Test-retest variability of brain morphometry analysis: an investigation of sequence and coil effects

Shuang Yan¹, Tianyi Qian², Bénédicte Maréchal³,⁴,⁵, Tobias Kober³,⁴,⁵, Xianchang Zhang², Jinxia Zhu², Jing Lei¹, Mingli Li¹, Zhengyu Jin¹

¹Department of Radiology, Peking Union Medical College Hospital, Peking Union Medical College, Chinese Academy of Medical Sciences, Beijing 100730, China; ²Department of MR Collaboration, Siemens Healthcare Ltd., Beijing 100102, China; ³Department of Advanced Clinical Imaging Technology, Siemens Healthcare, Lausanne, Switzerland; ⁴Department of Radiology, Lausanne University Hospital and University of Lausanne, Lausanne, Switzerland; ⁵LTS5, École Polytechnique Fédérale de Lausanne (EPFL), Lausanne, Switzerland

Contributions: (I) Conception and design: S Yan, T Qian; (II) Administrative support: Z Jin; (III) Provision of study materials or patients: S Yan, T Qian, M Li; (IV) Collection and assembly of data: J Lei, M Li, S Yan; (V) Data analysis and interpretation: B Maréchal, T Kober, X Zhang, J Zhu; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

Correspondence to: Zhengyu Jin, MD. Department of Radiology, Peking Union Medical College Hospital, Peking Union Medical College, Chinese Academy of Medical Sciences, No. 1 Shuaifuyuan, Dongcheng District, Beijing 100730, China. Email: jinzy@pumch.cn.

Background: Precise and reliable brain morphometry analysis is critical for clinical and research purposes. The magnetization-prepared rapid gradient echo (MPRAGE), multi-echo MPRAGE (MEMPRAGE) and magnetization-prepared 2 rapid acquisition gradient echo (MP2RAGE) sequences have all been used to acquire brain structural images, but it is unclear which of these sequences is the most suitable for brain morphometry and whether the number of coil channels (20 or 32) affects scan precision. This study aimed to assess the impact of T1-weighted image acquisition variables (sequence and head coil) on the repeatability of resultant automated volumetric measurements.

Methods: Twenty-four healthy volunteers underwent back-to-back scanning protocols with three sequences and two different coils (i.e., six scanning conditions in total) presented in a randomized order in a single session. MorphoBox prototype and FreeSurfer were used for brain segmentation. Brain structures were divided into cortical and subcortical regions for more precise analysis. The acquired volume and thickness values were used to calculate test-retest variability (TRV) values. TRV values from the six different combinations were compared for total brain structures, total cortical structures, total subcortical structures, and every single structure.

Results: The median TRV value for all brain regions was 1.23% with MorphoBox and 3.14% with FreeSurfer. When using FreeSurfer results to compare the six combinations, for total brain structures volume and total cortical structures volume and thickness, the MEMPRAGE-32 channel combination showed significantly lower TRV values than the others (P<0.01). Similar results were observed with MorphoBox. For total subcortical structures, the MP2RAGE-32 channel combination showed the lowest TRV values with both MorphoBox (lower about 0.01% to 0.17%) and FreeSurfer analyses (lower about 0.02% to 0.37%).

Conclusions: TRV values were generally low, indicating generally high reliability for every region. The MEMPRAGE sequence was the most reliable of the three sequences for total brain structures and cortical structures. However, MP2RAGE was the most reliable for subcortical structures. The 32-channel coil showed better repeatability results than the 20-channel coil.

Keywords: Test-retest variability (TRV); magnetic resonance imaging (MRI) repeatability; brain morphometry; post-processing robustness

Submitted Sep 16, 2019. Accepted for publication Nov 06, 2019.
doi: 10.21037/atm.2019.11.149

View this article at: http://dx.doi.org/10.21037/atm.2019.11.149
Introduction

The precise and robust visualization and delineation of brain structures are crucial for both clinical and research purposes (1-3). Automated morphometric analysis, based on high-resolution structural magnetic resonance imaging (MRI) data, has been widely used in cross-sectional and longitudinal research studies to visualize and delineate brain structures. The sensitivity of morphometry to detect changes in these studies depends on the reliability of the analysis (4). However, test-retest variability (TRV) can be affected by changes in pulse sequences and radio-frequency (RF) coils (5,6).

Several sequences can be used to acquire high-resolution 3D T1-weighted brain structural images which can then be used in morphometric analyses. One of the most widely used pulse sequences is the magnetization-prepared rapid gradient echo (MPRAGE) sequence (7), which is considered to be the standard sequence for structural brain imaging. This sequence can produce excellent T1-weighted contrast between gray matter (GM), white matter (WM), and cerebrospinal fluid (CSF), and delivers good overall image quality. MPRAGE sequence commonly consists of a magnetization preparation (MP) period, which is denoted as an inversion time (TI) after an inversion pulse, and a rapid gradient-echo (RAGE) readout, which is a FLASH (8) or a FISP (9) module. However, this sequence’s image quality is sensitive to the inhomogeneity of the transmit (B1+) and receive (B1−) RF fields (10). The inhomogeneity and distortion of the main magnetic (B0) field may also influence image quality (11).

Recently, new sequences have been developed based on the conventional MPRAGE sequence. One such sequence is the multi-echo MPRAGE (MEMPRAGE) sequence. A multi-echo sequence usually consists of four echoes with higher bandwidths instead of a single echo sequence that has a lower bandwidth. The higher bandwidths can help decrease B0-related distortions and enhance the reproducibility of morphometry measurements without sacrificing the signal-to-noise ratio (SNR), as the different echoes are typically combined. The MEMPRAGE sequence is also better able to segment anatomy from the dura to the cortex, resulting in structures with fine edges that allow for precise registration across contrasts (11,12), taking advantage of the largely different T2* characteristics of these tissues.

The magnetization-prepared 2 rapid acquisition gradient echo (MP2RAGE) sequence (13) is another such sequence. This sequence applies a time-efficient measure that can significantly lessen image biases and help prevent the impact of inhomogeneity induced by the reception and transmit RF fields. The sequence introduces the RAGE block twice and collects two full image sets during each of the two different TIs. The scan parameters are optimized so that one image can be synthesized by combining the two image sets and can provide eminent contrast across different brain tissues. The T1-weighted image, in which the cortical GM is efficiently nulled at the center of the k-space, is made by the first TI (TI1); the second TI (TI2) is long, thus it can approximately generate a proton density weighted contrast. Consensus has yet to be about which of these high-resolution 3D T1-weighted sequences is the most reliable and reproducible. Furthermore, few previous studies have compared these three sequences with each other.

Brain image quality is also influenced by the number of receiver channels in the utilized head coil. A phased-array coil with more channels can permit accelerated imaging and can significantly improve the SNR throughout the brain, especially in the cortex (14-16). However, it is unclear whether the number of coil channels affects scan and segmentation robustness.

The objectives of this study are to compare the repeatability of morphometric measures derived from the three aforementioned sequences, to investigate whether the results are consistent among different brain regions, such as between the cortical structures and subcortical structures, and to determine if RF coils impact the precision of the MRI scan and derived morphometric analysis.

Methods

Participants

The Institutional Review Board of our hospital approved this study, and informed consent was obtained from all the participants. Twenty-four healthy volunteers (12 males) with no history of severe head trauma, brain tumors, or neurological or psychiatric disorders were recruited. The mean age of the volunteers was 27.5 (range from 22 to 30) years old.

Imaging acquisition

Whole-brain images were acquired on a 3T MR scanner (MAGNETOM Skyra, Siemens Healthcare, Erlangen, Germany). Foam padding was used to help reduce head motion, and earplugs were used to attenuate noise. The acquisition protocol included three high-resolution 3D T1-
weighted sequences; scan parameters are shown in Table 1. All participants underwent scanning with all sequences, with both a 20-channel head-neck coil and a 32-channel head coil. Therefore, there were six experimental conditions: MPRAGE-20 channel, MPRAGE-32 channel, MEMPRAGE-20 channel, MEMPRAGE-32 channel, MP2RAGE-20 channel, and MP2RAGE-32 channel. The participants were assigned to undergo the six sequence-coil combinations in a randomized order. A rescan of each sequence-coil combination was performed immediately after the first scan, so that scan repeatability could be evaluated.

### Image processing

Both the MorphoBox prototype and FreeSurfer were used to automatically segment brain structures and measure brain region volumes.

#### MorphoBox

The MorphoBox prototype is a fast-inline volume-based morphometry algorithm (2,17-20). This algorithm is integrated into the image reconstruction chain of all three T1-weighted sequences so that brain structure volumes can be obtained directly in the scanner. It takes as input either a 3D MPRAGE, the result of root-mean-square combination of all four echoes in case of MEMPRAGE acquisition, or both the second inversion contrast and uniform original image in case of MP2RAGE acquisition. It combines atlas-free tissue classification and atlas-based brain segmentation to estimate the volume of 45 regions in approximately 2 minutes. Regions are further referred to as tissue [i.e., total intracranial volume (TIV), GM, WM, and CSF] or structures [i.e., subcortical nuclei, such as the thalamus, putamen, caudate, pallidum, and hippocampus; the GM and WM of the frontal, parietal, occipital, and temporal lobes; the ventricular system; the corpus callosum; the cerebellum; and the brainstem]. Additionally, we gathered regions into cortical and subcortical structures based on their anatomical position for further detailed comparisons.

#### FreeSurfer

FreeSurfer v5.3.0 (http://surfer.nmr.mgh.harvard.edu) (21) measures cortical thickness using a surface-based model and measures subcortical volume using a volume-based model. Detailed descriptions of the segmentation procedure have been reported in previous studies (22,23). Briefly, the default processing stream (recon-all) and the segmentation consist of the following steps: registering the images to a template, normalizing the intensity, skull stripping the images, segmenting the structures, and statistical analyses.

A total of 148 cortical structure regions, 45 subcortical structures, and 70 WM structures were automatically segmented; their volumes were calculated using FreeSurfer. Both the volume and thickness values of the cortical structure regions were acquired.

### Statistical analysis

TRV was used to evaluate the reliability of the measurements from both MorphoBox and FreeSurfer for each brain region generated by each sequence and coil combination. Lower TRV values indicate lower variability and higher repeatability. TRV was calculated using the following equation: \( \text{TRV} = \frac{|M1-M2| \times 2 \times 100\%}{(M1+M2)} \) (24), which was defined as the absolute difference divided by the mean of two measurements. M1 and M2 indicate the first (test) and the second (retest) measurements, respectively.
TRV values of all brain regions were analyzed together as TRV for total brain structures, TRV values of all cortical regions were analyzed together as TRV for total cortical structures, and TRV values of all the subcortical regions were analyzed together as TRV for total subcortical structures. TRV values for every single structure, total brain structures, total cortical structures, and total subcortical structures were then compared among the six combinations using the Friedman test. Differences between each pair of combinations were measured using Student-Newman-Keuls post hoc analysis. All P values were 2-tailed. P<0.05 was considered to be statistically significant. All statistical analyses were performed using SPSS20 (IBM, Armonk, NY, USA).

Results

Structural images acquired with different combinations of sequences and coils are displayed in Figure 1.

MorphoBox-based volumetry repeatability comparison

The median TRV calculated by the data acquired by MorphoBox was 1.23% (range, 0.00–9.89%) for all brain regions. Comparisons of the six different combinations and significant differences between the median TRV values of total brain structures, subcortical structures, brain tissue (TIV, GM, WM, CSF, cortical GM), and structural regions are shown in Table 2. All values are given in the form of median (first quartile, third quartile). TRV values were not significantly different between the different coil/sequence combinations for total brain structures and total subcortical structures. The median TRV value of the MEMPRAGE-32 channel combination for total brain structures was 1.10% (0.31%, 2.37%) and was slightly lower than that of other combinations. The MEMPRAGE sequence had the lowest TRV values among the three sequences for total brain structures, regardless of which coil was used. The MP2RAGE-32 channel combination had the lowest median TRV value [1.16% (0.21%, 2.11%)] for total subcortical structures. The TRV values of the MP2RAGE sequence were numerically lower than those for total subcortical structures regardless of which type of coil was used (lower 0.01% to 0.17%). The TRV values of the 32-channel coil were lower than those of the 20-channel coil, regardless of sequence, for both total brain structures and total subcortical structures.

Regarding single tissues and structures, the TRV values were only significantly different between the six combinations for the TIV, WM, left deep WM, and the left insular and right cingulum GM. For the TIV, the TRV of the MEMPRAGE-32 channel combination was the lowest [0.10% (0.07%, 0.24%)]. The following TRV values for the other combinations are listed here in ascending order: MPRAGE-32 channel [0.17% (0.05%, 0.23%)], MEMPRAGE-20 channel [0.20% (0.07%, 0.30%)], MPRAGE-20 channel [0.21% (0.05%, 0.33%)], MP2RAGE-32 channel [1.69% (1.00%, 3.23%)], and MP2RAGE-20 channel [2.54% (1.49%, 3.89%)]; the TRV values of the final two combinations were significantly higher than those of the other combinations. The TRV values for the MP2RAGE-32 channel and MP2RAGE-20 channel combinations were also significantly higher than those of the other four combinations for WM. However, these two combinations had significantly lower TRV values.
than those of the other four combinations for the left insular and right cingulum GM regions and the left deep WM regions. The MEMPRAGE-32 channel combination had the lowest TRV values in approximately 33.3% of all brain regions that had comparable TRV values across the six combinations; no other combination yielded optimal TRV values at a higher rate. The MP2RAGE-32 channel had the lowest TRV values among 30.8% of single regions within the subcortex; no other combination yielded optimal TRV values at a higher rate.

### FreeSurfer-based volumetry repeatability comparison

The median TRV value was 3.14% and all TRV values ranged from 0.00% to 16.46% in the data set generated by FreeSurfer. Comparisons and significant differences between TRV values for the total brain structures, total subcortical structures, and single subcortical structure are presented in Table 3. The MEMPRAGE-32 channel combination [2.83% (1.18%, 5.96%)] yielded the lowest TRV values, which were significantly lower than those of the MPRAGE-32 channel combination [3.01% (1.32%, 6.20%)] and the MEMPRAGE-20 channel combination [3.06% (1.31%, 6.40%)]. The TRV values for the MPRAGE-20 channel combination [3.24% (1.37%, 6.76%)] and MP2RAGE-32 channel combination [3.30% (1.41%, 6.91%)] were significantly higher than the values of the three aforementioned combinations but were significantly lower than the values of the MP2RAGE-20 channel combination [3.38% (1.48%, 7.02%)]. There were no significant differences in TRV values for the total subcortical structures across the six combinations, but the MP2RAGE-32 channel combination [2.29% (1.01%, 5.50%)] yielded the lowest TRV of all the combinations (lower about 0.02% to 0.37%). Regardless of coil, the TRV values for the MP2RAGE sequence were lower than those of the total subcortical structures. Additionally, regardless of sequence, the TRV values of the 32-channel coil were lower than those of the 20-channel coil. There were no significant differences between the six combinations for all the single subcortical and WM structures except for the left thalamus, in which the TRV for MP2RAGE-20 channel condition [1.12% (0.25%, 2.74%)] was significantly lower than that of other conditions.

### Table 2 TRV values of the six combinations with MorphoBox

<table>
<thead>
<tr>
<th>Brain structures</th>
<th>MPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total brain structures</td>
<td>1.33 (0.36, 2.67)</td>
<td>1.16 (0.35, 2.82)</td>
<td>1.17 (0.36, 2.44)</td>
<td>1.10 (0.31, 2.37)</td>
<td>1.41 (0.48, 2.82)</td>
<td>1.20 (0.37, 2.27)</td>
<td>0.328</td>
</tr>
<tr>
<td>Total subcortical structures</td>
<td>1.33 (0.22, 2.87)</td>
<td>1.20 (0.00, 2.66)</td>
<td>1.29 (0.00, 2.68)</td>
<td>1.17 (0.00, 2.67)</td>
<td>1.26 (0.30, 2.93)</td>
<td>1.16 (0.21, 2.11)</td>
<td>0.376</td>
</tr>
<tr>
<td>TIV</td>
<td>0.21 (0.05, 0.33)</td>
<td>0.17 (0.05, 0.23)</td>
<td>0.20 (0.07, 0.30)</td>
<td>0.10 (0.07, 0.24)</td>
<td>2.54** (1.49, 3.89)</td>
<td>1.69** (1.00, 3.23)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>GM</td>
<td>0.61 (0.31, 1.13)</td>
<td>1.63 (0.42, 3.09)</td>
<td>1.03 (0.26, 1.79)</td>
<td>1.07 (0.30, 1.98)</td>
<td>1.09 (0.00, 2.95)</td>
<td>1.02 (0.20, 2.04)</td>
<td>0.845</td>
</tr>
<tr>
<td>WM</td>
<td>0.48 (0.20, 1.23)</td>
<td>0.76 (0.52, 1.80)</td>
<td>0.24 (0.13, 1.24)</td>
<td>0.57 (0.16, 0.81)</td>
<td>1.41** (0.93, 4.05)</td>
<td>1.93** (1.03, 5.14)</td>
<td>0.005</td>
</tr>
<tr>
<td>CSF</td>
<td>1.26 (0.34, 4.85)</td>
<td>2.86 (1.88, 7.25)</td>
<td>1.23 (0.67, 3.30)</td>
<td>2.15 (0.83, 4.87)</td>
<td>2.40 (0.26, 3.66)</td>
<td>2.08 (1.16, 3.38)</td>
<td>0.581</td>
</tr>
<tr>
<td>Cortical GM</td>
<td>0.52 (0.33, 1.17)</td>
<td>1.67 (0.22, 3.34)</td>
<td>1.00 (0.12, 1.99)</td>
<td>1.04 (0.50, 2.24)</td>
<td>0.56 (0.10, 0.82)</td>
<td>0.43 (0.24, 1.21)</td>
<td>0.279</td>
</tr>
<tr>
<td>L insular</td>
<td>1.44 (1.24, 3.85)</td>
<td>1.49 (1.33, 2.65)</td>
<td>2.14 (1.28, 5.22)</td>
<td>2.65 (1.48, 4.61)</td>
<td>0.97* (0.39, 1.66)</td>
<td>1.06* (0.64, 1.84)</td>
<td>0.009</td>
</tr>
<tr>
<td>R cingulum GM</td>
<td>3.05 (1.38, 4.36)</td>
<td>2.90 (1.11, 4.92)</td>
<td>1.99 (1.55, 3.73)</td>
<td>2.05 (1.33, 3.11)</td>
<td>0.70* (0.61, 1.23)</td>
<td>1.17* (0.13, 2.45)</td>
<td>0.049</td>
</tr>
<tr>
<td>L deep WM</td>
<td>2.09 (0.74, 3.33)</td>
<td>0.97 (0.00, 2.94)</td>
<td>2.16 (1.14, 3.00)</td>
<td>1.71 (0.26, 2.72)</td>
<td>0.82* (0.35, 1.34)</td>
<td>0.67* (0.21, 0.98)</td>
<td>0.034</td>
</tr>
</tbody>
</table>

*, means the TRV values of this combination were significantly lower than those of other combinations; **, means the TRV values of this combination were significantly higher than those if other combinations. TRV, test-retest variability; MPRAGE, magnetization-prepared rapid gradient echo; MEMPRAGE, multi-echo MPRAGE; MP2RAGE, magnetization-prepared 2 rapid acquisition gradient echo; CH, channel; Q1, first quartile; Q3, third quartile; TIV, total intracranial volume; GM, gray matter; WM, white matter; CSF, cerebrospinal fluid; L, left; R, right.
structures and every single cortical structure across the six combinations are displayed in Table 4. Analyzed with volume data, the TRV values for total cortical structures were significantly different between the six combinations. The TRV for the MEMPRAGE-32 channel combination [3.43% (1.58%, 6.81%)] was the lowest. The TRV values for the MPRAGE-32 channel [3.70% (1.69%, 7.04%)] and MEMPRAGE-20 channel [3.83% (1.74%, 7.75%)] combinations were significantly higher than the values for the MEMPRAGE-32 channel combination and were significantly lower than the values for the MPRAGE-32 channel [4.18% (1.83%, 7.94%)], MP2RAGE-32 channel [4.30% (1.92%, 8.28%)], and MP2RAGE-20 channel [4.32% (1.99%, 8.40%)] combinations. The MPRAGE-32 channel combination had significantly lower TRV values than the other combinations, and the MP2RAGE-32 channel combination had significantly higher TRV values than the other combinations in the left temporal middle gyrus and right temporal middle gyrus. The TRV for the MP2RAGE-32 channel combination was significantly lower than that for other combinations, and the TRV for the MPRAGE-20 channel combination was significantly higher in the anterior left cingulum than that for other combinations. The MPRAGE-32 channel combination and the MEMPRAGE-32 channel combination had the lowest numerical TRV values in 25.9%, 35.3% and 40.0% of all brain regions, cortical structures and subcortical structures, respectively, which had comparable TRV values; and no other combinations yielded optimal TRV values at a higher rate.

The TRV values for thickness differed significantly among the different combinations. All combinations were significantly different from each other for total cortical structures; the TRV values in ascending order were: MEMPRAGE-32 channel [1.65% (0.73%, 3.02%)], MPRAGE-32 channel [1.73% (0.82%, 3.33%)], MPRAGE-20 channel [1.83% (0.84%, 3.40%)], MP2RAGE-32 channel [2.03% (0.93%, 3.84%)], MP2RAGE-20 channel [2.08% (0.92%, 3.83%)], and MP2RAGE-20 channel [2.30% (1.02%, 4.28%)]. Regarding single cortical structures, the TRV values for the MEMPRAGE-32 channel combination were significantly lower than those of the other combinations, and the TRV values for the MP2RAGE-20 channel combination were significantly higher than those in the right front middle gyrus, left temporal inferior gyrus, left parietal inferior supramarginal gyrus, and right parietal inferior angular gyrus. The MPRAGE-32 channel combination had TRV values that were significantly lower than those of the other combinations, and the MP2RAGE-20 channel combination had TRV values that were significantly higher than the other combinations in the left front inferior triangular gyrus and the left parietal inferior angular gyrus. The MPRAGE-32 channel combination had TRV values that were significantly lower than those of the other combinations, and the MP2RAGE-20 channel combination had TRV values that were significantly higher than the other combinations in the left front inferior triangular gyrus and the left parietal inferior angular gyrus. The TRV values for the MPRAGE-32 channel combination and the MEMPRAGE-32 channel combination were significantly lower than the values for the other combinations in the left temporal middle gyrus and right occipital middle gyrus; the TRV value for the MP2RAGE-32 channel combination was the highest in the left temporal middle gyrus; the TRV value for the MP2RAGE-20 channel combination was the

<table>
<thead>
<tr>
<th>Brain structures</th>
<th>MPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total brain</td>
<td>3.24 (1.37, 6.76)</td>
<td>3.01 (1.32, 6.20)</td>
<td>3.06 (1.31, 6.40)</td>
<td>2.83* (1.18, 5.96)</td>
<td>3.38** (1.48, 7.02)</td>
<td>3.30 (1.41, 6.91)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Total subcortical</td>
<td>2.66 (0.97, 5.60)</td>
<td>2.48 (0.91, 6.02)</td>
<td>2.45 (0.94, 5.51)</td>
<td>2.31 (0.92, 6.21)</td>
<td>2.44 (0.99, 5.82)</td>
<td>2.29 (1.01, 5.50)</td>
<td>0.514</td>
</tr>
<tr>
<td>CSF</td>
<td>4.36 (2.23, 6.57)</td>
<td>2.52 (6.44, 10.12)</td>
<td>4.95 (1.71, 7.81)</td>
<td>3.56 (2.45, 6.24)</td>
<td>2.53 (1.30, 8.22)</td>
<td>4.23 (2.63, 9.17)</td>
<td>0.686</td>
</tr>
<tr>
<td>L thalamus</td>
<td>2.60 (0.74, 5.07)</td>
<td>2.33 (1.01, 6.09)</td>
<td>2.34 (1.09, 6.44)</td>
<td>4.18 (2.17, 7.44)</td>
<td>1.12* (0.25, 2.74)</td>
<td>1.98 (0.80, 3.59)</td>
<td>0.022</td>
</tr>
</tbody>
</table>

*, means the TRV values of this combination were significantly lower than those of other combinations; **, means the TRV values of this combination were significantly higher than those of other combinations. TRV, test-retest variability; MPRAGE, magnetization-prepared rapid gradient echo; MEMPRAGE, multi-echo MPRAGE; MP2RAGE, magnetization-prepared 2 rapid acquisition gradient echo; CH, channel; Q1, first quartile; Q3, third quartile; CSF, cerebrospinal fluid; L, left.
<table>
<thead>
<tr>
<th>Brain structures</th>
<th>MPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MEMPRAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_20CH TRV (%), median (Q1, Q3)</th>
<th>MP2RAGE_32CH TRV (%), median (Q1, Q3)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cortical structures volume</td>
<td>4.18** (1.83, 7.94)</td>
<td>3.70 (1.69, 7.04)</td>
<td>3.83 (1.74, 7.75)</td>
<td>3.43* (1.58, 6.81)</td>
<td>4.32** (1.99, 8.40)</td>
<td>4.30** (1.92, 8.28)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>L temporal middle (V)</td>
<td>2.72 (1.01, 4.86)</td>
<td>1.62* (0.41, 3.31)</td>
<td>2.73 (0.79, 3.55)</td>
<td>2.32 (1.05, 4.37)</td>
<td>2.77 (1.27, 4.58)</td>
<td>4.74** (1.69, 8.09)</td>
<td>0.017</td>
</tr>
<tr>
<td>R temporal middle (V)</td>
<td>1.87 (0.98, 3.72)</td>
<td>1.64* (0.76, 2.32)</td>
<td>1.96 (1.04, 2.59)</td>
<td>1.96 (0.95, 2.77)</td>
<td>2.01(0.38, 4.42)</td>
<td>2.62** (1.76, 7.68)</td>
<td>0.043</td>
</tr>
<tr>
<td>L cingulum anterior (V)</td>
<td>5.65** (1.61, 7.07)</td>
<td>3.34 (1.76, 4.36)</td>
<td>2.06 (0.74, 4.25)</td>
<td>3.45 (2.06, 5.44)</td>
<td>4.50 (2.66, 8.12)</td>
<td>2.00* (1.35, 3.86)</td>
<td>0.035</td>
</tr>
<tr>
<td>Total cortical structures thickness</td>
<td>2.08 (0.92, 3.83)</td>
<td>1.73 (0.82, 3.33)</td>
<td>1.83 (0.84, 3.40)</td>
<td>1.65* (0.73, 3.02)</td>
<td>2.30** (1.02, 4.28)</td>
<td>2.03 (0.93, 3.84)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>L front inferior triangular (T)</td>
<td>2.07 (1.32, 3.67)</td>
<td>0.87* (0.33, 1.53)</td>
<td>2.12 (0.74, 3.29)</td>
<td>1.35 (0.55, 3.39)</td>
<td>2.31** (1.06, 4.37)</td>
<td>2.31** (0.89, 3.99)</td>
<td>0.011</td>
</tr>
<tr>
<td>R front middle (T)</td>
<td>1.55 (0.64, 3.19)</td>
<td>1.38 (0.66, 2.26)</td>
<td>0.98 (0.41, 2.01)</td>
<td>0.93* (0.31, 2.09)</td>
<td>1.75** (0.98, 3.10)</td>
<td>1.33 (0.41, 2.32)</td>
<td>0.040</td>
</tr>
<tr>
<td>L temporal middle (T)</td>
<td>1.57 (0.99, 2.09)</td>
<td>0.98* (0.16, 1.76)</td>
<td>1.42 (0.62, 2.35)</td>
<td>1.36* (0.49, 2.12)</td>
<td>1.56 (0.77, 3.45)</td>
<td>3.01** (1.35, 5.83)</td>
<td>0.010</td>
</tr>
<tr>
<td>L temporal inferior (T)</td>
<td>1.87 (0.90, 2.71)</td>
<td>1.59 (0.51, 2.40)</td>
<td>1.17 (0.62, 2.07)</td>
<td>0.87* (0.33, 1.48)</td>
<td>1.93** (1.31, 4.13)</td>
<td>1.86 (0.53, 3.44)</td>
<td>0.026</td>
</tr>
<tr>
<td>L parietal inferior supramarginal (T)</td>
<td>1.49 (0.83, 2.46)</td>
<td>1.38 (0.64, 2.58)</td>
<td>1.26 (0.35, 2.22)</td>
<td>0.71* (0.17, 2.36)</td>
<td>2.43** (1.45, 3.87)</td>
<td>1.34 (0.89, 2.30)</td>
<td>0.024</td>
</tr>
<tr>
<td>L parietal inferior angular (T)</td>
<td>1.77 (0.84, 3.86)</td>
<td>1.05* (0.49, 1.87)</td>
<td>1.28 (0.58, 2.78)</td>
<td>1.53 (0.57, 3.04)</td>
<td>2.50** (1.47, 4.79)</td>
<td>1.65 (0.83, 3.42)</td>
<td>0.044</td>
</tr>
<tr>
<td>R parietal inferior angular (T)</td>
<td>1.80 (0.86, 3.80)</td>
<td>1.43 (0.30, 2.98)</td>
<td>1.26* (0.48, 2.59)</td>
<td>1.11* (0.41, 2.28)</td>
<td>3.14** (1.43, 4.74)</td>
<td>1.99 (0.59, 3.51)</td>
<td>0.005</td>
</tr>
<tr>
<td>R occipital middle (T)</td>
<td>2.31 (0.39, 3.51)</td>
<td>1.09* (0.90, 1.91)</td>
<td>1.19 (0.60, 2.63)</td>
<td>1.15* (0.32, 1.92)</td>
<td>2.85** (1.71, 4.58)</td>
<td>1.77 (0.58, 3.32)</td>
<td>0.018</td>
</tr>
</tbody>
</table>

*, means the TRV values of this combination were significantly lower than those of other combinations; **, means the TRV values of this combination were significantly higher than those of other combinations. TRV, test-retest variability; MPRAGE, magnetization-prepared rapid gradient echo; MEMPRAGE, multi-echo MPRAGE; MP2RAGE, magnetization-prepared 2 rapid acquisition gradient echo; CH, channel; Q1, first quartile; Q3, third quartile; R, right; L, left; V, volume; T, thickness.
highest in the right occipital middle gyrus.

Discussion

The main goal of this research was to investigate which 3D T1-weighted sequences yielded the best scan and segmentation repeatability for brain structures. The TRV values for the MEMPRAGE-32 channel combination were the lowest for the total brain structures and total cortical structures. When using FreeSurfer segmentation, the TRV values for the MEMPRAGE-32 channel combination were significantly lower than those of all the other combinations. Although the TRV values for the MEMPRAGE-32 channel combination were not significantly lower than those of the other combinations for the total brain structures when using the MorphoBox, it was still the lowest one numerically. We also found that, regardless of the coil type, MEMPRAGE outperformed the other two sequences regarding brain volume repeatability, especially for cortical structures.

The TRV values in the present study were similar to those reported in previous studies which also indicated high reliability for the same sequences. Our results showing that the TRV values ranged from 0.00% to 9.89% using the MorphoBox prototype and 0.00% to 16.46% using the FreeSurfer are comparable to those of previous studies indicating high reliability for similar sequences. Seiger et al. (25) reported TRV values that were under 3% in 78% of scanned regions when using the MPRAGE sequence. Okubo et al. reported coefficients of variation (COV) values (2× COV = TRV) for deep GM that were 0.50% to 3.31% when using the MP2RAGE sequence and 0.62% to 4.12% when using the MPRAGE sequence (26). Our results may have been slightly higher than those reported in these two studies because we used a different analysis procedure. These studies used voxel-based morphometry (VBM) analysis for brain segmentation, while we used MorphoBox and FreeSurfer. However, the median TRV values in our study were 1.23% with MorphoBox and 3.14% with FreeSurfer, which were low and comparable with those in former studies.

The results of our TRV comparisons between different sequences were also consistent with previous studies. We found that the MEMPRAGE sequence usually performed more reliably than the other two sequences for total brain structures and cortical structures. Another study also showed that the MEMPRAGE sequence had numerically but not significantly higher repeatability for cortical thickness measures and cortical surface placement than the MP2RAGE sequence (20). In addition, another study that solely compared the MPRAGE and MP2RAGE sequences at 7T found that MP2RAGE had a larger TRV than MPRAGE especially in the inferior temporal region (25). Although a study demonstrated that MPRAGE was more reliable than MEMPRAGE, the reliability of MEMPRAGE in that study was still generally high and was comparable with the reliability of MPRAGE (27). Furthermore, there were some regions in our study in which the MPRAGE sequence performed more reliably than MEMPRAGE.

Differences between our results and the results of other studies may have occurred because we used a higher bandwidth for MEMPRAGE than the other studies, which could have resulted in enhanced repeatability.

Similar to other studies, our TRV results for subcortical structures were different from our results for total brain and cortical structures. Both the results from MorphoBox and FreeSurfer demonstrated that the TRV values for the MP2RAGE sequence, regardless of coil type, were the lowest of the three sequences for total subcortical structures numerically, but this difference was not significant. Similar results were also found for some single subcortical structure, such as the left deep WM when MorphoBox was used, and the left thalamus when FreeSurfer was used. MP2RAGE also performed more reliably than the other sequences for certain deep cortical structures, such as the insula and cingulum. These findings are similar to those of a previous study that found that MPR2RAGE had significantly better reliability than MPRAGE in the bilateral putamen and had comparable reliability with MPRAGE in other deep GM areas (26). The superior reliability of the MP2RAGE sequence for subcortical and deeper cortical structures may arise because this sequence is less influenced by B1 inhomogeneity, allowing it to more easily distinguish the subcortical WM and GM, which is challenging because subcortical GM has a lower T1 values than cortical GM (28).

Our results did not consistently indicate a better coil for the majority of single regions. However, regardless of sequence, scanning with the 32-channel coil resulted in more repeatable volumetric measures compared to the 20-channel coil for total structures, total cortical structures, and total subcortical structures. This result may occur because coils with more channels improve the SNR in the brain, especially in the cortex (14,16) and the SNR positively correlates with the reproducibility of MR images (29,30). Therefore, by improving image quality, coils with higher numbers of channels may improve brain scan and segmentation reliability.
This study has a few limitations. First, this study did not compare the absolute volume values of different brain regions between the sequence and coil combinations. This lack of information regarding absolute volumes may have detracted from the accuracy of the resulting measurements used in our studies because previous studies have reported that different sequences and coils lead to variable measurements of brain structure volume and surface region thickness, indicating that some sequences and/or coils may over- or under-estimate the aforementioned measurements (25,31). However, this study was more concerned with repeatability comparisons than accuracy. Second, we obtained repeated acquisitions within a single scan session, which means that factors affecting reliability, such as differences in head positioning, auto shimming, and operational error were not taken into account (4). Future studies should include rescanned acquisitions acquired across different sessions. However, previous studies have reported comparable reliability between images acquired in the same session and images acquired in different sessions (20).

Conclusions

In conclusion, our study showed that overall the MEMPRAGE sequence provides better scan and segmentation repeatability for brain structures, especially cortical structures, than MPRAGE or MP2RAGE sequences. However, the MP2RAGE sequence may perform better when targeting subcortical structures volumetry. Our results also suggest that the 32-channel coil may have better scan reliability than the 20-channel coil. These findings may serve as a reference for choosing more reliable sequence and coil combinations in future studies and, potentially, in clinical practice.

Acknowledgments

We are very grateful to Dr. Ang Li and Prof. Bing Liu of Brainnetome Center, Institute of Automation, Chinese Academy of Sciences, for helping us in the data extraction and analysis.

Footnote

Conflict of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The Institutional Review Board of our hospital approved this study (No. ZS-1137), and informed consent was obtained from all the participants.

References

9. Mugler JP 3rd, Epstein FH, Brookeman JR. Shaping the signal response during the approach to steady state in...


