Acute Pressure Changes in the Brain are Correlated With MR Elastography Stiffness Measurements: Initial Feasibility in an In Vivo Large Animal Model

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Purpose: The homeostasis of intracranial pressure (ICP) is of paramount importance for maintaining normal brain function. A noninvasive technique capable of making direct measurements of ICP currently does not exist. MR elastography (MRE) is capable of noninvasively measuring brain tissue stiffness in vivo, and may act as a surrogate to measure ICP. The objective of this study was to investigate the impact of changing ICP on brain stiffness using MRE in a swine model.

Methods: Baseline MRE measurements were obtained, and then catheters were surgically placed into the left and right lateral ventricles of three animals. ICP was systematically increased over the range of 0 to 55 millimeters mercury (mmHg), and stiffness measurements were made using brain MRE at vibration frequencies of 60 hertz (Hz), 90 Hz, 120 Hz, and 150 Hz.

Results: A significant linear correlation between stiffness and ICP in the cross-subject comparison was observed for all tested vibrational frequencies (P < 0.01). The 120 Hz (0.030 ± 0.004 kilopascal (kPa)/mmHg, P < 0.0001) and 150 Hz (0.031 ± 0.008 kPa/mmHg, P = 0.01) vibrational frequencies had nearly identical slopes, which were approximately two- to three-fold higher than the 90 Hz (0.017 ± 0.002 kPa/mmHg, P < 0.0001) and 60 Hz (0.009 ± 0.002 kPa/mmHg, P = 0.001) slopes, respectively.

Conclusion: In this study, MRE demonstrated the potential for noninvasive measurement of changes in ICP. Magn Reson Med 79:1043–1051, 2018. © 2017 The Authors Magnetic Resonance in Medicine published by Wiley Periodicals, Inc. on behalf of International Society for Magnetic Resonance in Medicine. This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

Key words: intracranial pressure; magnetic resonance elastography; brain stiffness; viscoelasticity

INTRODUCTION

The homeostasis of intracranial pressure (ICP) is of paramount importance for maintaining normal brain function. Several pathological conditions can lead to a loss of homeostasis that results in elevated ICP. High levels of ICP can compress important brain regions, specifically the brain stem, triggering high-risk side effects such as abnormal breathing, cardiovascular stress, and potentially death. Maintaining normal ICP is of utmost importance for patient care, including postsurgical monitoring in patients with neurosurgical conditions. Currently, a noninvasive method for measuring and monitoring ICP does not exist.

A lumbar puncture (LP) is a valuable diagnostic tool that can be used to measure ICP. During the procedure, cerebrospinal fluid (CSF) samples also can be obtained for treatment planning; however, medical conditions such as obstructive hydrocephalus can contraindicate performing a LP. Hydrocephalus is a common condition that results from the obstruction of CSF flow or resorption. However, different types of hydrocephalus exist, which look similar radiologically but are caused by different mechanisms. For instance, obstructive or noncommunicating hydrocephalus results from the blockage of CSF flow anywhere from the foramen of Monro to the outlet foramina of the fourth ventricle. Common causes include benign and malignant tumors, aqueductal stenosis, and hematomas. Communicating hydrocephalus, however, may develop following meningitis, subarachnoid hemorrhage, or idiopathic causes such as normal pressure hydrocephalus, which allow a LP to be performed safely. The hallmark of hydrocephalus is increased volume of CSF associated with enlarged ventricles. Increased CSF volume often elevates ICP. A noninvasive method to measure ICP could significantly improve patient care.

Several groups have investigated the role of MR elastography (MRE), a noninvasive quantitative stiffness imaging technique capable of measuring the viscoelastic properties of tissues in vivo, for the diagnosis of
neurological diseases. The MRE approach requires an external source of vibration to generate shear waves within the tissue of interest. The displacement field generated by the shear waves is imaged with MR phase-contrast sequences. Finally, displacement fields are converted into quantitative stiffness maps through mathematical inversion techniques (1). MRE techniques estimate a map of the absolute shear modulus (stiffness) of the brain in kilopascal (kPa), which have shown to significantly change in brain tumors (2,3), normal pressure hydrocephalus (4,5), multiple sclerosis (6,7), and different types of dementia (8,9). Furthermore, groups have established baseline measurements of brain stiffness in healthy volunteer populations by investigating cross-sectional changes in brain viscoelasticity with respect to age and sex (10–12).

Recently, a study has reported that MRE stiffness is sensitive to changes due to altered venous drainage during jugular compression (13), which the authors suggest are due to neurovascular pressure changes in the brain. The authors were not able to report that jugular compression resulted in an increase in brain stiffness, but instead reported that volunteers who do not divert venous blood through extrajugular pathways have higher brain stiffness than those who do. The hypothesis of our study is that changing ICP alters the biomechanics or effective viscoelasticity of brain parenchyma, and brain stiffness may serve as a surrogate to ICP. Therefore, the objective of this study was to establish a swine model to investigate the relationship between acute changes in ICP and brain stiffness as measured with MRE.

METHODS

This study was approved by our Institutional Animal Care and Use Committee. Four domestic swine (Manthei Hog Farm, LLC, Rochester, Minnesota, USA) were used for the investigation of this study (35 ± 3 kg). The first animal was used to determine the optimal MRE baseline parameters without undergoing any surgery. In the other three animals, the relationship between stiffness and ICP was investigated by increasing ICP through two surgically implanted catheters while monitoring acute stiffness changes with MRE.

Presurgical MR Imaging

For surgical planning, a MRI-compatible stereotactic frame with a nine fiducial marker localizer box (14,15) was mechanically attached with screws to the porcine skull. This stereotactic frame subsequently was used as the platform to couple the source of vibration or driver to the skull to generate shear waves within the brain. With the frame attached, an anatomical T1-weighted 3D magnetization-prepared rapid gradient-echo sequence was obtained for surgical planning and catheter targeting. Imaging was performed on a 3T closed-bore MRI scanner (GE, Signa HDx, Waukesha, Wisconsin, USA) with the following imaging parameters: repetition time (TR) = 8 ms; echo time (TE) = 3.3 ms; slice thickness = 0.6 mm; number of excitations = 3; matrix size = 320 × 320; field of view (FOV) = 24 cm; and acquisition time = 6 min, with a custom-made four-channel receive-only surface coil placed on the top of the pig’s head within the stereotactic frame, as previously described (15).

Following anatomical imaging, and within the same scan session, MRE was performed to measure baseline brain stiffness presurgery at a range of vibration frequencies. MRE was conducted at 60 hertz (Hz), 90 Hz, 120 Hz, and 150 Hz vibration frequencies; and the image acquisition included a modified spin-echo echo-planar imaging sequence with the following parameters: TR/TE = 3,600 to 3,800/46 to 62 ms; FOV = 24 cm; matrix size = 72 × 72; 35 contiguous 3-mm-thick axial slices; vibration frequency matched motion-encoding gradient; motion-encoding in all three directions; and eight-phase offsets. MRE postprocessing involved: 1) applying a [5 × 5 × 5] quartic smoothing kernel (16), 2) calculating the curl of the displacement images, and 3) calculating the stiffness with a previously described amplitude squared weighted direct inversion algorithm (1,17), which can be described mathematically as:

\[ G^* = -\rho \omega^2 [\nabla^2 \mathbf{u}]^{-1}, \]

where \( G^* \) is the complex shear modulus; \( \rho \) is the tissue density (assumed to be 1,000 kg/m³); \( \omega \) is the angular velocity of the vibration; \( \mathbf{u} \) is the 3D (3 × 1) displacement vector of a single voxel; \( \nabla^2 \) is the 3D Laplace operator; and \( \mathbf{u}^\dagger \) is the pseudo inverse. This is derived from solving a system of simultaneous linear equations—one for each sensitization direction—for the quantity 1/\( \mu \), and is equivalent to least-squares fitting, assuming no noise in the displacement. It also is equivalent to amplitude-squared weighting of separate 1/\( \mu \) results for each sensitization direction. Stiffness over the entire brain volume was reported as the median magnitude of the complex shear modulus \( [G^*] \), where \( G^* \) is defined as a complex number with real part equal to the storage shear modulus \( \text{G}^* \) and imaginary part equal to the loss shear modulus \( \text{G}^0 \).

Surgical Catheter Placement

To simultaneously monitor and be able to intermittently increase ICP, silicon 1.5-mm lumen pediatric catheters (BACTISEAL EVD, Codman, Raynham, Massachusetts, USA) were implanted in the left and right lateral ventricles of the animals using a procedure similar to that previously described for an intraparenchymal drug-delivery system technique by MR image-guided stereotactic targeting (18). Briefly, this technique involves three steps: 1) an MRI-compatible stereotactic frame with a nine fiducial marker localizer box (14,15) was mechanically attached with screws to the porcine skull; 2) a high-resolution T1-weighted anatomical MRI of the pig brain with the stereotactic frame was acquired; 3) the DICOM image data was then transferred to a stereotactic planning computer for catheter projection planning.

Two burr holes were made through the skull for catheter placement. To secure a tight seal and minimize CSF leak, the burr holes were minimized to be ~8 mm, with an internal groove in the bone layer. Dental etch was used on top of the dry skull surface around the burr.
holes. Four to five titanium skull screws (3 mm) were used around the burr holes and etch layer. Then, a dental cement was used to secure the pediatric catheters to the skull (Fig. 1A). Unique to this study, a catheter was placed in both the left and right ventricles of the brain (Fig. 1B). Communication between the two catheters was confirmed during surgery by attaching a saline drip to one catheter and draining CSF from the other catheter. An inline pressure monitor was then connected to the drainage line to continuously monitor pressure throughout the experiment (Fig. 2). Prior to postsurgical imaging, the baseline was measured to be 0 millimeters mercury (mmHg) in the first two pigs and 28 mmHg in the third pig. In the third pig, free communication between the ventricles did not occur, which resulted in the higher baseline pressure. Sedation was maintained with 1.5% to 3% isoflurane during surgery and 1.5% to 1.75% isoflurane during the experiments. Vital signs continuously were monitored throughout the procedures.

**Postsurgical MR Imaging**

Following surgery, to measure brain stiffness as a function of ICP, the anesthetized animal with the stereotactic frame in place was repositioned within the MRI bore. The inline pressure monitor was kept outside the scanner room and connected by a drainage line that traveled through the waveguide of the scanner and attached to one of the pediatric catheters. The second catheter was connected to a saline bag kept inside the scanner room on an MRI-compatible intravenous (IV) pole next to the scanning table. ICP values were increased by raising the height of the saline bag with the extendable IV pole and monitoring the pressure values outside the scanner (Fig. 2). Each animal had three to four steps of increasing pressure. Imaging was conducted only after the pressure had stabilized for 1 min on the pressure monitor.

At each pressure level, and prior to any MRE acquisition, anatomical T₁-weighted images of the porcine brain were obtained to document any morphological changes.

**FIG. 1.** (A) Schematic diagram of surgical dental cement cap and catheter placement through the skull. (B-D) Catheter trajectory planning using high-resolution T₁-weighted images. (B) Oblique-coronal and (C) axial reconstructed image of pig brain, with the prescribed catheter placements shown in red. (D) Postsurgical axial reconstructed image after catheter placement. One catheter was used to increase ICP, whereas the other catheter was used to continuously monitor ICP. ICP, intracranial pressure.

**FIG. 2.** MRI schematic diagram of postsurgical MR elastography, imaging, and pressure-monitoring setup. Hz, hertz.
of the brain. A short, 4-min and 20-s inversion recovery–spoiled gradient recalled pulse sequence was used with the following parameters: slice thickness = 1.2 mm; matrix size = 256 × 256; FOV = 27 cm; sagittal acquisition plane; TR/TE = 7 ms/2.8 ms; flip angle = 11 degrees; and acceleration factor = 1.7, with a custom-made four-channel receive-only surface coil placed on the top of the pig’s head within the stereotactic frame. MRE was conducted postsurgery at the same frequencies as for presurgery for three (1 pig) and four (2 pigs) different ICP values ranging from 0 mmHg to 55 mmHg.

The same MRE imaging acquisition and postprocessing protocol was followed for the postsurgical scans, as described prior to surgery. A pairwise comparison on a voxel-by-voxel level was not possible due to ventricular deformations caused by catheter placement, as well as ventricular enlargement due to increased ICP. Therefore, to measure within-subject changes in brain stiffness with respect to ICP, the stiffness distribution of all voxels in the brain volume was compared between the postsurgical baseline and the states of elevated ICP using a one-sided Mann-Whitney U test of significance (19). Effect sizes were calculated for each comparison by calculating the $r$ value, as described by Fritz et al. (20), for which the guidelines for $r$ values are such that a large effect size is 0.5; a medium effect size is 0.3; and a small effect size is 0.1. To assess whether a linear correlation existed between brain stiffness and ICP across subjects, a multiple linear regression was applied to all the data, with pressure as a continuous independent variable, pig number as a nominal independent variable, and stiffness as the dependent variable. No significant correlation between stiffness and pig number was observed, but a significant correlation existed between stiffness and ICP. Therefore, another linear regression was applied comparing only the median brain stiffness and ICP across subjects at each frequency. A $P$ value of 0.05 was considered statistically significant.

RESULTS

Postsurgical Stiffness and Morphological Changes with Increased ICP

At baseline, prior to surgery, the median $|G^*|$ of the entire brain across all three pigs ranged from 1.58 to 1.92 kPa, 2.76 to 3.22 kPa, 4.09 to 4.74 kPa, and 5.70 to 5.8 kPa for frequencies of 60 Hz, 90 Hz, 120 Hz, and 150 Hz, respectively. Ventricle size increased progressively with stepwise increased ICP, as shown in Figure 3 for pig 1. At a pressure of 20 mmHg, an observable increase in only one ventricle is present (compare the initial ventricle size overlays in the bottom row of Fig. 3). At a pressure of 36 mmHg, both ventricles increased in size. Postsurgery, after the drainage of CSF and prior to elevating ICP, the median $|G^*|$ decreased in all three pigs. The decrease in $|G^*|$ across all three pigs was in the ranges of 0.015 to 0.025 kPa, 0.011 to 0.13 kPa, 0.20 to 0.42 kPa, and 0.71 to 0.78 kPa at vibration frequencies of 60 Hz, 90 Hz, 120 Hz, and 150 Hz, respectively. Ventricle size increased progressively with stepwise increased ICP, as shown in Figure 3 for pig 1. At a pressure of 20 mmHg, an observable increase in only one ventricle is present (compare the initial ventricle size overlays in the bottom row of Fig. 3). At a pressure of 36 mmHg, both ventricles increased in size. Postsurgery, after the drainage of CSF and prior to elevating ICP, the median $|G^*|$ decreased in all three pigs. The decrease in $|G^*|$ across all three pigs was in the ranges of 0.015 to 0.025 kPa, 0.011 to 0.13 kPa, 0.20 to 0.42 kPa, and 0.71 to 0.78 kPa at vibration frequencies of 60 Hz, 90 Hz, 120 Hz, and 150 Hz, respectively. In the third pig, good communication between the two catheters was not achieved. As a result, not as much CSF was drained from the ventricles of the third pig; however, the amount of CSF that was drained still produced a decrease in stiffness after surgery.

Within- and Cross-Subject Stiffness Changes with Increased ICP

The within-subject median $|G^*|$ values over the entire brain volume have been plotted in Figure 4 and listed in Table 1 as a function of ICP for vibration frequencies of 60 Hz, 90 Hz, 120 Hz, and 150 Hz. The $P$ values and corresponding effect sizes from the one-sided Mann-Whitney U

FIG. 3. Anatomical images of ventricles under different pressure levels postcatheter implantation in pig 1 (top row). The black dashed boxes in the top row have been magnified in the bottom row. For comparison purposes, the ventricles at 0 mmHg have been overlaid in red in all three magnified images. mmHg, millimeters mercury.
Figure 4, but no distinction has been made between pigs. For all vibration frequencies, a statistically significant linear correlation \((P \leq 0.01)\) was found between median brain stiffness and ICP. Sample elastograms illustrating the observed changes in brain stiffness over the span of the experiment for all three pigs are shown in Figure 6.

DISCUSSION

This study demonstrated both decreased ICP results in lowered brain stiffness and increased ICP results in increased brain stiffness within the same subject and across subjects in a swine model. Postsurgical baseline measurements with the associated drainage of CSF and accompanying lowered ICP were shown to decrease the median brain stiffness by 0.01 to 0.03 kPa, 0.01 to 0.12 kPa, 0.20 to 0.42 kPa, and 0.71 to 0.78 kPa at vibration frequencies of 60 Hz, 90 Hz, 120 Hz, and 150 Hz, respectively. A significant correlation between increased ICP and increased brain stiffness in the cross-subject comparison for all tested vibrational frequencies (60 Hz, 90 Hz, 120 Hz, and 150 Hz) was observed. The 120 Hz \((0.030 \pm 0.004 \text{ kPa/mmHg})\) and 150 Hz \((0.031 \pm 0.008 \text{ kPa/mmHg})\) vibrational frequencies had nearly identical slopes, which were approximately two- to three-fold higher than the 90 Hz \((0.017 \pm 0.002 \text{ kPa/mmHg})\) and 60 Hz \((0.009 \pm 0.002 \text{ kPa/mmHg})\) slopes, respectively. This suggests that, with current stiffness inversion methods, higher frequencies may be more sensitive to changes in pressure. Although this is a small sample population, it is encouraging to see that a linear relationship exists between absolute ICP and stiffness across subjects.

It appears that for pig 1 there was a pressure range from 0 to 20 mmHg in which brain stiffness did not increase with ICP (Fig. 4) (Table 1). Once past this range, the median brain stiffness for pig 1 showed a significant increase with increased ICP, which was especially noticeable at higher vibration frequencies. For the other two pigs, significant increases in stiffness were observed at all pressure levels tested. From a mechanical standpoint, increases in tissue stiffness result when a tissue is compressed such that the relationship between stress and strain becomes nonlinear, causing a rise in stiffness.

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pressure. These normal ICP values may be different than those observed in human subjects in whom ICP values greater than 15 mmHg sometimes can be considered above average (21). Future studies in humans will be needed to investigate this.

It may appear from the elastograms in Figure 6 that the distribution of brain stiffness is not uniform throughout the entire brain, which may suggest that only regions of tissue are increasing in stiffness with acute changes in pressure, and that these do not become significant enough to affect the entire brain volume until higher ICP values are reached. However, the linear correlation between median stiffness and ICP suggests that this is a global phenomenon because it is affecting the entire distribution. It is not within the scope of this paper to evaluate if MRE is capable of measuring localized regions of increased pressure in this animal model. However, future studies could help support or disprove the hypothesis that some forms of hydrocephalus result from localized CSF accumulation, leading to local compression in certain regions in the brain (5,22).

To the knowledge of the authors, this is the first study to investigate the relationship of $G^*$ and absolute ICP in vivo. Hatt et al. (13) performed brain MRE on nine healthy normal volunteers with and without jugular compression. They acquired a single slice in the brain using a vibration frequency of 30 Hz, and were not able to report a significant difference in brain stiffness between these two conditions. Pattison et al. (23), building on Shapiro et al.'s (24) feline hydrocephalus model, performed MRE at a vibration frequency of 85 Hz in 18 female feline subjects, 12 of which received kaolin injections to induce an acute form of hydrocephalus. Pattison et al. did not report ICP values, and no formal statistics are reported on their stiffness results; however, they did report a mean increase in parenchymal stiffness from 5.4 ± 2.8 kPa to 9.3 ± 5.1 kPa between the baseline values and the hydrocephalic state, respectively. Mousavi et al. (25), with a strain MRI-imaging approach, demonstrated that volumetric strain correlated with whole-brain dilation in five healthy volunteers by using the Valsalva maneuver to alter ICP. Hirsch et al. (26) used MRE and phase-contrast MRI measurements to show that volumetric strain in the brain could be increased by 45% with abdominal muscle contractions. Lastly, Weaver et al. (27), with MRE and a poroelastic model, showed that fluid pressure carries a portion of the stress that contributes to the mean shear modulus in the brain. However, because ICP was not recorded during any of these studies, it currently is not possible to compare these to our results. In addition, previous studies conducted in the liver and spleen have reported supporting evidence that a direct correlation with MRE stiffness and different levels of pressure and static strain does exist (28,29).

An effect size is considered small, medium, or large when $r$ equals 0.1, 0.3, and 0.5, respectively.

Hz, hertz; mmHg, millimeters mercury; not acquired, NA.
patients with chronically elevated ICP. Some of the advantages of MRE over the described technique are that MRE is capable of producing localized stiffness measurements, does not depend on the accuracy of lumen segmentation, and is an independent measurement of the tissue’s response to pressure and not a direct derivative of the pressure in the CSF itself. Several other groups also have used invasive pressure-volume measurements, with the addition of different aliquots of artificial CSF, to estimate an elasticity slope from elastance measurements. These techniques have been investigated in a series of animal experiments (31–33) and in patient populations (34,35) in various disorders of the CSF system. Again, these measurements are made on the response of the lumen and not necessarily the underlying tissue mechanics within the entire brain. Although the elastance index is a different metric than $G^*$, these results give support that elasticity may be a valuable biomarker for disease differentiation and motivate further exploration of MRE in human subjects.

The limitations of this study were the small sample size, the measurement of only acute changes in ICP versus chronic ICP elevations, and low 3-mm isotropic MRE acquisition resolution. In addition, proper drainage of CSF was not possible in pig 3, resulting in a baseline ICP of 26 mmHg instead of 0 mmHg. However, similar responses in stiffness still were observed, both directly after surgery and with elevations in ICP in pig 3. Although the number of subjects used was small, the experiment was designed to elicit a graded response within each animal, with multiple within-animal correlations being acquired. Thus, the number of data points was larger than the number of animals used, improving statistical power. We previously also have shown, despite small sample sizes, that consistent results with high statistical power can be achieved in stereotactic swine surgeries (36–38). Lastly, the objectives of this study were to demonstrate the feasibility of measuring stiffness in vivo, while simultaneously increasing and monitoring ICP through two implanted catheters in the left and right ventricles of the

| Vibration Frequency | 0 mmHg Intercept (kPa) | Stiffness/Pressure Slope (kPa/mmHg) | $P$ Value | R Squared |
|---------------------|------------------------|------------------------------------|-----------|-----------|
| 60 Hz               | 1.55 ± 0.06            | 0.009 ± 0.002                      | 0.001     | 0.74      |
| 90 Hz               | 2.62 ± 0.08            | 0.017 ± 0.002                      | < .0001   | 0.86      |
| 120 Hz              | 3.62 ± 0.15            | 0.030 ± 0.004                      | < .0001   | 0.85      |
| 150 Hz              | 4.83 ± 0.24            | 0.031 ± 0.008                      | 0.01      | 0.73      |

Hz, hertz; kPA, kilopascal; mmHg, millimeters mercury.

FIG. 6. Elastograms ($G^*$) of presurgery and postsurgery at baseline and postsurgery at maximum ICP in three pigs and at a 120 hertz vibration frequency. ICP, intracranial pressure; kPA, kilopascal; mmHg, millimeters mercury.
CONCLUSION

MRE demonstrated a direct relationship between brain stiffness and ICP across the entire brain volume. A statistically significant increase in stiffness was observed both within and across subjects in response to elevations in ICP. This study motivates future investigation in human subjects to determine if MRE has a role in the diagnosis and management of patients with changes in ICP and hydrocephalus.

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