4548

Postmortem Imaging with Reusable 3D Printed Ex Vivo Brain Enclosures/Cutting Guide for MRI Registration with Gross Anatomy Photographs at 7T

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Synopsis

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The ability to continuously prototype and 3D print new containers as well as using an improved transmit coil has led to major improvements in ex-vivo image quality. This has also allowed for a more automated registration process of the MR images to the gross anatomy

Introduction

Since the creation of the 3D printed ex vivo brain enclosure and cutting guide more than 40 brains have been scanned at 7T and registered to the pathology as well as in vivo scans when available. This novel approach to scanning, cutting and registering MRI images to the gross anatomy has proven to be robust and reusable.

Previously this study utilized the 16-channel Tic-Tac-Toe (TTT) Coil [1,2] and since has moved to the newly designed 60-channel single-transmit/32 channel receive TTT Coil [3]. There have also been updates to the container to increase the usable volume inside the container while decreasing the volume it takes up inside the rigid receive coil.

Methods

The updated container, visible in Figure 1.b, makes use of a domed lid design which allows for an overall smaller container base while increasing the usable volume the brain occupies in the container. The container has also been rounded to eliminate all sharp corners and fit easier in the oblong head shaped receive coil. The previous fill port and nozzle connector have been replaced with a watertight screw cap. The larger hole allows for hemisphere volume between the top of the container base and the peak of the domed lid to be filled entirely eliminating the possibility for there to be a layer of air between the lid and agar.

The left hemisphere of the brain is separated from the right and cerebellum and embalmed in 4% PFA until the day of scanning. An agarose mixture of water, 1.5% agarose and 30% sugar by weight is created and heated to a liquid. The brain is placed in the cutting guide and the cutting guide is placed into the base of the container. Agarose is poured into the container and around the brain carefully ensuring minimal air bubbles are left in the sulci of the brain and ventricles. The lid is then attached, more agarose is added until the container is filled, and the screw cap is secured. The container is wrapped in a plastic bag prior to scanning as an added security measure protecting the coil from the possibility of liquids damaging the coil and providing a barrier between the human tissue/PFA and coil.

The container is then placed into the coil as shown in Figure 1.c, note that it is rotated slightly for improved B1+. The first MR images acquired is a B1+ map using a Turbo-FLASH sequence with TR/TE = 2000/1.16ms; TA = 12min; flip angle from 0° to 90° in 18° increments; and 3.2mm isotropic resolution. Next is an Mp2rage with TA = 32min; TR/TE = 6000/4.1ms; Flip Angle = 6 and 7 degrees; TI1 = 514; TI2 = 2020ms; and 0.4mm isotropic resolution. A GRE is then acquired with TA = 59min TR = 40ms; TE1 = 8ms; TE2 = 15ms; TE3 = 21ms; and .37mm isotropic resolution. A T2-SPACE is acquired with TA = 1hr 38min TR/TE = 3400/363ms; and .36mm isotropic resolution. When transitioning to the 60-channel coil the voltage for the T2-SPACE voltage was scaled proportionally for the higher B₁⁺ field intensity, the new coil provides

Results

In Figure 2.a b and c are the 60-Channel T2-SPACE, GRE, and MP2RAGE respectfully. The quality of these images can be compared to those in Figure 3.a b and c from the 16-channel coil MP2RAGE and GRE respectively.

The image registration process has been automated making use of a photograph provided from the pathologist of cutting locations in order to compute structural similarity index and applying a position gate to guarantee alignment of edge cases. The registered MR images to gross anatomy are visible in Figure 3 and are a near perfect match at all locations. The image quality from the 16-channel to 60-channel TTT coil can be seen in Figure 4. Masks of two separate brains were created and used to analyze the B1+ only where the brain occupies space within the container and are visible in Figure 4.

Discussion

A major benefit our method provides is that the brain is scanned intact with minimal deformations using agarose as a medium to suspend the tissues [4]. This replicated in vivo condition also allows the pathologist to review the images prior to cutting. After deciding where to cut, the design of the cutting guide also provides the pathologist with the freedom to cut at any location with a variety of angles through any singular location.

The slightly smaller container size allows for more flexibility when positioning the container in the rigid receive coil. This is important because the 60-channel TTT coil is tuned for a human head.

Conclusion

The improved container shape and lid has resulted in significant improvements, specifically minimizing the manual labor by the pathologist as well as image quality improvement by reducing air bubbles during embedding. The use of the 60-channel single-transmit TTT Coil has also improved the image quality.

Future work may include slightly altering the container shape further to make it smaller still and refining the agarose doping concentrations to mimic the dielectric properties of the human head to achieve better homogeneity as seen in vivo [3].

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Figures



Figure 1: The cutting guide has not changed from the first a) design to the updated version b). The exploded view of the 5-piece assembly fit together as shown and the lid is screw to the base. The lid has a recessed gasket that fills a groove sealing it to the base. The container can be seen places inside the coil in c). Note the slight tilt of the container used to obtain a more ideal B₁⁺.



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ISMRM 2023

Figure 2: The images shown are a a) T2-Space b) GRE and c) MP2RAGE acquired using the 60-channel single-transmit/32-channel receive Tic-Tac-Toe coil.



Figure 3: The images shown are a a) MP2RAGE and b) GRE acquired using the 16-channel single-transmit/32-channel receive Tic-Tac-Toe coil.



Figure 4: Shown are two different examples of B1+ maps for two postmortem brains of slightly different size and positioning. The signal from the agar has been masked out to show only the B₁⁺ where the brain is.



Figure 5: Shown is a few slices of the registered gross anatomy with the acquired T2-SPACE and MP2RAGE.

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4548

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