Erythropoietin attenuates the brain edema response following experimental traumatic brain injury

Jonas Blixt<sub>ab</sub>, Eli Gunnarson<sub>cd</sub>, Michael Wanecek<sub>bd</sub>

<sub>a</sub>Perioperative Medicine and Intensive care, Karolinska University Hospital;

<sub>b</sub>Department of Physiology and Pharmacology;

<sub>c</sub>Department of Women’s and Children’s Health;

<sub>d</sub>all: Karolinska Institutet, SE-171 77, Stockholm, Sweden;

<sub>b</sub>Shared last authorship

Jonas.blixt@sll.se
Eli.gunnarson@ki.se
Michael.wanecek@ki.se

Corresponding author: Jonas Blixt, Perioperative Medicine and Intensive care, Karolinska University Hospital, SE-171 76 Stockholm, Sweden. Tel: +46 (0)707398 598, +46 (0)8 517 791 87. E-mail address: jonas.blixt@sll.se

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ABSTRACT

Erythropoietin (EPO) has neuroprotective effects in multiple CNS injury models; however, EPO’s effects on traumatic brain edema are elusive. To explore EPO as an intervention in traumatic brain edema, male SD-rats were subjected to blunt, controlled traumatic brain injury (TBI). Animals were randomized to EPO 5000IU/kg or saline (control group) intraperitoneally within 30 minutes after trauma and once daily for 4 consecutive days. Brain MRI, immunohistofluorescence, immunohistochemistry and quantitative protein analysis were performed at day 1 and 4 post trauma. EPO significantly prevented the loss of the tight junction protein zona occludens 1 (ZO-1) observed in control animals after trauma. The decrease of ZO-1 in the control group was associated with an IgG increase in the perilesional parenchyma, indicating BBB dysfunction and increased permeability. EPO treatment attenuated decrease in apparent diffusion coefficient (ADC) after trauma suggesting a reduction of cytotoxic edema, and reduced the IgG leakage indicating that EPO contributed to preserve BBB integrity and attenuated vasogenic edema. Animals treated with EPO demonstrated conserved levels of aquaporin 4 (AQP4) protein expression in the perilesional area while control animals showed a reduction of AQP4. We show that post TBI administration of EPO decreases early cytotoxic brain edema and preserves structural and functional properties of the BBB, leading to attenuation of the vasogenic edema response. The data support that the mechanisms involve preservation of the tight junction protein ZO-1 and the water channel AQP4 and indicate that treatment with EPO may have beneficial effects on the brain edema response following TBI.
INTRODUCTION

Traumatic brain injury (TBI) is a major cause of early death and neurological disability throughout the world and remains a major public health problem.\(^1,2\) Injury mechanisms in brain trauma include several cascades of events such as release of excitotoxic molecules, metabolic challenges, inflammation and apoptosis, accompanied by breakdown of the blood-brain barrier (BBB) and cell swelling, which can finally lead to cell death. Despite several recent clinical studies based on promising experimental evidence, targeting injury mechanisms, there are currently no available pharmacological treatments that convincingly counteracts secondary injuries and/or improve long term functional recovery after human TBI.\(^3-9\)

Erythropoietin (EPO) is a glycoprotein cytokine that has been investigated in TBI research. EPO is produced mainly in the kidney but also locally in brain cells under hypoxic conditions.\(^10,11\) EPO receptors have been found throughout the brain in various cell types such as neurons, astrocytes and endothelial cells.\(^12-14\) In the literature there is a substantial body of evidence in support of non-hematopoietic, neuroprotective effects of EPO\(^11,15\) in models of CNS injury such as ischemia, trauma, intracranial hemorrhages and diffuse axonal injuries. The proposed mechanisms include decreased inflammatory response,\(^16-18\) decrease of oxidative stress and anti-apoptotic effects.\(^16,19,20\)

The effects of EPO treatment in clinical CNS lesions/injuries have varied. A recent randomized, prospective, placebo-controlled clinical trial in ischemic stroke showed that EPO significantly improved long term neurological outcome.\(^21\) Following moderate or severe TBI results on EPO treatment have been conflicting. One study found decreased mortality and improved neurological outcome\(^22\) while the recent study by Nichol et al did not find that EPO resulted in significant improvement of functional outcome when assessed 6 months after TBI.\(^23\)
The clinical course that follows severe TBI is characterised by brain edema development and an associated increase in intracranial pressure contributing to most of the early deaths following trauma. Using a traumatic brain injury model we have previously shown that the dynamic course of edema development include co-existing vasogenic and cytotoxic brain edema. The aim of the present study was to investigate whether EPO treatment can modulate the dynamic brain edema response following experimental focal brain injury.

**MATERIALS AND METHODS**

**Animals**

Animal care and experimental procedures were conducted in accordance with European Communities Council Directive of 24 Nov. 1986 (86/609/EEC). Experimental protocols were approved by the Northern Stockholm Laboratory Animal Review Board. All studies were performed on male Sprague-Dawley rats (Scanbur, Sweden) 250-280 g, housed at the Karolinska University Hospital (Stockholm, Sweden) in polystyrene cages containing aspen wood shavings and with free access to water and standard rodent chow with regulated 12-h light–dark cycles. All procedures and housing throughout the study were at room temperature (21ºC±1ºC) with normal air humidity.

**Experimental groups**

A total of 75 male rats were used in this study and were randomized by drawing numbers. They were divided into a naive group with animals without TBI, a control group that received NaCl after trauma, and a treatment group that received EPO after trauma. The naive group (n=15) was further divided in groups of five to be used in different experiments. The NaCl or EPO groups (n=30/group), were further randomized to be sacrificed after 1 day or 4 days (n=5/group/day) in three different experimental settings.
Drug and dose

EPO (recombinant human Erythropoietin, Eprex, 4000 IU/ml, Janssen-Cilag, France) was administered intraperitoneally 30 minutes after trauma and thereafter once daily in a dose of 5000 IU/kg throughout the study. Animals in the control group received an identical volume of saline (isotonic NaCl 0.9%, Fresenius Kabi, Sweden) intraperitoneally.

Traumatic brain injury model

After randomization to either naive or TBI, the TBI rats were anesthetized by intramuscular injections of 0.15 ml of Hypnorm® (10 mg/ml fluanisone and 0.315 mg/ml fentanyl citrate; Janssen Pharmaceutical, Beerse, Belgium) and 0.2 ml Dormicum® (midazolam, 1 mg/ml, F. Hoffmann-La Roche, Basel, Switzerland). Before skin incision 0.1 ml of Marcin-Adrenalin® (bupivacaine 5 mg/ml and adrenaline 5 µg/ml, Astra Zeneca, Södertälje, Sweden) was injected subcutaneously in the sagittal midline of the skull. The rats were placed in a stereotactic frame. Eye-gel (Fucithalmic®) applied for eye protection and isotonic saline (NaCl 9mg/ml) was used to clean and rinse the scalp wound throughout the experiment. Under aseptic conditions a craniotomy (2 mm in diameter) was drilled under microscopic guidance at a point 2-3 mm posterior and 2-2.5 mm lateral to the right side of bregma. A standardized parietal contusion was produced by a weight-drop model after Feeny that has been further developed in our lab. A weight of 24 g was dropped from a height of 7 cm on a 1.8 mm diameter piston that was set to allow for a maximal tissue compression of 3 mm. The craniotomy was not resealed. Spontaneous ventilation was monitored by pulse oximetry (Datex-Ohmeda, Helsinki, Finland) and oxygen was supplied through a facemask to keep oxygen saturation above 90%. The body temperature was kept at 37.5-38°C by means of a feedback regulated heating pad. Heart rate and respiratory rate was continuously observed and within physiological range throughout the experimental period (Table 1). The skin was sutured using Vicryl 4-0 (Ethicon, Johnson & Johnson, New Brunswick, USA) and the animals were allowed to recover under at heating lamp with extra supply of oxygen. The TBI animals showed no deficit in motor, balance or sensory function during the posttraumatic phase and they showed normal grooming and feeding behaviour.
Euthanasia

Animals intended for immunohistochemistry were anaesthetized and sacrificed by transcardial perfusion. All other animals were anaesthetized as described above and then immediately sacrificed by decapitation.

Magnetic Resonance Imaging

MRI of the brain (Naive n=5, Control/EPO 1 and 4 days n=5/day/group) was performed at 1 day and 4 days post-injury. Animals were re-anaesthetized with the same doses of Midazolam and Hypnorm. After intubation with a 16-gauge intravenous catheter (45mm length, 20mm hub, 1.7mm ID) anaesthesia was maintained with 1.5-2% isoflurane and animals mechanically ventilated with a small animal ventilator SAR-830P (Stoelting Co, Chicago, USA). The ventilator was set at a tidal volume of 8ml/kg and a respiratory rate of 40 breaths/min. Rectal temperature was maintained between 36.5°C and 37.5°C by using a feedback-regulated heating pad. The eyes were lubricated and covered to prevent dehydration. MRI was performed with a Bruker Biospec 4.7 T scanner (Bruker, Karlsruhe, Germany) fitted with a 12cm inner diameter self-shielded gradient system (max. gradient strength 200 mT/m). A 38 mm diameter head coil birdcage was used for imaging.

Qualitative T2-weighted images were obtained using a rapid acquisition with relaxation enhancement (RARE) (TR = 2,500 ms; effective echo time = 30 ms; RARE factor = 8; FOV = 3.5 × 3.5 cm; matrix 256 × 256) five contiguous slices of 1 mm, two averages.

At the centre of the lesion a 1.5mm thick diffusion-weighted image was taken. In order to reach an acceptable signal-to-noise ratio without dramatically increasing the acquisition time, a 128 × 64 matrix for FOV = 3.5 × 3.5 cm was used. Diffusion images were obtained using a pulsed gradient stimulated echo sequence with six diffusion weighting factor b = 39.049, 93.557, 274.708, 401.350, 665.853, 2391.773 s/mm². The diffusion-sensitizing gradient was placed only along the direction of the slice selection gradient (i.e. rostro-caudally).
MRI - Mid line shift

Midline shifts (MLS) were determined on T2-weighted images with the use of analysis software ImageJ® 1.48v (Rasband, W.S., ImageJ, U. S. National Institutes of Health, Bethesda, Maryland, USA). After optical adjustment of contrast and brightness, the distance between the outer border of the cortex and the middle of the third ventricle was measured from the ipsilateral (I) and contralateral (C) sides. MLS was calculated using the equation MLS = (I-C)/2.

MRI - Edema analysis

ADC maps were calculated by ImageJ and the use of MRI Analysis Calculator, a plugin provided for free by Karl Schmidt. Further analysis was performed in region of interest (ROI) with ROIs being placed in the perilesional cortex area, and at the same site in the contralateral hemisphere. Assessments of both MLS and ADC were performed blinded by one investigator and calculated as mean of three repeated measurements. Repeated analysis with more than two weeks separation did not show any statistical significant difference.

Immunohistochemistry - IgG staining protocol

Transcardial perfusion was performed (Naive n=5, Control/EPO 1 and 4 days n=5/day/group) with cooled saline followed by cold 4% paraformaldehyde (PFA) in 0,15M phosphate buffer (pH 7.3-7.4). Brains were removed and post-fixated for 90 minutes in the same fixative followed by wash over night in 0,15M phosphate buffer with 17% (w/v) sucrose at 4°C, thereafter snap frozen in isopentane and stored at −70°C. Whole brain sections 14µm were cut horizontally through the length of the contusion using a Leica cryostat and then thaw mounted onto Super Frost/Plus object glasses, dried at room temperature for 1-2 hours and stored at −20°C prior to use.
The sections were incubated in 0.3% hydrogen peroxide for 30 min to quench endogenous peroxidase. Before incubation with primary antibody, blocking serum was used to prevent non-specific conjugate binding. An avidin-biotin blocking step was then performed with Vectastain Elite avidin-biotinylated enzyme complex (ABC) peroxidase kit (Vector Laboratories, Burlingame, CA, USA) to prevent non-specific conjugate binding to endogenous biotinylated proteins. Sections were then incubated with primary anti-IgG antibody overnight at +4°C. In negative controls (data not shown) slides were incubated with PBS. After washing, the indirect peroxidase method was used for detection of the primary antibodies. Biotin-conjugated donkey anti-goat immunoglobulin was used as conjugate. Sections were incubated with a standard ABC kit and the bound peroxidase was detected via incubation with a DAB substrate kit. Sections were then dehydrated and mounted with DPX (Distrene 80, di-butyl phthalate, xylene, BDH Laboratory Supplies Pool, UK).

**Immunohistochemistry - IgG Analysis**

Stained sections were scanned with a Nikon Super Coolscan 4000 ED (32bit, 4000dpi, sRGB) and saved as TIF images. After adjustment of contrast and brightness in ImageJ, parameters then remained unchanged through the series and the pictures were converted to grayscale. Three ROIs of the same size were placed in representative areas in both hemispheres in three consecutive sections per animal, and the mean staining intensity for each animal was calculated. We chose to measure the monochromatic intensity of the IgG signal rather than the area, as the dissemination of IgG could be affected unevenly by local factors, e.g. edema. We analyzed the contralateral side in all sections, both within groups, between groups and against sections from naive animals without any statistical differences in staining intensity. To diminish potential effects of small differences in staining, corresponding areas in the contralateral hemisphere were used for signal normalisation, and the results were presented as ipsilateral /contralateral values (“ipsi/contra”).
Immunohistofluorescence - Staining protocol

The brains for immunohistofluorescence analysis (Naive n=5, Control/EPO 1 and 4 days n=5/day/group) were quickly removed, snap frozen in isopentane and stored in −70°C. Whole brain 14 μm cryosections were cut coronally through the centre of the impact using a Leica cryostat (CM 3000, Leica Instruments GmbH, Nussloch, Germany). The sections were thaw mounted onto Super Frost/Plus object glasses (Menzel–Gläzer, Braunschweig, Germany) dried at room temperature for 1-2 hours and stored at −20°C prior to analysis.

Prior to staining the sections were dried at room temperature for 30 minutes, rehydrated in PBS and fixed in 4% buffered paraformaldehyde at room temperature for 10 min and then rinsed in PBS. Treatment for 60 min with blocking serum (PBS, 1% bovine serum albumin, 0.3% triton X-100) was used to prevent non-specific conjugate binding to the primary antibodies. Primary antibodies (anti-ZO-1 or anti-AQP4) were diluted in blocking serum and added for incubation overnight at +4 °C either in solitude or in mixtures. Sections were washed in PBS, incubated for 1 hour with secondary antibodies, washed again and cover-slipped with ProLong Gold anti-fade reagent with DAPI (Invitrogen Life Technologies, Carlsbad, CA, USA). Negative control staining with omitted primary antibody, or with depletion of the AQP4 primary antibody by an excess of the specific peptide (Chemicon International, Temecula, CA, USA), did not result in any detectable labelling.

Immunohistofluorescence - Analysis of AQP4 and ZO-1

Fluorescent microscope images were acquired on a Vslide® slide scanning microscope (MetaSystems, Altlussheim, Germany). Whole microscope slides were scanned at x2.5 to adjust focus and to generate a tissue map. Focus and tissue depth were detected based on the DAPI signal. Upper half of brains were scanned using x10 objective. Images were stitched together to generate large 3-channel fluorescence images with microscopic resolution. Acquired images were extracted as TIFF-files using the software Metaviewer® (MetaSystems, Altlussheim, Germany).
Immunohistofluorescence - Quantification of AQP4 and ZO-1

ImageJ was used to analyse antibody labelling in the cortical area. Three quadrant ROIs of 550µm width were manually placed in the medial part of the perilesional area in the cortex, and in the corresponding area on the contralateral side. The thresholds were manually set to each individual antibody but were then kept fixed throughout the experiment. The staining was automatically quantified with batch processing, using a by us created ImageJ macro with thresholding and background subtraction. Three ROIs per section and three sections per animal were used to generate a mean value per animal. All sections for each experiment were stained at the same time to minimize differences between different staining sessions. Contralateral hemisphere was used for signal normalization and the results presented as ipsilateral/contralateral values (“ipsi/contra”). In control experiments, multiple localizations were tested to evaluate that the selected ROIs in both hemispheres were representative and that the areas in the ipsilateral side represented viable tissue. Selected areas were analysed in higher magnification (60x) to ensure that the tissue morphology appeared intact according to GFAP-positive astrocytes, number of nuclei, and the density of blood vessels (data not shown). Any increase of GFAP immunoreactivity within the ROI was taken as an indication that astrocytes in the area were viable. To ascertain that control areas in the contralateral side did not differ from corresponding areas in naive animals the values of ADC, AQP4, ZO1, GFAP and IgG were analysed and found not to differ.

STATISTICAL ANALYSIS

Evaluation of data was performed by an observer blinded for the randomization data. Intraobserver variability was calculated after three measurements, each separated by two weeks for the first 5 measurements in all settings. No statistical differences between the observations were found, with a variability of 3-7% depending on type of measurement. Statistical analysis was performed using two-sided Student’s t-test. All data were normally distributed. P-values <0.05 was considered statistically significant. Calculations were performed using SPSS 20 (IBM SPSS Statistics, Armonk, NY) and graphs were made using Prism 6.02 (GraphPad Software, La Jolla, CA, USA). Data are expressed as mean ±SD.
RESULTS

All animals presented normal behaviour without apparent neurological deficits after the brain trauma. Physiological parameters (body weight, temperature, saturation, heart rate and respiratory rate) were within the physiological range throughout the experiment (Table 1).

Focal brain contusion resulted in brain edema evident in T2 MRI sections

Evaluation of time course of brain swelling and type of edema development following TBI were assessed by analysing MRI images with regard to mid line shift (MLS) and ADC values. ADC values were measured in the perilesional area lateral to the traumatic brain lesion, and in the corresponding area in the contralateral, uninjured, cortex. Brain MRI images revealed a focal cortical contusion in the right hemisphere of similar magnitude in all animals exposed to TBI. T2 weighted MRI images showed hyper-density in the perilesional area of the focal injury, indicating presence of brain edema. No T2 changes were evident in other brain areas in either the ipsi- or contralateral hemisphere (Fig. 1A).

EPO treatment resulted in less ADC decrease following TBI

ADC values are presented as relative change (in percent) of ipsilateral/contralateral values under the assumption that ADC in naive, uninjured, animals is equal in both hemispheres. TBI in the control group resulted in an ADC decrease in the perilesional area by 12% and 15% at day 1 and 4, respectively. EPO treatment prevented the ADC decrease seen in the control group and did not differ significantly compared to naive animals at either time point (Fig. 1B). There were no significant changes of ADC in the corresponding areas on the contralateral side at any time.
**EPO did not alter the Midline Shift**

The focal TBI caused a significant MLS (Fig. 1C) of the ipsilateral brain hemisphere towards the non-traumatized hemisphere by approximately 8% both at day 1 and day 4 after trauma in the control group. MLS was not significantly affected by EPO treatment (Fig. 1D).

**EPO improved BBB morphology**

BBB morphology was assessed by analysing the amount of the endothelial tight junction marker ZO-1 in the perilesional area. ZO-1 was decreased by more than 25% at day 1 and day 4 in control animals. EPO significantly reduced the loss of ZO-1, as animals treated with EPO only had an 8% decrease at day 1 and showed a 10% net increase of ZO-1 at day 4 after trauma (Fig. 2).

**EPO improved blood brain barrier integrity**

BBB permeability was assessed by quantifying the presence of IgG in the perilesional area. Focal TBI in control animals resulted in an increase of IgG by 23% and 21% at day 1 and 4, respectively. There was no significant change in IgG abundance at day 1 in animals that received EPO, however EPO treatment significantly decreased IgG in the perilesional area to 11% at day 4 post injury (Fig. 3).

**EPO attenuated the trauma induced decrease of AQP4**

In control animals focal TBI caused a significant decrease of AQP4 protein expression in the perilesional area, as analysed by fluorescence intensity in ipsilateral/contralateral hemisphere and compared to naive animals. AQP4 protein was decreased by 17% and 14% at day 1 and day 4, respectively. In EPO treated animals the expression of AQP4 did not decrease and was similar to naive animals (Fig. 4). AQP4 was present in perivascular end-feet surrounding vasculature as confirmed with the endothelial marker RECA-1 (data not...
shown). The polarized distribution of AQP4 in predominantly perivascular areas was not overtly changed in either TBI controls or EPO-treated animals as compared to naive animals.

DISCUSSION

In this experimental TBI model we show that daily administration of EPO post injury resulted in improved structure and function of the BBB, preserved AQP4 protein expression in the perilesional area and MRI changes consistent with decreased cytotoxic brain edema. Mechanisms proposed to be involved in the neuro-protective properties of EPO are mostly attributed to direct effects on neurons and are expected to be mediated via binding to an alternative EPO-receptor, the IRR-receptor (innate repair receptor). This receptor is typically not expressed in normal tissue but upregulated under hypoxic and ischemic conditions as can be present in brain injury. The IRR-receptor activates the Janus-tyrosine-kinase-2 (JAK-2) leading to several non-hemapoetic downstream signaling pathways. Alterations in nitric oxide (NO) production by endothelial nitric oxide synthase (eNOS) and enhanced NO production have also been shown to be protective routes for EPO in animal models such as heart ischemia-reperfusion, TBI and intracerebral hemorrhages.

Several different mechanisms are suggested to be involved in edema development after traumatic brain injury. According to the traditional view there is a primary and immediate opening of the BBB after TBI that peaks at 4-6h and then resolves within 6-7 days after trauma in humans. This opening is considered to be caused by the direct physical trauma, disruption of the BBB, alternations in the expression of AQP4, complement activation, dysregulation of matrix metalloproteinases and release of cytokines.

In a previous study we showed that the current focal TBI model causes BBB dysfunction as indicated by structural (loss of ZO-1) and functional changes (increased IgG) of the BBB. The findings support a development of vasogenic brain edema as a result of disrupted BBB integrity.
In the present study we find that administration of EPO after TBI and once daily preserves ZO-1 levels. The effect was observed as early as day 1 after trauma and persisted at day 4 with a net increase in ZO-1 compared to controls. At day 4, there was a markedly decreased IgG extravasation in the perilesional area in EPO-treated animals. Taken together, the present findings indicate that EPO attenuates trauma-induced BBB dysfunction and thus decreases vasogenic brain edema. ZO-1 is localized at tight junction (TJ) sites and interacts directly with a majority of the transmembrane TJ proteins. In accordance with our findings, dissociation of ZO-1 from the TJ complex has been shown to increase permeability of the BBB. Further, epithelial cells with ZO-1 deficiency do not form TJs due to lack of claudin polymerization and establishment of the barrier function is therefore delayed. In TBI, neuroinflammatory mediators such as metalloproteinase-9 cleave extracellular matrix including TJ-associated proteins such as ZO-1. This induces alterations of BBB integrity leading to changes in TJ architecture, allowing for the development of vasogenic edema. Our findings thus indicate that treatment with EPO after TBI may preserve BBB function and reduce vasogenic edema by attenuating loss of ZO-1. Such an effect of EPO on BBB disruption in focal TBI has previously not been well characterised. In a weight drop TBI model EPO was found to decrease permeability to Evans blue, but BBB integrity or morphology was not evaluated. EPO treatment has however been reported to preserve BBB function in other, non-TBI, brain injury models such as cryogenic injury, MCA occlusion and diffuse axonal injury with hypoxia. The temporal changes in our study show that ZO-1 levels are normalized by day 1 after trauma, while the effect on IgG leakage is evident at day 4. We speculate that the focal TBI cause an immediate increase in BBB permeability and that EPO treatment, due to preserved or early restoration of ZO-1 levels, attenuates a continued IgG leakage as shown by the significantly reduced parenchymal IgG at day 4 after injury. Small amounts of immunoglobulins, including IgG, may be present in normal CNS perivascular tissue and it has been shown that active transportation of IgG over BBB may occur. However, a significant increase in IgG in brain tissue indicates vasogenic edema as a result of BBB dysfunction.
ADC mapping is recognised to differ between relative changes of the two entities of brain edema; cytotoxic edema which is associated with low ADC values and vasogenic edema, which is associated with high ADC values. Here we found that EPO treated TBI animals showed ADC values similar to naive animals with no injury. In contrast, control animals subjected to TBI showed a decrease in ADC at both day 1 and 4 post injury, findings in consistency with our previous study.\(^{24}\) The ADC decrease indicates a presence of cytotoxic edema in the area surrounding the focal injury. EPO treatment thus seems to oppose the development of cytotoxic edema. This effect was observed as early as day 1 and was sustained to day 4 after trauma.

Astrocyte foot processes, with their abundance of AQP4, are susceptible to cytotoxic edema formation. It has been shown that AQP4 is involved in the development of cerebral cytotoxic edema in various pathological conditions. In accordance, AQP4-deficient mice are less prone to develop cytotoxic edema in experimental ischemia and bacterial meningitis.\(^ {56, 57}\) However, the consequences of AQP4 deficiency vary depending on the type of brain edema studied, as AQP4-null mice are more prone to develop vasogenic edema.\(^ {58}\) In studies of traumatic brain injury models there have been reports of both increased\(^{59-61}\) and reduced\(^ {41, 62}\) AQP4 protein expression.

In the present model of focal TBI we found a decrease in AQP4 protein expression in the perilesional area where the gross and microscopic anatomy otherwise appeared unaffected (present study and\(^ {24}\)). The quantification of AQP4 in the perilesional area was performed using immunohistofluorescence, which allowed us to assess the distribution of AQP4 in the same sections. Interestingly, treatment with EPO counteracted the trauma-induced decrease in AQP4 protein and, in fact, AQP4 levels were indistinguishable from naive animals without TBI. Moreover, both in control animals and EPO treated animals the polarized localization of AQP4 in astrocyte end-feet facing cerebral blood vessels appeared unaffected.

We interpret the absence of ADC decrease after EPO treatment as a decrease in cytotoxic edema. One possible mechanism for the reduced cytotoxic edema may be increased water clearance due to the preserved protein levels of AQP4 in EPO treated animals.\(^ {63}\) Other
mechanisms may include downregulation of AQP4 water permeability by EPO and/or improved cellular ionic and metabolic functions, counteracting water accumulation.

We cannot rule out that the attenuation of ADC decrease induced by EPO represents an increase in vasogenic edema, however we find this unlikely as the BBB integrity was concomitantly improved by EPO as shown by a significant decrease in IgG leakage.

Since the craniotomy was not resealed the current injury model may be considered to resemble a penetrating brain injury leading to a brain contusion. Previous studies has ascertained that this injury model results in edema development and hemisphere enlargement but no brain protrusion, which may be due to the small size of the craniotomy (<5mm²) as compared to e.g. Dixons original work (estimated area 80-100mm²). Pre hoc we decided to focus on local effects and evaluation of edema types in the pericontusional area in this study, and decided not to analyze total brain water accumulation or contusion volume. To ensure that the contusions were homogenous and comparable, we compared AQP4, ZO-1, ADC and IgG values within the same pericontusional areas in the control animals and found very small deviations.

The focal cortical contusion and edema resulted in a significant mid line shift of the brain towards the uninjured side. This shift was not significantly affected by EPO treatment. MLS is a crude estimate of brain swelling and is not considered a sensitive marker of the underlying cause of brain swelling – hyperemia, vasogenic edema and/or cytotoxic edema. Despite this crudeness MLS is thought to be a relevant prognostic tool in clinical practice. The lack of statistically significant effect on MLS by EPO treatment here may reflect that this injury estimate has a low sensitivity for changes in small-volume injuries, relatively small sample sizes or the fact that MLS does not identify shifts of volume between different tissue compartments.

EPO has been shown to be beneficial in multiple experimental models of neurological injury. In a recent study, Operation Brain Trauma Therapy (OBTT) consortium tested EPO in three different TBI models in rat, including a focal cortical injury model. In the study a single dose of EPO was administered directly after trauma. The investigators did not find any effect on either histopathological, cognitive or biomarker parameters. In the present
study of focal TBI, we administered multiple post-injury doses of EPO and observed improved BBB function and an effect on the dynamic edema response following trauma. We did not estimate whether injury volume or neurocognitive function was improved by this dosing regimen of EPO, however this will be interesting for future studies.

In clinical trials of CNS insults the results of EPO treatment have been diverse. EPO significantly improved long term neurological outcome after ischemic stroke in recent study by Tsai et al,\textsuperscript{21} while results have been inconclusive in subarachnoid haemorrhage.\textsuperscript{71} Following moderate or severe traumatic brain injury, Nichol and coworkers could not show that EPO reduced the number of patients with severe neurological dysfunction on outcome scales 6 months after trauma.\textsuperscript{23} Although not significant, there was an interesting trend towards a favorable outcome with regard to mortality in the EPO treatment group.

In a smaller study, using a different EPO dosing regime, Aloizos et al found that EPO decreased mortality and improved neurological outcome at 6 months post severe TBI.\textsuperscript{22}

Limitations in the present study include relatively low sample sizes, which may have affected the negative MLS results, and a limited time resolution, which might conceal details in the sequence of events and fluctuations in the estimated parameters in between the analyzed time points. The study does not explore intracellular signaling pathways involved in the effects of EPO on the BBB and brain edema.

CONCLUSION

In this experimental traumatic brain injury model we show that post traumatic administration of daily doses of EPO preserved both structural and functional properties of the BBB by preventing loss of the tight junction component ZO-1 and reducing IgG permeability. Furthermore, EPO treatment inhibited the ADC decrease following TBI, indicating that EPO reduced traumatic cytotoxic edema. A summary of the data is presented in Fig 5. In conclusion, the findings in this study support that EPO may have
positive effects on traumatic brain edema. Further examination of both dose and timing of EPO administration is warranted.

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AUTHOR DISCLOSURE STATEMENT

No competing financial interests exist for any author
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Abbreviations

ADC, apparent diffusion coefficient; AQP, aquaporin; AQP4, aquaporin-4; BBB, blood-brain barrier; C, contralateral; EPO, Erythropoietin; GFAP, Glial fibrillary acidic protein; I, ipsilateral; IF, immunofluorescence; IgG, immunoglobulin G; MLS, midline shift; MRI, magnetic resonance imaging; TBI, traumatic brain injury; ZO-1, zona occludens 1.
### Table 1. Physiological data during operation

| Group          | Naive     | Control   | EPO       |
|----------------|-----------|-----------|-----------|
| Heart Rate     | 351±11    | 359±10    | 360±7     |
| Respiratory Rate| 80±2      | 85±3      | 83±5      |
| Saturation     | 94±3      | 95±1      | 94±2      |
| Temperature (°C)| 37.5-38   | 37.5-38   | 37.5-38   |
| Body Weight (g) | 258±5     | 252±4     | 255±3     |
Figure legends

**Fig. 1.** (A) Representative T2 weighted coronal MRI section of adult rat brain at day 4 after traumatic brain injury (contusion) in the right hemisphere. The focal injury in the ipsilateral hemisphere (I) is surrounded by a hyper-dense area, indicating edema. No hyper-dense MRI changes are observed in the contralateral hemisphere (C). (B) Relative changes of ADC in the perilesional region at day 1 and 4 in control animals and EPO-treated animals (EPO); ipsilateral/contralateral (I/C) values in percent. ADC I/C was significantly decreased at both day 1 and day 4 after trauma in controls. EPO treatment attenuated the ADC decrease at both time points. Resultant ADC values were similar to ADC in naive animals without trauma.

(C) Schematic illustration of a coronal section of rat brain. Midline shift (MLS) was estimated in percent using the distance between the outer border of the cortex and the middle of the third ventricle in the ipsilateral and contralateral hemispheres, respectively. The dark grey area illustrates the contusion and the light grey quadrants depict the perilesional regions analyzed in all other experiments. (D) MLS at day 1 and 4 after trauma in control animals and EPO-treated animals. A significant MLS towards the uninjured side was present at both time points in both groups; i.e. EPO treatment did not affect MLS. Values presented as mean ± SD. (MLS, mid line shift; ADC, apparent diffusion coefficient; I/C, ipsilateral/contralateral; d, days after trauma; # p<0.05)
Fig. 2. (A) High magnification confocal immunofluorescence image of ZO-1 (red), the glial cell marker GFAP (green) and the nucleus marker DAPI (blue) from brain area with blood vessels (longitudinal view of blood vessel to the left), to illustrate the specificity and localization of the antibody binding. (B) Monochromatic presentation of GFAP immunofluorescence. (C) Monochromatic presentation of ZO-1 immunofluorescence. (D) Summary data on ZO-1 immunofluorescence in the traumatized, ipsilateral, hemisphere normalized against the contralateral side at day 1 and 4 after trauma in control and EPO-treated animals. ZO-1 was significantly reduced after trauma in the control group. EPO prevented the loss of ZO-1 at both time points after trauma. Values presented as mean ± SD. (ZO-1, Zona Occludens 1; GFAP, glial fibrillary acidic protein; DAPI, 4',6-diamidino-2-phenylindole; #p<0.05, ##p<0.01 compared to naive animals; **p<0.01, ***p<0.001 between groups)
**Fig. 3.** IgG immunohistochemistry on representative axial brain sections. (A) Control animals day 1 and 4 post trauma, (B) EPO treated animal day 1 and day 4 post trauma, (C) Naive animal without trauma. IgG positive immunoreactivity (dark area) was observed around the site of the trauma in the right hemisphere. (D) Relative changes (in percent) of IgG immunoreactivity at day 1 and 4 after trauma and compared to naive (base line). The perilesional area is normalized against the contralateral side. IgG was significantly increased in control animals. EPO treatment significantly decreased IgG in the perilesional area at day 4 after trauma. Values presented as mean ± SD. (d, days post trauma; I/C, ipsilateral/contralateral; IgG, Immunoglobulin G; ###p<0.001 compared to naive; **p<0.01 between groups)
Fig. 4. (A) Representative monochromatic immunofluorescence image with AQP4 (white) on part of coronal brain section after blunt controlled parietal trauma (*) in the right hemisphere in control animal. The core of the injury shows disrupted microscopic structure. Measurements of AQP4 protein abundance were performed in the perilesional area (as shown in Fig. 1(C)) with visually intact structure. (B) High magnification of representative contralateral and (C) perilesional and areas. (D) Summary data on relative changes of AQP4 immunofluorescence intensity in the perilesional area at day 1 and 4 after trauma in control and EPO-treated animals. AQP4 was significantly reduced at both time points in control animals. EPO treatment normalized AQP4 protein levels in the perilesional area at both day 1 and 4 after trauma. Values presented as mean ± SD. (AQP4, aquaporin 4; I/C, ipsilateral/contralateral; d, days post trauma; ##p<0.01, ### p<0.001 compared to naive; **p<0.05, ***p<0.001 between groups)
Fig. 5. Schematic presentation of relative changes in the perilesional region at day 0, 1 and 4 after trauma in control animals (NaCl) and animals treated with EPO (EPO). Values are compared to naive animals without trauma (point zero) (IgG, ZO-1, AQP4), or to the assumption of no difference between the hemispheres (ADC). The arbitrary continuous time line is created from the separate measurements for each parameter at the indicated time points. (IgG, immunoglobulin G; AQP4, aquaporin 4; ADC, apparent diffusion coefficient; ZO-1, zona occludens 1; d, days after trauma)