pH-weighted molecular MRI in human traumatic brain injury (TBI) using amine proton chemical exchange saturation transfer echoplanar imaging (CEST EPI)

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ABSTRACT

Cerebral acidosis is a consequence of secondary injury mechanisms following traumatic brain injury (TBI), including excitotoxicity and ischemia, with potentially significant clinical implications. However, there remains an unmet clinical need for technology for non-invasive, high resolution pH imaging of human TBI for studying metabolic changes following injury. The current study examined 17 patients with TBI and 20 healthy controls using amine chemical exchange saturation transfer echoplanar imaging (CEST EPI), a novel pH-weighted molecular MR imaging technique, on a clinical 3T MR scanner. Results showed significantly elevated pH-weighted image contrast (MTRasym at 3 ppm) in areas of T2 hyperintensity or edema (P < 0.0001), and a strong negative correlation with Glasgow Coma Scale (GCS) at the time of the MRI exam ($R^2=0.4777$, $P=0.0021$), Glasgow Outcome Scale - Extended (GOSE) at 6 months from injury ($R^2=0.5334$, $P=0.0107$), and a non-linear correlation with the time from injury to MRI exam ($R^2=0.6317$, $P=0.0004$). This evidence suggests clinical feasibility and potential value of pH-weighted amine CEST EPI as a high-resolution imaging tool for identifying tissue most at risk for long-term damage due to cerebral acidosis.

1. Introduction

Traumatic brain injury (TBI) is a major world health problem (Ghajar, 2000; Maas et al., 2008) and it is one of the leading causes of mortality and disability (Finfer and Cohen, 2001). A cascade of biological events occurs following traumatic insult to the brain leading to long-term secondary brain damage, including tissue ischemia from impaired cerebral perfusion (Coles et al., 2002; Diringer et al., 2002), amino acid excitotoxicity (Faden et al., 1989), and interstitial acidification (Ham and Traystman, 1996; Siesjo et al., 1993) from the resulting energy depletion and metabolic crisis (Vespa et al., 2005). Each of these physiologic changes have been shown to be directly or indirectly related to the degree of neuronal damage following injury and may be useful for predicting outcome. Reduced cerebral perfusion has been shown to be linked with outcome (Kirkness et al., 2005; Prabhasakar et al., 2014), while the degree of excitotoxicity has also been linked with outcome (Timofeev et al., 2011) and is the target of a variety of treatment strategies (Willis et al., 2004). Extracellular acidosis within the brain following TBI occurs due to metabolic dysfunction and results in a variety of subsequent injury mechanisms including increased permeability of calcium acid-sensing ion channels (Yermolaieva et al., 2004), further excitotoxic (McDonald et al., 1998) or oxidative (Ying et al., 1999) injury, reduced glutamate uptake by astrocytes (Swanson et al., 1995), impairment of astrocytes to maintain glucose metabolism (Swanson and Benington, 1996), and blood-brain barrier disruption (Nagy et al., 1985). Further, measures of extracellular pH, which is intimately linked with intracellular pH (Mellergard et al., 1994), has been shown to be a strong marker of metabolic dysfunction and has been linked with increased mortality following TBI (Clausen et al., 2005; Timofeev et al., 2013; Yokota et al., 2000). Despite human and...
experimental evidence suggesting acidification of the brain may occur after TBI (McIntosh et al., 1987; Shiogai et al., 1999; Zauner et al., 2002), there remains a critical need for tools to non-invasively identifying regions of the brain at risk for low extracellular pH in patients following TBI.

Amine chemical exchange saturation transfer echo planar imaging (amine CEST EPI; Fig. 1A) is a fast, noninvasive molecular MRI technique with sensitivity to amino acid concentration and pH (Harris et al., 2016; Harris et al., 2018). By preparing the MRI signal through selective saturation of the fast exchanging, pH-dependent amine protons (-NH2) on glutamine or glutamate (Fig. 1B), amine CEST imaging results in signal contrast in tissues with high glutamine or glutamate concentration and low pH. Since there is a well-documented increase in both extracellular glutamate (Bullock et al., 1998; Chamoun et al., 2010; Folkersma et al., 2011; Vespa et al., 1998) and mobile glutamine (Harris et al., 2012; Palmer et al., 1993; Xu et al., 2011) following TBI from damage to neurons and astrocytes (Fig. 1C), respectively, we theorized amine CEST EPI may provide unique insight into the biologic status of the brain independent of other MR image contrasts. We hypothesized areas of T2 hyperintensity adjacent to contusion would represent the water and amine protons, respectively; and $k_a$ is the exchange rate of protons from pool $a$ to pool $b$ as given by $\left(M_{ba}/M_{ab}\right)k_b$; $\omega_1$ is the RF pulse amplitude as given by $\omega_1 = \gamma B_1(t)$, where $\gamma$ is the gyromagnetic ratio and $B_1(t)$ is given in $\mu T$; $\delta a = \omega - \omega_a$ and $\delta b = \omega - \omega_b$, where $\omega$ is the applied radiofrequency (RF) irradiation frequency, $\omega_a$ is the water proton resonance frequency, and $\omega_b$ is the amine proton resonance frequency; $T_{1a}$ and $T_{1b}$ are the longitudinal relaxation times of water and amine protons, respectively; and $C_{1a} = (1/T_{1a}) + k_\alpha$, $C_{2a} = (1/T_{2a}) + k_\alpha$, $C_{1b} = (1/T_{1b}) + k_\beta$, $C_{2b} = (1/T_{2b}) + k_\beta$ represent the sum of exchange and relaxation rates. Eq. 1 can be solved analytically to yield

$$ M(t) = e^{\omega_1 M_0(X/e) - (X/e)} $$

(3)

The effects of pH on the CEST signal can then be modeled by accounting for the chemical exchange rate between amine and water protons. The chemical exchange between amine and water protons can be characterized as a base-catalyzed process, governed by the equation:

$$ k_b = k_\alpha + k_{base} * 10^{-(14-pH)} $$

(4)

where $k_\alpha$ is the default exchange rate, $k_{base}$ is the base-catalyzed rate constant, and $k_b$ is the exchange rate of protons from the metabolite proton pool to the water pool (Liepinsch and Otting, 1996).

### 2. Materials and methods

#### 2.1. Theory

The magnetization of water protons undergoing two-compartment chemical exchange with amine protons is described by the Bloch-McConnell equations (Woessner et al., 2005) in the form of:

$$ \frac{dM(t)}{dt} = X \cdot M(t) - c $$

(1)

where

$$ M = \begin{bmatrix} M_{aa} \\ M_{ba} \\ M_{ab} \\ M_{bb} \end{bmatrix}, \quad X = \begin{bmatrix} C_{aa} & k_b & 0 & 0 \\ k_a & C_{ab} & 0 & -\delta b \\ 0 & 0 & C_{bb} & -\delta a \\ 0 & 0 & 0 & C_{ba} \end{bmatrix}, \quad c = \begin{bmatrix} 0 \\ 0 \\ 0 \\ 0 \end{bmatrix} $$

(2)

pool $a$ and pool $b$ represent the water and amine protons, respectively; $M_{aa}$ and $M_{bb}$ are the equilibrium magnetizations of water and amine protons, respectively; $k_b$ is the exchange rate of protons from pool $b$ to pool $a$; $k_a$ is the exchange rate of protons from pool $a$ to pool $b$ as given by $\left(M_{ba}/M_{ab}\right)k_b$; $\omega_1$ is the RF pulse amplitude as given by $\omega_1 = \gamma B_1(t)$, where $\gamma$ is the gyromagnetic ratio and $B_1(t)$ is given in $\mu T$; $\delta a = \omega - \omega_a$ and $\delta b = \omega - \omega_b$, where $\omega$ is the applied radiofrequency (RF) irradiation frequency, $\omega_a$ is the water proton resonance frequency, and $\omega_b$ is the amine proton resonance frequency; $T_{1a}$ and $T_{1b}$ are the longitudinal relaxation times of water and amine protons, respectively; and $C_{1a} = (1/T_{1a}) + k_\alpha$, $C_{2a} = (1/T_{2a}) + k_\alpha$, $C_{1b} = (1/T_{1b}) + k_\beta$, $C_{2b} = (1/T_{2b}) + k_\beta$ represent the sum of exchange and relaxation rates. Eq. 1 can be solved analytically to yield

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#### 2.2. Phantom preparation and estimation of exchange rate constants

To determine the constants $k_\alpha$ and $k_{base}$ for amino acid amines, a series of glutamine and glutamate phantoms with pH ranging from 4.8 to 7.5 were prepared for MRI evaluation. Glutamine or glutamate were combined with phosphate-buffered saline (PBS) in 50 mL falcon tubes to reach a concentration of 50 mM. 0.5 mM Gd-DTPA (Magnevist®) was then injected into each phantom solution to reduce relaxation times (i.e. $T_1$ and $T_2$) near physiologic levels. The pH values of sample solutions were then titrated to 18 different values ranging from 4.8 to 7.5, with acid (1 N HCl) and base (1 N NaOH) solution. Phantom pH values were re-recorded after the MRI scanning and used as the final pH values for data fitting using the Bloch-McConnell equations described previously.

#### 2.3. Mathematical simulations of CEST effects

To demonstrate the relationship between amine CEST image contrast and pH for both glutamine and glutamate we employed the Bloch-McConnell simulation framework for CEST EPI previously described (Harris et al., 2016), using exchange rate values estimated from phantom experiments. Specifically, we simulated the z-spectrum from pH values ranging from 5.5 through 8.0 by first assuming the concentration of amine protons was 50 mM, which is reasonable
considering 1) the concentration of amino acids in normal neural tissues has been estimated around 20–25 mM (Perry et al., 1971); 2) many amino acid derivative metabolites including norepinephrine, 5-hydroxytryptophan, levodopa and other neurotransmitters possess a similar amine functional group that will also contribute to the CEST signal at 3.0 ppm; and 3) we have previously shown these amino acids exhibit similar pH-dependent exchange rate characteristics and CEST effects (Harris et al., 2015). Additionally, we assumed the MR characteristics of water protons were similar to values in normal white matter (Harris et al., 2015; Wright et al., 2008), assumed the MR characteristics of the amine protons were similar to those in the literature (T1b = 0.2 s and T2b = 0.1 s) (Choi et al., 2006; Li et al., 2008; Traber et al., 2004; Voessner et al., 2005), and used a set of 3 × 100 ms Gaussian radiofrequency CEST saturation pulses with an amplitude of 6 μT, similar to previous studies (Harris et al., 2016; Harris et al., 2018). Lastly, using similar values of T1 and the exchange rates estimated from the phantom experiments, we simulated the CEST signal for a range of concentrations from 5 mM to 50 mM and a range of transverse relaxation times from T2 = 50 ms to 500 ms.

2.4. Quantification of CEST contrast

During a CEST imaging experiment, the RF saturation frequency ω is swept across a range of values to obtain a “z-spectrum” (Bryant, 1996). To reduce the effects of T1 and T2 weighting along with other variables, the attenuation of bulk water magnetization following a saturation pulse is often described by the magnetization transfer ratio (MTR), given by:

\[ MTR(\omega) = \frac{S(\omega)}{S_0} \]  

where \( S(\omega) \) is the magnitude of water proton MR signal available after the saturation pulse with frequency \( \omega \) and \( S_0 \) is the MR signal available without application of any RF saturation pulses. Since MTR can be affected by symmetric effects of direct water saturation and conventional magnetization transfer effects, CEST contrast is often described by the asymmetry in the magnetization transfer ratio (MTR asym) with respect to water proton resonance:

\[ MTR_{\text{asym}}(\omega) = \frac{S(-\omega) - S(\omega)}{S_0} \]  

For the current simulation, MTR at spectral points between −5.0 and +5.0 ppm were calculated to obtain simulated z-spectra in these tissues, and the MTR asym at 3.0 ppm was used as a final measure of pH-dependent amine CEST contrast for use in subsequent patient studies.

2.5. Human participants

A total of 17 patients with traumatic brain injury and 20 healthy volunteers were enrolled in this prospective, Health Insurance Portability and Accountability Act (HIPAA) compliant study approved by the Institutional Review Board at our institution. All patients and volunteers signed informed written consent to have advanced imaging collected. A summary of TBI patient demographics is shown in Table 1.

2.6. Magnetic resonance imaging

All patients underwent anatomic and amine CEST-EPI on a 3T MR scanner (Siemens Trio, Prisma, or Skyra; Siemens Medical; Erlangen, Germany). All patients received a standardized anatomic imaging protocol consisting of at least T2-weighted turbo spin echo images, diffusion tensor images (DTI), susceptibility-weighted gradient echo images (SWI), fluid-attenuated inversion recovery (FLAIR) images, and 1 mm isotropic parameter matched 3D inversion-recovery gradient recalled echo (IR-GRE) images. Amine CEST-EPI was collected with the following scan parameters: field-of-view (FOV) = 256 × 256 mm,
In order to estimate the static magnetic field distortion, $\Delta B_0$, we employed a novel method using iterative down-sampling with Lorentzian fitting (Yao et al., 2018). Briefly, all CEST data were first preprocessed with motion correction (mcflirt function, FSL; Oxford) (Jenkinson et al., 2002). Next, a slice-by-slice $\Delta B_0$ map estimation was achieved by first normalizing the raw CEST data with respect to the $S_0$ reference scan data, resulting in $z$-spectral measurements, then performing $k$-means clustering cluster the pixels with similar patterns in the $z$-spectrum. The number of clusters for each iteration was set to $3^m$, where $m$ is the iteration number. Next, the average $z$-spectral data was estimated across all pixels within the same cluster, and then the mean $z$-spectral data were fit to the Lorentzian function in the form: $c - \frac{\alpha}{c^2 - x^2 + \beta x}$, where $c$ indicates the shift in $z$-spectral center point, $\beta$ relates to the width of the Lorentzian shape, and $\alpha$ and $\beta$ together determine the $y$-scaling of the function. The constraint bound for $\Delta B_0$ estimation was set to $\pm 1$ mppm. The magnetic field distortion, or inhomogeneity in the static magnetic field ($\Delta B_0$), was estimated by the value of $\alpha$ in the Lorentzian model and subsequently stored for all pixels. Lastly, a 3D Gaussian filter with standard deviation of unity was applied to the final $\Delta B_0$ map to remove potential erroneous estimations.

### 2.8. Region of interest selection

All images were registered to high-resolution, 1 mm isotropic T1-weighted images using a 6° of freedom rigid body transformation and mutual information cost function (FSL, FMRIB, Oxford, UK) for subsequent analyses. $MTR_{asym}$ at 3 ppm and magnetic field fluctuations ($\Delta B_0$) were measured within areas of T2 hyperintensity along with areas of contusion as defined by signal voids on susceptibility ($T2^*$) weighted imaging. Additionally, regions with apparent diffusion coefficient (ADC) $<0.6 \text{um}^2/\text{ms}$, suggestive of ischemia, were included. Measurements in patients with multiple sites of T2 hyperintensity, contusion, or potential ischemia were averaged together resulting in a single measurement per patient. As a reference, a 2 cm diameter spherical region of interest was identified within normal-appearing white matter (NAWM), distal from any apparent lesion, hemorrhage or contusion as well as a similar region of interest in white matter within the brains of healthy volunteers. Line plots ranging from 13 to 20 mm were created by traversing the edge of the T2 hyperintense lesion in TBI patients in order to document any apparent spatial colocalization of acidic and ischemic tissue using $MTR_{asym}$ and ADC, respectively.

### 2.9. Clinical examination and outcome measures

The Glasgow Coma Scale (GCS) is a commonly used clinical measure of initial TBI severity, has been shown to be an early prognostic marker for outcome (Jiang et al., 2002), and is inversely correlated with likelihood of significant morbidity and mortality (Jiang et al., 2002; Lawrence F. Marshall et al., 1991). Average GCS at the time of MRI was $7.4 \pm 0.9$ S.E.M. (range = 3 to 14). Additionally, we evaluated the extended Glasgow outcome scale (GOSE) at 6 months from the time of injury as a widely accepted clinical measure of outcome in patients with TBI (Jiang et al., 2002; Marshall et al., 1991; Levin et al., 2001; Woischneck and Firsching, 1998). Six-month GOSE scores were available for 11 of the 17 patients in the current study and averaged $4.2 \pm 0.6$ S.E.M. (range = 1 to 8). In the current study we tested the hypothesis that CEST measures of $MTR_{asym}$ at 3 ppm within T2 hyperintense areas adjacent to contused tissue correlated with GCS at the time of the MRI examination and could predict GOSE 6 months following initial injury.

### 3. Statistical analyses

All values are presented as mean $\pm$ standard error of the mean (S.E.M.). A linear mixed effects model and expected marginal means estimations were used to compare measurements of $MTR_{asym}$ at 3 ppm in regions of normal appearing white matter (NAWM), T2 hyperintense regions, areas of contusion, areas of potential ischemia as indicated by ADC $<0.6 \text{um}^2/\text{ms}$, and healthy white matter from control participants. Similarly, a linear mixed effects model and expected marginal means estimations were used to compare measurements of $\Delta B_0$ at 3 ppm in regions of normal appearing white matter (NAWM), T2 hyperintense regions, areas of contusion, and measurements from white matter in healthy volunteers. A total of 5 comparisons were performed for $MTR_{asym}$ and $\Delta B_0$ at 3 ppm, so a Bonferroni-corrected $p$-value of
0.01 was used to indicate statistical significance. Additionally, linear regression was used to compare CEST measures of MTR_{asym} at 3 ppm within areas of T2 hyperintensity to both GCS at the time of MRI examination and GOSE at 6 months from injury in patients with TBI. We used nonlinear regression to test whether there was a logarithmic association between the time between injury and MRI exam and MTR_{asym} at 3 ppm within areas T2 hyperintensity, which may suggest a higher degree of excitotoxicity and acidosis in the early acute phases of TBI. Lastly, we used a multiple linear regression using a combination of CEST measures and the time between injury and MRI exam in order to predict both GCS at the time of the MRI exam and GOSE at 6 months from injury.

4. Results

4.1. Phantom and simulation results

Phantom experiments involving CEST imaging of 50 mM vials containing glutamine and glutamate exhibited distinct changes at lower pH values (Fig. 2A–B), consistent with previous studies. Results clearly showed
a higher MTR<sub>asym</sub> at 3 ppm for pH values below around 7.0 in for glutamine and around 7.5 for glutamate, with an estimated pH-dependent exchange rate, in Hz, for glutamine and glutamate of

\[
k_{ex}(\text{Gln}) = 163.2 + 1.562 \times 10^{[\text{pH}-3]}
\]

and

\[
k_{ex}(\text{Glu}) = 781 + 2.522 \times 10^{[\text{pH}-4]}
\]

respectively (Fig. 2C).

Consistent with previous simulations and other studies (Harris et al., 2016; Harris et al., 2015; Harris et al., 2018; Jin and Kim, 2014; Jin et al., 2012; Sun et al., 2013), CEST measures of MTR<sub>asym</sub> at 3 ppm from bulk water for both glutamine and glutamate demonstrated sensitivity to pH, concentration, and T2 relaxation time. In particular, the z-spectrum showed a slight decrease in normalized MR signal intensity with decreasing pH assuming a constant glutamine concentration of 50 mM (Fig. 3A–B). The MTR<sub>asym</sub> around 2-3 ppm showed the optimal sensitivity to pH-related amine proton exchange (Fig. 3C), which increased with increasing glutamine and glutamate concentration (Fig. 3D) and increasing tissue T2 (Fig. 3E), which supports the hypothesis that amine CEST EPI may provide value in identifying tissue at risk of acidity due to excitotoxicity, particularly in areas of T2 hyperintensity.

4.2. MTR<sub>asym</sub> at 3 ppm within NAWM, T2 hyperintensity, contusion and ischemia in TBI

Whole brain CEST-EPI data within healthy volunteers exhibited relatively uniform, low values of MTR<sub>asym</sub> at 3 ppm (Fig. 4), averaging 0.58% ± 0.05% standard error of the mean (S.E.M.), which was similar to average values of MTR<sub>asym</sub> at 3 ppm in a 2 cm diameter region of relatively NAWM on anatomic MRI in TBI patients (Fig. 7A; 0.66% ± 0.05%). No difference in MTR<sub>asym</sub> at 3 ppm was observed between NAWM in TBI patients and white matter in healthy volunteers (P = 0.2371). This was in stark contrast to MTR<sub>asym</sub> at 3 ppm within lesions in TBI patients (Fig. 4). As illustrated in Fig. 5A–C and quantified in Fig. 7A, regions of T2 hyperintensity near areas of contusion exhibited a significantly higher estimate of median MTR<sub>asym</sub> at 3 ppm (1.62% ± 0.09%) compared with NAWM (Fig. 7A; P < 0.0001). Areas of potential ischemia as indicated by ADC < 0.6 um²/ms exhibited an average MTR<sub>asym</sub> at 3 ppm around 0.86% ± 0.34%, which was not significantly different from NAWM (Fig. 7A; P = 0.4988). No significant association was observed between ADC and MTR<sub>asym</sub> at 3 ppm in areas of T2 hyperintensity (Fig. 7B; R² = 0.1267, P = 0.1609), as also illustrated by the line plots through the T2 hyperintense lesions in Fig. 5A–C, nor between ADC and MTR<sub>asym</sub> at 3 ppm in areas of contusion (Fig. 7C; R² = 0.1438, P = 0.2013).

4.3. Magnetic field distortions (ΔB₀) within NAWM, pericontusion, and hemorrhage in TBI

Perturbations or distortions in the homogeneous, static magnetic field within the MRI system can lead to errors in MTR<sub>asym</sub> measurements and may provide unique sensitivity to regions of resolving blood products. The median ΔB₀ within NAWM averaged 0.042 ppm ± 0.011 ppm S.E.M., which was not significantly different (ANOVA, P = 0.1287) from the ΔB₀ estimated within areas of T2 hyperintensity (Fig. 7D; 0.039 ppm ± 0.018 ppm) or contusion (Fig. 7D; 0.077 ppm ± 0.012 ppm). No difference was observed between NAWM in TBI patients and white matter from healthy volunteers (P = 0.8213), which averaged 0.045 ppm ± 0.009 ppm S.E.M.

4.4. Correlation between MTR<sub>asym</sub> at 3 ppm and clinical outcome

A strong negative correlation was observed between GCS at the time of MRI and MTR<sub>asym</sub> at 3 ppm within contusion plus pericontusional tissue (Fig. 8A; R² = 0.4777, P = 0.0021), suggesting patients with the highest degree of brain acidity had worse function. Similarly, MTR<sub>asym</sub>
at 3 ppm was inversely correlated with GOSE at 6 months (Fig. 8B; \( R^2 = 0.5334, P = 0.0107 \)), also suggesting patients with the lowest pH had worse long-term outcomes. We also observed a significant non-linear (logarithmic) association between MTR_{asym} at 3 ppm and the time between the injury and MRI exam (Fig. 8C; Model: \( \text{log}_{10}(\text{time from injury}) = B \times \{\text{MTR}_{asym}@3\text{ppm}\} + C; \ R^2 = 0.6317, \ P = 0.0004 \)), implying a higher acidity during acute injury compared with more subacute or chronic stages of TBI. Lastly, multiple linear regression suggested a combination of MTR_{asym} at 3 ppm along with the time between injury and MRI could better predict GCS at the time of the MRI exam (Fig. 8D; Model: GCS = \( B_1 \times \text{log}_{10}(\text{time from injury}) + B_2 \times \{\text{MTR}_{asym}@3\text{ppm}\} + C; \ R^2 = 0.6539, \ P = 0.0026 \)) and GOSE at 6 months (Fig. 8E; Model: GOSE = \( B_1 \times \text{log}_{10}(\text{time from injury}) + B_2 \times \{\text{MTR}_{asym}@3\text{ppm}\} + C; \ R^2 = 0.6539, \ P = 0.0026 \)).

5. Discussion

Following TBI, a series of secondary brain damage occurs including reduced cerebral blood flow, hypoxia, and excitotoxicity (Bullock et al., 1995; Bullock et al., 1998; Miller, 1985), all of which result in cerebral acidosis (Clausen et al., 2005; Hovda et al., 1992; Kalimo et al., 1981). Results from the current study are consistent with the hypothesis that areas of T2 hyperintensity adjacent to contusions exhibit significant acidosis, arising from excitotoxicity. Further, our results suggest the degree of acidosis is highly correlated with both clinical function as well as outcome, as evidenced by correlation with GCS at the time of the MRI examination and GOSE at 6 months, respectively. Eloquent studies using multiparametric sensors inserted into the brain in patients with TBI have verified a potential association between low extracellular pH and outcomes including mortality (Gupta et al., 2004; Timofeev...
Interestingly, we also observed a correlation between time from initial injury and degree of acidosis as evidenced by CEST contrast, which appears to be consistent with these same studies involving temporal pH monitoring using intracranial sensors, which showed acidic pH measurements reversing or normalizing in subsequent days following injury (Timofeev et al., 2013). Few studies have used molecular MR imaging to evaluate pH changes following TBI, and the current study represents the first to utilize amine CEST EPI on 3T clinical MR systems to evaluate acidosis in human TBI at high spatial resolution. A number of preclinical studies have shown alterations in intra- and extracellular pH using phosphorous nuclear magnetic resonance ($^{31}$P NMR) spectroscopic techniques (McIntosh et al., 1987; Vink et al., 1987). However, $^{31}$P NMR has extremely low sensitivity, therefore data must be acquired using large voxels and long scan times, making it less suitable for routine clinical examination and differentiation of acidic subregions in heterogeneous lesions. A recent study by Wang et al. (2017) utilized amide proton transfer (APT) CEST imaging on a preclinical MR system (4.7T) in untreated rats following TBI, noting a characteristic decrease in MTR$_{sym}$ at 3.5 ppm (for amide protons), which has been shown to similarly reflect a decreasing pH (Note: decreasing pH results in increased MTR$_{sym}$ at 3 ppm for amine protons and decreased MTR$_{sym}$ at 3.5 ppm for amide protons (Jones et al., 2013)). Further, the approach in the current study focusing on amine CEST imaging has specific advantages over APT imaging including short radiofrequency irradiation times due to the fast exchange rates of the amine protons, so fast readout strategies including

![Fig. 6. T2-weighted anatomic MRI, pH-weighted amine CEST EPI, T2*-weighted MRI, and estimated magnetic field distortion ($\Delta B_0$) (Left-to-Right) for A) a 45 year old male patient (Subject #1) with diffuse axonal injury and white matter microhemorrhages after suffering a fall from hang gliding; B) a 35 year old male patient (Subject #2) who was struck by a motor vehicle and suffered bilateral subarachnoid and intraparenchymal hemorrhage; and C) a 28 year old male subject (Subject #3) who suffered bilateral hemorrhagic contusions, subarachnoid hemorrhage in the frontal lobe, and subdural hematomas along the cerebral convexities after being in a motor vehicle accident. Red arrows highlight areas of microbleeds and/or diffuse axonal injury within the corpus callosum. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)](image)
EPI can be used to obtain high resolution, whole brain images within clinically realistic scan times.

It is important to point out a few limitations to the current study. First, no correction for B1 inhomogeneities were performed. As B1 can vary as much as ± 20% (Sengupta et al., 2017), our simulations (not shown) suggest that acidic tissue within areas of high amino acid concentration (~50 mM) can have MTRasym at 3 ppm that varies ± 15% (e.g. from 3.65% to 4.3%). Additionally, the results observed in the current study should be taken as preliminary, since the relatively small number of patients included may not represent the strength of trends observed in a larger cohort of patients with different injury profiles. Future studies with a larger number of patients and inclusion of B1 correction are warranted.

In summary, the current study demonstrates the feasibility and potential clinical value of pH-weighted amine CEST EPI as a high-resolution imaging tool for identifying tissue most at risk for long-term damage due to cerebral acidosis. Results suggest low pH in contusion and pericontusional regions correlate strongly with clinical function and 6-month outcome measures.

Author disclosure statement

Ellingson – Advisory Board – Hoffman La-Roche; Siemens; Nativis; Medicenna; MedQIA; Bristol Meyers Squibb; Imaging Endpoints; Agios. Paid Consultant – Nativis; MedQIA; Siemens; Hoffman La-Roche; Imaging Endpoints; Medicenna; Agios. Grant Funding – Hoffman La-Roche;
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