor on CD4+ T cells (Figure 5C; Figure IXC in the Data Supplement). To test this directly, we treated WT and Cd84−/− T cells with CD84-Fc or a control-Fc protein and compared their migratory behavior to that of untreated CD4+ T cells (control). Notably, WT but not Cd84−/− T cells responded to the presence of CD84-Fc with an increase in velocity and migrated distance compared with the control groups, indicating that the homophilic CD84 interactions are critical for the T-cell modulating effects of sCD84 (Figure 5D; Figure IXD in the Data Supplement). Of note, no significant difference in T-cell motility was observed in the presence of WT or Cd84−/− platelets (Figure IXE and I邢 in the Data Supplement).
Supplement). In contrast, CD84−/− PLT-R supplemented with CD84-Fc restored CD4+ T-cell motility to a level comparable to that induced with WT PLT-R (Figure 5E; Figure IXG in the Data Supplement) demonstrating that platelet-derived sCD84 enhances CD4+ T-cell migration. Notably, the migration-promoting effect of sCD84 on WT T cells was also observed in a migration assay in the presence of primary brain-derived microvascular endothelial cells, and results were indistinguishable on WT or CD84−/− endothelial cells (Figure 5F; Figure IXH in the Data Supplement). Together, these data demonstrated that platelet CD84 exerts a (co)stimulatory effect on CD4+ T cells and is a critical determinant of thromboinflammatory infarct progression in a murine model of IS.

Downregulation of Platelet CD84 Surface Levels in the Ischemic Arterial Compartment

Finally, we applied 2 strategies of clinical observation to analyze the potential role of platelet CD84 in acute stroke patients. First, we established an endovascular clinical method of local human in vivo sampling of ischemic
arterial blood already during the (hyper)acute stage of stroke with the samples taken directly within the occluded arterial vascular field. In this manner, local aspirates of 1.0 mL ischemic arterial blood were obtained via transfemoral arterial access immediately after endovascular microcatheter navigation into the center of the ischemic cerebral vascular field and before vessel reopening. This step was immediately ensued by therapeutic recanalization (ie, MTE). It was considered a particularly relevant a priori criterion for eligibility that the occlusion location had to precisely match the experimental situation (ie, ICA-T/M1 segment embolic occlusions). In a prospective cohort of large vessel occlusion stroke patients (n=23), we could then show that platelet surface abundance of CD84 (Table 1) was reduced locally within the ischemic circulation as compared with nonischemic intraindividual systemic platelet CD84 surface abundance indicating local shedding of CD84 from the platelet surface.

High Platelet CD84 Surface Levels Correlate With Poor Outcome in Stroke Patients

Second, local findings during the hyperacute stage were further extended by observation of a large prospective cohort in which we analyzed platelet CD84 expression in 98 participants of the SICFAIL study, to assess a possible link between platelet CD84 expression and disease course in acute IS patients. Median age was 67.0 years.

| Table 1. Downregulation of Platelet CD84 Surface Levels in the Ischemic Arterial Compartment |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Sampling Location | F               | C               | D               | F/C/D           | F vs C          | F vs D          | C vs D          |
| MFI, median (IQR)       | 21.5 (18.6–28.9) | 21.3 (17.9–27.4) | 20.7 (18.4–26.7) | 0.0239         | 0.1960         | 0.0296         | >0.9999         |

Platelet CD84 MFI measured by flow cytometry of arterial blood samples in IS, labeling according to the sampling location (common femoral artery; internal carotid artery; ischemic circulation distal to the occlusion site. Statistical significances analyzed by Friedman test with Dunn post hoc test; n=23 triplicates (F/C/D). C indicates carotid; D, distal; F, femoral; IQR, interquartile range; and MFI, mean fluorescence intensity.
(IQR, 55.0–76.0 years), 66% of the patients were men, and the median NIHSS score on admission was 3 (IQR, 2–6; Table IV in the Data Supplement). Nineteen patients had poor outcome at day 3 of hospitalization defined as NIHSS ≥5. Patients with poor outcome tended to be older, had a history of hypertension, and had higher NIHSS scores on admission. Although not statistically significant in univariate analysis (P=0.18), platelet CD84 surface abundance was identified to be independently associated with poor outcome after adjustment of age and baseline NIHSS score (odds ratio, 1.29 [95% CI, 1.05–1.60]; Table 2; Table IV in the Data Supplement) in an exploratory multivariable logistic regression approach. After further adjusting the model for comorbidities, CD84 remained independently associated with NIHSS score ≥5 at day 3 (odds ratio, 1.27 [95% CI, 1.02–1.59]) in sensitivity analyses (data not shown).

**DISCUSSION**

Our data strongly suggest that the homophilic interaction of platelet-derived sCD84 and CD4+ T cell–expressed CD84 represents a key pathogenetic mechanism in cerebral I/R injury. In T cells, CD84 has been shown to support interactions with B cells and TCR-induced IFNγ secretion. We and others have reported that platelet activation results in rapid ectodomain shedding of CD84, and here we show that sCD84 directly acts on CD4+ T cells and modulates their migratory behavior indicative of a (co)stimulatory effect. Studies in mice have shown that platelets are recruited very early to the postischemic cerebral microvasculature, and this appears to be in line with ectodomain shedding of CD84 in activated platelets, locally within the center of the human ischemic brain. Nevertheless, our results should be validated using further preclinical studies applying additional transient and permanent stroke models. In addition, it is important to note that, although the overall structure of the immune system in mice is similar to that in humans, existing differences in innate and adaptive immunity between mice and humans could interfere with clinical translation when targeting the interactions of T cells and platelets in acute stroke. Despite these potential differences, the concept of thromboinflammation as driving force in cerebral I/R injury has opened new therapeutic avenues based on the targeting of immunomodulatory functions of platelets while leaving their hemostatic response largely unaffected. Recently, several randomized controlled pilot trials could be completed testing the immunosuppressant drug FTY720, which rapidly reduces the number of circulating lymphocytes and is approved for mechanistic link between T cells and platelets in acute IS that may represent a new therapeutic target to reduce neuronal damage in acute IS. It appears as a paradox that platelet CD84 is not critical for arterial thrombosis but contributes to tissue damage during I/R in the brain. However, the pathophysiology of I/R injury does not require rethrombosis as substantiated by the fact that blockade of GPIIb/IIIa is ineffective. To investigate the relevance of this novel mechanism for acute stroke in humans, considerable efforts of clinical observation were undertaken. First, very early during the hyperacute phase of IS, and at the exact same occlusion location as in the experimental setting (ie, ICA-T/M1 segment occlusions), we observed a reduction of local platelet CD84 surface abundance. This reduction of CD84 is in line with ectodomain shedding of CD84 in activated platelets, locally within the center of the human ischemic arterial brain circulation.

Furthermore, platelet CD84 surface abundance in peripheral venous blood of patients with acute IS was identified to be independently associated with poor outcome after adjustment of age and baseline NIHSS in 98 participants of the prospective SICFAIL cohort study in an exploratory approach. Despite the limitations of this analysis, including the small sample size and relatively mild affected IS patients, these results clearly point to a pathogenic role of CD84 in the development of IS in human patients.

Collectively, our data indicate that targeting CD84 might be effective in reducing antithromboinflammatory neuronal damage in the acute ischemic brain with the potential to improve stroke outcomes. In this regard, it is noteworthy that attempts to further improve clinical outcome after thrombolysis by conventional antiplatelet drugs, for example, acetylsalicylic acid, clearly failed due to an increase in intracranial hemorrhages. In our study, both, the data from complementary in vivo and in vitro experimental mouse models and the observations in human stroke patients point toward a role of CD84 in cerebral ischemia. Nevertheless, our results should be validated using further preclinical studies applying additional transient and permanent stroke models. In addition, it is important to note that, although the overall structure of the immune system in mice is similar to that in humans, existing differences in innate and adaptive immunity between mice and humans could interfere with clinical translation when targeting the interactions of T cells and platelets in acute stroke. Despite these potential differences, the concept of thromboinflammation as driving force in cerebral I/R injury has opened new therapeutically avenues based on the targeting of immunomodulatory functions of platelets while leaving their hemostatic response largely unaffected. Recently, several randomized controlled pilot trials could be completed testing the immunosuppressant drug FTY720, which rapidly reduces the number of circulating lymphocytes and is approved for

**Table 2. Multivariable Logistic Regression to Assess the Association of Platelet CD84 Expression and Poor Outcome (NIHSS Score, ≥5) at Day 3 of Hospitalization, Adjusted for Age and NIHSS at Admission**

|          | Odds Ratio | 95% CI       | P Value |
|----------|------------|--------------|---------|
| Age, per y | 1.07       | 1.01–1.14    | 0.02    |
| Baseline NIHSS | 1.38       | 1.16–1.65    | 0.0004  |
| CD84 MFI | 1.29       | 1.05–1.60    | 0.02    |

CIs and P values were calculated by Wald test statistic. CD indicates cluster of differentiation; MFI, mean fluorescence intensity; and NIHSS, National Institutes of Health Stroke Scale.
the treatment of relapsing-remitting multiple sclerosis.46 Stroke patients treated immediately with FTY720 alone or in conjunction with r-tPA thrombolysis or thrombectomy showed decreased infarction.47–49 Our study now shows that targeting T-cell/platelet CD84 may become an alternative therapeutic strategy, which similarly does not affect hemostasis in the ischemic brain that is highly susceptible to bleeding complications. Our findings are likely to have broader implications also beyond the brain, since T cells have been identified as key players in I/R injury in many organs system such as the heart, kidney, and lung.

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