Contribution of polymorphonuclear neutrophils in the blood periphery to ischemic brain injury

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Three major lines of argumentation presently support a role of polymorphonuclear neutrophils (PMNs) in ischemic brain injury:

1. PMNs abundantly accumulate in ischemic brain tissue in response to stroke both in rodents and human patients, where they massively release enzymes, such as myeloperoxidase and elastase, and reactive oxygen species (ROS), which are known contributors to ischemic injury.^{1–3}
2. In mouse models of ischemic stroke induced by intraluminal middle cerebral artery occlusion (MCAO), the selective depletion of PMNs by anti-Ly6G antibody or prevention of PMN brain entry by CXCR2 or VLA-4 blockade significantly reduces ischemic injury and neurologic deficits.^{4,5}
3. In patients with ischemic stroke, high neutrophil counts or high neutrophil to lymphocyte ratios in the blood on admission are associated with poor neurologic outcome even when adjustments for age, sex, vascular risk factors, and stroke severity are made.^{6,7}

The combined evidence of these studies has prompted the idea that blood-derived PMNs are attracted into the ischemic brain, where they aggravate brain injury.⁸

In this issue of *Neurology: Neuroimmunology & Neuroinflammation*, Weisenburger-Lile et al.⁹ further strengthened and expanded this view, providing a detailed and comprehensive analysis of peripheral blood PMN characteristics in a cohort of 41 patients with acute ischemic stroke, which were compared with 22 healthy control subjects of the same age who were close relatives of patients. Blood samples were collected within 6 hours after stroke onset and, in a subgroup of patients, after 2 and 7 days. Flow cytometry studies revealed hyperactivation of circulating PMNs in the acute stroke phase, that is, within 6 hours after stroke, indicated by decreased CD62L and increased CD11b expression on PMNs, increased ROS production by unstimulated and stimulated PMNs, and increased circulating elastase levels in peripheral blood. The number of necrotic PMNs was increased from 2 to 7 days after stroke, whereas the concentration of neutrophil extracellular trap components in the serum was decreased. An increased percentage of senescent, that is, CXCR4^{bright}/CD62L^{dim} PMN was noted at all 3 time points examined. PMNs with the capacity to reversely transmigrate from inflamed tissues back into the blood, defined as CD54^{high}/CXCR1^{low}, were increased in patients with stroke. States of hyperactivation, senescence, and reverse migration were particularly pronounced in PMNs from patients exhibiting a high NIH Stroke Scale score (>12) on admission. The authors hypothesize that changes in PMN homeostasis may instrumentally contribute to ischemic brain injury, e.g., by promoting systemic inflammation, promoting blood-brain barrier breakdown, or inducing immunomodulation. The authors suggest that rebalancing PMN subsets or preventing reverse PMN transmigration might alleviate stroke consequences.

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Harmful neutrophil subsets in patients with ischemic stroke: Association with disease severity

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The clear strength of this study is a meticulously conducted characterization of PMN subsets that provides a first in-depth characterization of PMN changes in ischemic stroke. A downside is the choice of control subjects, which, representing healthy humans, do not exhibit a similar vascular risk profile. Differences in vascular risk factors and associated diseases (coronary heart disease and large artery atherosclerosis) may at least partly account for the observations made. Hence, confirmation studies in an independent patient cohort matched for risk factors and comorbidities will be required. Also, functional readouts of PMN activity such as migration should be studied in more detail,¹⁰ as they are further indicators of disease states.

What can we learn from these studies? In the past, we have perhaps focused too exclusively on the PMN-associated aggravation of ischemic injury inside the brain. By modulating immune responses in peripheral blood, reversely transmigrated PMNs might potentially deregulate the neurovascular unit or exert bystander effects on additional immune cells such as T cells in the blood, which on brain entry may aggravate ischemic damage.¹¹ In line with this hypothesis, we have previously shown after intraluminal MCAO in mice that the delivery of antibodies inhibiting the integrin VLA-4, which mediates the entry of T cells and PMNs into the brain, did not show any additional protective effects when PMNs had been depleted.⁴ The depletion of PMNs alone provided maximum protection. However, anti-VLA-4 acted synergistically with a selective T cell-depleting antibody to provide maximum protection.⁴ Hence, in this model, PMNs appeared to be a master switch that was responsible for the injury-promoting capacity of both PMNs and T cells. The combined evidence of the now presented study and our earlier study suggests that PMNs might act as an amplifier of detrimental immune responses that

compromise stroke outcome. Further studies on this issue are warranted.

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