Advances in MRI-Based Detection of Cerebrovascular Changes after Experimental Traumatic Brain Injury

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Abstract Traumatic brain injury is a heterogeneous and multifaceted neurological disorder that involves diverse pathophysiological pathways and mechanisms. Thorough characterization and monitoring of the brain’s status after neurotrauma is therefore highly complicated. Magnetic resonance imaging (MRI) provides a versatile tool for in vivo spatiotemporal assessment of various aspects of central nervous system injury, such as edema formation, perfusion disturbances and structural tissue damage. Moreover, recent advances in MRI methods that make use of contrast agents have opened up additional opportunities for measurement of events at the level of the cerebrovascular tree, such as blood–brain barrier permeability, leukocyte infiltration, cell adhesion molecule upregulation and vascular remodeling. It is becoming increasingly clear that these cerebrovascular alterations play a significant role in the progression of post-traumatic brain injury as well as in the process of post-traumatic brain repair. Application of advanced multiparametric MRI strategies in experimental, preclinical studies may significantly aid in the elucidation of pathomechanisms, monitoring of treatment effects, and identification of predictive markers after traumatic brain injury.

Keywords Animal models · Contrast agents · Magnetic resonance imaging · Neuroinflammation · Traumatic brain injury

Introduction

Magnetic resonance imaging (MRI) is a versatile modality for in vivo imaging of the central nervous system (CNS). The technique relies on detection of MR signals from the highly abundant water protons in tissues. Its versatility stems from the ability to sensitize MRI acquisitions to different parameters, such as proton density, MR relaxation times (T_1, T_2), susceptibility contrast, diffusion, perfusion and flow. This enables multiparametric imaging studies on different pathophysiological events that are involved in the evolution of CNS injury, e.g., after neurotrauma. Furthermore, the availability of MRI in hospital and laboratory settings facilitates the translation between clinical and preclinical research. Hence, MRI studies in experimental models of CNS injury can contribute to the characterization of pathomechanisms and treatment effects, which may lead to improvements in diagnosis and therapeutic strategies after CNS injury.

MRI of Tissue and Perfusion Status

MRI in animal models of traumatic brain injury and other CNS diseases most often involve T_2*- diffusion- and/or perfusion-weighted imaging to measure changes in brain tissue and perfusion status (for a review, see Dijkhuizen and Nicolay [1]). The transverse relaxation time of the MR signal, T_2, is dependent on the interstitial water content and increases when intercellular or vasogenic edema develops. This has been demonstrated with T_2-weighted MRI in rats in the first days after controlled cortical impact injury [2–5], closed-head injury [6, 7] and lateral fluid percussion injury [8]. While T_2-weighted MRI serves as a reliable tool to determine the extent and location of CNS lesions at
subacute and chronic stages, earlier tissue changes may be detected with diffusion-weighted MRI. Reduction of the apparent diffusion coefficient (ADC) or diffusivity of tissue water, most likely reflective of cytotoxic cell swelling [9], has been detected within hours after experimental traumatic brain injury, which precedes severe tissue damage, e.g., loss of cellularity, and resultant ADC elevations at later stages [6–8, 10–12]. Furthermore, with diffusion tensor imaging [13], it has been shown that changes in diffusion anisotropy and axial diffusivity in gray and white matter can inform on the extent of pericontusional and diffuse axonal injury after controlled cortical impact in mice [14] and head acceleration in rats [15], respectively. These studies reported permanent loss of diffusion anisotropy and initial decline, followed by rise of diffusivity, which correlated with axonal degeneration and demyelination.

The combination of T2- and/or diffusion-weighted MRI with perfusion-weighted MRI allows concurrent assessment of tissue lesions and perfusion alterations. Two perfusion-weighted MRI strategies that are frequently applied are dynamic susceptibility contrast (DSC)-enhanced MRI and arterial spin labeling (ASL) [16–18]. DSC MRI allows calculation of various hemodynamic parameters from MR signal changes caused by first passage of an intravascular bolus of contrast agent. ASL is based on spatially selective inversion or saturation of incoming arterial spins and subsequent detection of labeled blood water in perfused tissue, which enables quantification of cerebral blood flow. Both methods have been applied in rodent models of traumatic brain injury, and have provided insights into the development of focal as well as global post-traumatic cerebral hypoperfusion [7, 11, 19–22]. Figure 1 demonstrates how multiparametric MRI can inform on the spatiotemporal profile of tissue and perfusion changes after experimental traumatic brain injury. T2, diffusivity and cerebral blood flow (CBF) maps, acquired in studies by Immonen et al. [8] and Hayward et al. [20, 23], reveal first signs of secondary injury which slowly evolves from the primary contusion site at a subacute stage after lateral fluid percussion injury in rats, eventually resulting in an enlarged lesion at chronic stages.

**Post-traumatic Cerebrovascular Changes**

Abovementioned conventional MRI strategies allow depiction of edema formation and perfusion disturbances, which can be useful for diagnosis and treatment evaluation after traumatic brain injury. Nevertheless, the pathophysiology of neurotrauma is highly heterogeneous and complex, and involves various primary and secondary neuropathological pathways [24, 25]. In particular, increasing evidence points towards a major role for inflammatory events in the development of secondary tissue damage following traumatic brain injury, while inflammation has also been implicated in post-traumatic regenerative processes [26–29]. Associated cerebrovascular changes, such as blood–brain barrier (BBB) breakdown, upregulation of cell adhesion molecules, leukocyte infiltration and vascular remodeling, may strongly contribute to secondary hypoperfusion and edema formation after the primary insult, but might also play a part in post-traumatic tissue repair mechanisms [25, 27, 30, 31]. Elucidation of such pathways may contribute to improvement of assessment of injury severity, prediction of lesion progression and identification of targets for interventions after traumatic brain injury. The following paragraphs will briefly describe how specific contrast-enhanced MRI strategies may aid in gaining insights in abovementioned events in relation to neurotrauma.

**MRI of BBB Permeability**

Assessment of BBB integrity with MRI is based on detection of leakage of MR contrast agent from vascular into parenchymal space in CNS tissue [32, 33]. Intravenously administered paramagnetic contrast agent that leaks out and accumulates in brain parenchyma when microvascular barriers are disrupted, will result in local changes in MR signal intensity. The most frequently used contrast media are gadolinium-containing compounds, such as Gd-DTPA chelate, that locally shorten the longitudinal relaxation time, T1, of the MR signal. Contrast agents are usually injected as a bolus; however, step-down infusion may improve contrast enhancement through higher contrast agent concentrations in plasma and tissue [34]. Contrast-enhanced BBB permeability imaging usually involves a static approach, in which T1-weighted MRI is executed at a specific time-point (typically 10–30 min) after contrast agent injection, to detect hyperintense signal intensities generated by extravasated contrast agent. This is a fast, simple and sensitive method, but it merely provides qualitative information on the integrity of the BBB. Quantification of BBB leakage is feasible with dynamic contrast-enhanced (DCE) imaging, involving repetitive T1-weighted MR image acquisition, and analysis with a tracer kinetic model to assess contrast agent concentration changes in tissue and vasculature as a function of time [33]. This allows calculation of the blood-to-tissue volume transfer constant, Ktrans, as well as other parameters, such as the extravascular extracellular space volume. Analytical approaches that are frequently applied include Tofts’ compartmental analysis [35] and multiple time-graphic, or Patlak plot analysis [36]. Quantification of BBB permeability with these analysis methods, however, relies on certain physiological assumptions that may not
invariably hold under pathological conditions, for instance because of deviations in vascular delivery and cellular uptake of the tracer.

Serial BBB permeability MRI studies, particularly in cerebral ischemia–reperfusion models, have provided evidence for early and late BBB breakdown after experimental CNS injury [37–39]. However, the exact temporal profile of BBB permeability changes is still unclear, as studies include reports of persistent [39], as well as biphasic BBB opening [38]. Acute, transient BBB leakage, as detected with post-contrast T1-weighted MRI, has been shown after experimental traumatic brain injury, and may be prolonged by a post-traumatic secondary insult [40, 41]. In rabbits with weight-drop induced traumatic brain injury, serial DCE MRI revealed increasing $K_{\text{trans}}$ in the focal lesion area from 3 h to 3 days post-trauma [42] (Fig. 2). This was followed by a gradual decline, but $K_{\text{trans}}$ remained elevated up to 30 days as compared to animals with sham injury.

**MRI of Leukocytes**

Cellular MRI is based on the detection of cells that have been labeled with MR contrast agent, such as paramagnetic chelates, superparamagnetic particles or $^{19}$F MRI-detectable fluorinated particles (for reviews, see Modo et al. [43] and Stoll and Bendszus [44]). MRI-based visualization of various types of hepatobiliary, hematopoietic, immune and stem cells, has most frequently relied on labeling with superparamagnetic iron oxides [43]. These particles, generally ranging in size from a few nanometers to more than a micrometer, can induce very strong contrast effects by local shortening of the T$_2$ or T'$_2$, resulting in hypointense signal

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**Fig. 1** Maps of T$_2$ (a–c), D$_{av}$ (‘average diffusivity’ = 1/3 of the trace of the diffusion tensor) (d–f), and CBF (g–i) of a coronal brain slice from control rats (left column), and rats at 2–3 weeks (middle column) and 8 months after lateral fluid percussion injury (right column). Data are acquired from a quantitative MRI study by Immonen et al. [8] (a and d, b and e, and c and f are map pairs from the same animal, respectively) and CBF studies by Hayward et al. [20, 23], executed on a 4.7-T MR system with quadrature surface coil as radio-frequency transmitter/receiver (for T$_2$ and D$_{av}$ mapping), or a combination of an actively decoupled linear volume transmission coil and quadrature surface receiver (for CBF mapping with ASL). The parametric MRI maps demonstrate that at the cortical contusion site (*), a lesion/cyst was formed which slowly expanded (together with enlarging ventricles) up to several months post-injury. The perilesional cortical area (white arrows) and remaining tissue in the hippocampus ipsilateral to the contusion site (black arrows) show gradual increase in T$_2$ and diffusivity from 2 to 3 weeks post-injury onwards, indicative of progression of neurodegeneration (acute alterations in T$_2$ and diffusivity during the first week post-injury are edema-related and resolved by 2 weeks; data not shown). CBF was decreased in the perilesional area and in the ipsilateral hippocampus 2 weeks post-injury, and remained lowered 8 months later. At this chronic stage increased CBF is apparent in the ipsilateral thalamus (arrowhead). Scale bar = 1 mm (Courtesy of Drs. Riikka Immonen, Nick Hayward and Olli Gröhn)
intensity on $T_2^*$- (spin echo) or $T_2^*$-weighted (gradient echo) MR images. Basically two approaches for cellular labeling have been employed: (1) in vivo uptake of intravenously injected contrast agent by (circulating) phagocytic cells or (2) in vitro labeling of isolated cells that are subsequently reinjected. Ultrasmall particles of iron oxide (USPIOs) with a long blood half-life have been mostly applied for in vivo uptake by monocytes or macrophages, whereas larger particles of iron oxide (such as small particles of iron oxide (SPIOs) or micron-sized particles of iron oxide (MPIOs)), which provide maximum contrast effects, may be effectively used for in vitro cell labeling [43, 44]. Both cell labeling strategies have been applied in a number of experimental studies that sought to characterize the spatio-temporal profile of leukocyte infiltration in neuroinflammatory diseases, such as multiple sclerosis and stroke (see Stoll and Bendszus [44] and references therein). These studies have, for instance, shown that cellular infiltration is not necessarily linked to BBB disruption. In a controlled cortical impact injury model in mice, intravenous MPIO injection resulted in discrete, punctuate areas of MRI hypointensity in and around contusion sites (see Fig. 3), which correlated histologically with presence of MPIO-containing macrophages [45]. This was evident even when MPIOs were administered 6 days before neurotrauma, which suggests that previously labeled macrophages migrate to the trauma site and that the hypointense MR signals were not caused by nonspecific accumulation of MPIO particles that traversed an acutely leaky BBB.

**MRI of Cell Adhesion Molecules**

Advances in design and synthesis of contrast agents have recently initiated MRI studies that take advantage of molecular targeting with tailored contrast agents to measure particular disease markers [46–49]. This kind of molecular imaging with MRI typically employs (super)paramagnetic nano- or microparticles with conjugated ligand, e.g., an antibody. Besides a (super)paramagnetic agent, e.g., gadolinium
or iron oxide, particles may contain other labels, such as fluorescent dye, for detection with alternative imaging modalities. In addition, polyethylene-glycol coating is often used to increase circulation time and bioavailability.

Molecular MRI contrast agents can be constructed by direct functionalization of (super)paramagnetic agents or particles (e.g., MPIOs) with targeting peptides or antibodies [50]. An alternative platform for the design and formation of molecular imaging probes makes use of lipid formulations, such as liposomes and micelles, that are build up from amphiphilic lipids [51] (Fig. 4). These lipid-based structures can carry various contrast agents in relatively high amounts, which, in combination with the ease of preparation, flexibility and biocompatibility, make these nanoparticles highly adequate for preclinical cellular and molecular imaging studies as well as for drug delivery in patients [52].

Despite the potential of targeted contrast agents to bind and depict molecular markers, molecular MRI studies in the brain are complicated due to the presence of the BBB, which restricts the blood-to-brain transfer of relatively large intravascular molecular MRI probes. For that reason...
molecular MRI in CNS injury models have largely focused on
detection of markers that are expressed on the luminal side of
cerebral blood vessels, such as cell adhesion molecules.
Studies in rodent models of neuroinflammatory diseases have
demonstrated that paramagnetic or superparamagnetic com-
pounds targeted against endothelial cell adhesion molecules,
such as P- and E-selectin and VCAM-1, induce local contrast
enhancement on $T_1$- or $T_2^*$-weighted MR images, respective-
ly, reflective of activated cerebral endothelium [53–56]. In a
model of lateral fluid percussion in rats, Chapon et al. [57]
recently showed that injection of selectin-targeted iron oxide
particles resulted in persistent $T_2^*$ shortening in the trauma-
tized brain hemisphere, while initial $T_2$ shortening recovered
within 1 h after injection of non-targeted iron oxide particles.

**MRI of Vascularity**

There is increasing evidence that chronic vascular remodel-
ing can occur after CNS injury, which may contribute to

Fig. 4 Schematic representation of two types of lipid-based nano-
particles, i.e., micelles and liposomes, that can be formed with
various constituents, such as MR contrast agent, fluorescent dye,
polyethylene-glycol (PEG), and a targeting ligand for molecular
imaging purposes. Courtesy of Drs. Geralda A. van Tilborg and
Willem J. Mulder

Fig. 5 $\Delta R_2$ (a) and Q map (b) calculated from steady-state contrast-
enhanced MRI of a coronal rat brain slice with a subcortical infarct at
7 days after transient unilateral middle cerebral artery occlusion, and
corresponding histological sections with vessel staining of the ipsi-
(c) and contralesional hypothalamic region (d). Relatively high $\Delta R_2$ after
administration of a superparamagnetic blood pool agent (USPIO) indicates a high density of microvessels with a relatively small
diameter in the ipsilesional hypothalamic region (black arrow) as
compared to contralesional (white arrow) (a). This area also revealed a
high $Q$ value (b), reflective of enhanced microvessel density, which
was confirmed by more vessel staining with von Willebrand Factor
(DAB-enhanced; brown) in the ipsi- (c) as compared to contralesional
hypothalamic region (d) (nuclei were stained with hematoxylin; blue).
Modified from Seevinck et al. [59] (Courtesy of Peter R. Seevinck,
Lisette H. Deddens and H. Elga de Vries)
reestablishment of perfusion, removal of waste product or tissue repair [31, 58]. Preclinical studies have reported traumatic brain injury-induced angiogenesis, which could be promoted with specific therapeutic interventions, leading to improvement of functional outcome [31]. Hayward et al. [23] have recently detected brain region-specific increases in vascular density at 9 months after lateral fluid percussion injury in rats, which, however, showed no clear correlation with restoration of CBF or recovery of neurological function. Evidently, the pattern of changes in brain tissue vascularity and its relation to pathophysiology or plasticity after neurotrauma require further investigation.

Various MRI methods are available for detection of vascular changes in injured brain (for a review, see Seevinck et al. [59]). A particularly powerful approach is offered by steady-state contrast-enhanced MRI, which involves concurrent gradient echo and spin echo MRI acquisition in combination with injection of a superparamagnetic blood pool agent (e.g., USPIOs). This enables measurement of changes in MR relaxation rates $R_1$ and $R_2$ that are dependent on local cerebral blood volume (CBV), which allows calculation of different vascular parameters, such as total and microvascular blood volume fraction, microvessel density, and vessel size index [59, 60]. A serial steady state contrast-enhanced MRI study in rats after experimental stroke has reported chronically increased microvascular CBV and microvessel density subsequent to subacute vascular compression, pointing toward active revascularization of post-ischemic tissue [61]. Figure 5 shows steady state contrast-enhanced MRI data obtained from a rat at 7 days after subcortical stroke. The elevated $\Delta R_2$ (a measure of microvascular CBV) and $Q (\Delta R_2/\Delta R_2^{2/3};$ a measure of microvessel density) in part of the lesion reflect increased vascularization, which was confirmed histologically [59].

**Conclusion**

Recent MRI strategies that focus on detection of specific cerebrovascular alterations associated with neurological disorders have advanced the abilities of integrative in vivo studies on the pathophysiology of CNS injury. In particular, the developments in contrast agent chemistry have provided unique opportunities for in vivo MRI-based detection of cellular and molecular processes at the level of the cerebrovasculature, such as the invasion of leukocytes and upregulation of cell adhesion molecules. The application of these cellular and molecular MRI methods in combination with other structural and functional MRI techniques, allows direct correlation of different pathophysiological events in an experimental setting, which can contribute to elucidation of the interaction between the various pathways through which traumatic brain injury develops. Furthermore, preclinical experiments with multiparametric MRI protocols may lead to the identification of exclusive disease biomarkers and distinctive therapeutic targets that can be translated to the clinic. The potential of advanced MRI techniques, such as susceptibility-weighted MRI and diffusion tensor imaging, to aid in the classification of traumatic brain injury patients has recently been described by Kou et al. [62]. These methods, in combination with functional MRI, may also inform on brain remodelling or neural plasticity at later post-traumatic stages [63], which could lead to identification of key processes that contribute to functional recovery and may be stimulated with therapeutic strategies.

In conclusion, studies with novel contrast-enhanced MRI strategies in animal models will provide more insights into the potential of these methods in terms of their sensitivity and specificity to detect distinctive characteristics of traumatic brain injury. The outcome of such studies, as well as adequate evaluation of safety issues, will be critical to determine whether these contrast-enhanced MRI techniques have potential for clinical application in neurotrauma patients.

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