In-room ultrasound fusion combined with fully compatible 3D-printed holding arm - rethinking interventional MRI

Friebe, Michael; Sanchez, Juan; Balakrishnan, Sathish; Illanes, Alfredo; Nagaraj, Yeshaswini; Odenbach, Robert; Matooq, Marwah; Krombach, Gabriele; Vogele, Michael; Boese, Axel

Published in:
Medical devices-Evidence and research

DOI:
10.2147/MDER.S150459

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2018

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):

Copyright
Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

Take-down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): http://www.rug.nl/research/portal. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.
In-room ultrasound fusion combined with fully compatible 3D-printed holding arm – rethinking interventional MRI

Michael Friebe,1 Juan Sanchez,1 Sathish Balakrishnan,1 Alfredo Illanes,1 Yeshaswini Nagaraj,2 Robert Odenbach,1 Marwah Mataoq,1 Gabriele Krombach,3 Michael Vogele,4 Axel Boese1

1Chair of Intelligent Catheter, Otto-von-Guericke-University, Magdeburg, Germany; 2University of Groningen, University Medical Center Groningen, Center for Medical Imaging North East Netherlands, Groningen, the Netherlands; 3Universitätsklinikum Giessen, Radiologische Klinik, Giessen, Germany; 4Interventional Systems GmbH, Kitzbühel, Austria

Abstract: There is no real need to discuss the potential advantages – mainly the excellent soft tissue contrast, nonionizing radiation, flow, and molecular information – of magnetic resonance imaging (MRI) as an intraoperative diagnosis and therapy system particularly for neurological applications and oncological therapies. Difficult patient access in conventional horizontal-field superconductive magnets, very high investment and operational expenses, and the need for special nonferromagnetic therapy tools have however prevented the widespread use of MRI as imaging and guidance tool for therapy purposes. The interventional use of MRI systems follows for the last 20+ years the strategy to use standard diagnostic systems and add more or less complicated and expensive components (eg, MRI-compatible robotic systems, specially shielded in-room monitors, dedicated tools and devices made from low-susceptibility materials, etc) to overcome the difficulties in the therapy process. We are proposing to rethink that approach using an in-room portable ultrasound (US) system that can be safely operated till 1 m away from the opening of a 3T imaging system. The live US images can be tracked using an optical inside–out approach adding a camera to the US probe in combination with optical reference markers to allow direct fusion with the MRI images inside the MRI suite. This leads to a comfortable US-guided intervention and excellent patient access directly on the MRI patient bed. This was combined with an entirely mechanical MRI-compatible 7 degrees of freedom holding arm concept, which shows that this test environment is a different way to create a cost-efficient and effective setup that combines the advantages of MRI and US by largely avoiding the drawbacks of current interventional MRI concepts.

Keywords: interventional MRI, fusion imaging, ultrasound/MRI hybrid, MRI-compatible, medical holding arm ultrasound/ultrasound guided MRI intervention

Introduction to interventional MRI

The unparalleled soft tissue contrast and excellent image quality of magnetic resonance imaging (MRI) have made it the imaging modality of choice, especially for neurological and oncological applications. The obtained MRI images are then frequently used as relevant preoperative images for surgery planning or as a monitor-presented visual reference in the surgery suite.

MRI systems directly employed for therapeutic guidance or surgical monitoring are rarely available. The standard imaging systems in the surgery room are typically X-ray, ultrasound (US), or endoscopy (video) and are used to perform the patient monitoring, controlling, and device guiding functions.

To use these preoperative MRI images for therapy purposes as well usually requires fusion with the available imaging data in the surgery room. That is either
not done at all or happens only in the head of the clinician, so called cognitive fusion. This requires experience and training and leads to inaccuracies, subjectivity, variability, and lack of reproducibility. It can also be accomplished by more or less accurate and complicated special software tools in combination with optical and/or electromagnetic device tracking systems.1–4

For the fusion via software-based image coregistration, the images are manually or semi-automatically registered to each other. A preoperative MRI image – as the wording implies – was taken before the surgery, however, sometimes even on a bed with different curvature than the surgery table or in a different patient position.

For prostate MRI, for example, the imaging is typically done with the patient in a straight supine position, while the actual biopsy or therapeutic treatment is performed with the patient prone or supine with the legs angled toward the torso.

And for breast MRI, the patient typically lies in an elevated prone position while surgery is performed on a surgical table in supine position or on the side. You easily recognize the difficulties using these preoperative MRI images for accurate image fusion with the available imaging modalities available in the therapy or surgery room.

For both procedures, there is relatively little space left in a conventional horizontal-field MRI system, not leaving maneuverable space for the surgeon or longer instruments, especially if they are orthogonal to the patient body.

Real precise image registration (US to MRI) is also not possible in this way, with typical image registration errors of the order of 3–7 mm, and in case of missing fiducial marker structures sometimes even significantly larger. That may be sufficient for actual therapies using an additional imaging system or for coarse orientation, but is too large for tissue biopsies and other precise therapies on small structures. This can be reduced to 2–3 mm by adjusting the MRI imaging position to the actual surgery position.5

Using real-time MRI as a guidance device for intraoperative or interventional therapies (eg, biopsies and treatment of prostate cancer, liver laser ablation/radiofrequency ablation/or cryoablations, brain interventions) could be very beneficial and would obviously eliminate or greatly reduce these registration errors.

This requires that the therapeutic procedure is performed in the MRI suite or the MRI system is placed in a dedicated surgery room.

The latter is an option that has been explored by several institutions, but comes with very high investments and requires very highly skilled interdisciplinary staff (Figures 1 and 2). These systems are ideally suited for research institutions that explore image guidance options for developing advanced procedures, but are not an option for the large majority of clinical providers at the moment.

Conventional diagnostic MRI for interventions

Difficult patient access in conventional horizontal-field superconductive magnets and the need for special nonmagnetic therapy tools have also prevented the widespread “occasional” interventional use of conventional diagnostic MRI systems.

This means that an MRI system that was primarily purchased and installed for diagnostic imaging purposes is also used for selected easy biopsy or needle aspiration procedures. But even these relatively simple interventions require special tools that are safe for the patient (MR safe) and do not cause significant imaging artifacts (MR compatible).

A typical artifact is caused by the susceptibility of the devices material, which can lead to a significant increase in the diameter of the device. In a high-field MRI system, an MRI-compatible needle is, for example, shown with a 10 mm or greater diameter, but actually only has a diameter of 2 mm. It is obvious that with these artifacts, guidance and secure placement in smaller structures is very difficult and sometimes impossible.

Any external system that is used in or close to the MRI magnet is exposed to the strong magnetic fields, and so it must be ensured that the devices are not pulled into the magnet bore. Also, the strong electrical and magnetic fields of the MRI can lead to system malfunction or patient safety issues. But the devices itself also can influence the performance of the MRI if not properly shielded. A lot of things to consider.

The interventional and therapeutic use of MRI follows for the last 20+ years the strategy to employ conventional standard high-field systems and add more or less complicated and expensive components.2

MRI-compatible robotic systems were, for example, proposed to solve the patient access issue in combination with specially shielded in-room monitors, dedicated nitinol or plastic devices with no or little susceptibility-related imaging artifacts, and in-room optical tracking systems.5 These components are quite expensive, and so the possible therapies are still limited because of the tight space in the MRI bore and the issues around surgical sterility.
The robotic systems are even needed with the short-bore magnets of the newest MRI generation. With a magnet length of 1.20 m, it is still 60 cm (an arm’s length) to the center of the magnet, and with a bore diameter of 70 cm – with a mid-size patient placed – there is only 10–15 cm of available vertical space for a therapy tool (Figure 3).

Open vertical-field magnets do provide some benefits for interventional procedures, but typically have only magnetic field strength of 0.2 T to 0.4 T, with currently only one system over 1 T and another one at 1 T that is unfortunately phased out (Figure 2). Lower field strength comes with increased acquisition times, which is not good for therapy applications, and a lower image quality. On the other hand, the lower field strength also reduces susceptibility artifacts and comes with a much lower magnetic attraction force making the surgery generally safer. The authors do believe, however, that the low-field systems could be an excellent base for easier, dedicated procedures, especially when preoperative high-field and high-quality MRI images are used as a base of image fusion.

**High-field MRI with in-room US**

The difficulty of patient access in the high-field MRI systems lead to the question of whether or not it would make more sense to rethink the interventional MRI procedure by combining MRI with US inside the MRI imaging room. For many procedures, US guidance is the gold standard that could benefit from additional, near real-time MRI guidance and imaging.5,6

The general idea is to acquire comprehensive and high-quality MRI images of the area of interest, move the patient out of the tight bore remaining on the MRI patient bed, and then continue with the therapy under US guidance. For further MRI images or updates, the patient is moved back into the MRI system. This may need to be done several times during the placement of the device to ensure that the target site
has been reached and should also be done at the end of the procedure to record the therapy result for quality assurance. The MRI image is much better suited to give details on the therapy region and to show the effects of the procedure.

The images of the MRI and US are either cognitively fused (Figure 4B), or automatically coregistered and overlaid to the just-obtained MRI images using special tracking and navigation techniques (Figure 5). Several commercially available US systems could theoretically be used even as close as 1 m to a 3 T magnet opening (Figure 4A).

While these systems do not have an official MRI-safe label and are not tested and approved for use in an MRI imaging suite, research institutions like ours have used these systems in phantom setups to rethink interventional MRI procedures. The US system that we used (GE Venue 50) with a commercial 3T MRI (Siemens Skyra) turned itself off once it was moved too close to the magnetic field – at around 25 mT – but we did not experience major magnetic attraction forces even at the 50 mT magnetic field line.

A more accurate alternative to the cognitive fusion using the MRI images and the live US images (Figure 4B) is to use a marker-based approach in combination with a navigation system that is attached to the US probe. The marker, for example, a combination of an MRI-visible component (eg, Vitamin B pills) and an optical marker, allows for registration of live US with the just acquired MRI images. This approach also has the advantage compared to the normally used outside–in optical tracking systems in that there is no or only a small issue with obstructing the line of sight between the camera and the marker and it is rather cheap and easy to use. The disadvantage is the rather small field of view, but that may not be as relevant as the area of procedure is known and limited in size.
This setup and process (Figure 5) allows a rather comfortable US-guided intervention for the surgeon standing next to the patient, with plenty of working space in all directions and excellent patient access directly on the MRI patient bed.

Low-cost interventional MRI holding arm
External intervention on the MRI patient bed may need additional MRI scanning during the procedure for verification or monitoring, which in turn requires that the therapeutic
devices are fixed and held in position while the patient is moved back into the MRI magnet. This is ideally achieved using a completely MRI-compatible holding arm that is purely mechanical and manufactured from nonferrous or even plastic materials to avoid any magnetic attraction and to ensure that neither is the arm influenced by the MRI nor is the MRI negatively affected by the holding arm while maintaining a safe therapy setup for the patient.

We developed and 3D printed a completely new and modular holding arm concept from plastic material with 7 degrees of freedom that can be attached to the MRI bed railing and adjusted based on procedure needs and patient shape (Figure 6).

An initial MRI scan would provide the initial target site details used for the insertion path planning and the baseline scan for the US fusion.

Depending on the complexity, a quick MRI may need to be done to verify that the target site has been reached, and another scan can be done at the end of the procedure to inspect and record the therapy results (quality assurance).

**Discussion and conclusion**

MRI/US fused and used together and in parallel combines the excellent contrast and details of the MRI image with the real-time high-resolution device guidance of US and shifts the intervention space from inside the MRI to the MRI table.

Combining state-of-the-art high-field MRI with relatively low-cost US would provide superior baseline image quality with a fast and easy to use imaging system for guidance.

MRI/US fusion is already offered by several vendors for prostate biopsy applications. These systems generally, however, fuse preoperative MRI images with live and real-time US. Registration errors predominantly resulting from different patient position during imaging can be of the order of 7 mm.

The presented setup showed the feasibility of using an US system as the primary guidance and monitoring device for interventional MRI procedures. To avoid the limitations of cognitive fusion and the cost and complexity of conventional optical tracking systems, we suggest a fusion of the MRI and US images using an inside–out tracking approach with a stereo camera system mounted to the US transducer camera in combination with MRI and US visible markers on the patient’s body. Obviously when using this, the procedure should rather be called “Interventional US/MRI fusion procedure.”

To make this procedure feasible for occasional use of the MRI system as an interventional and therapy guidance system, all the involved components need to be relatively inexpensive, easy to use, and easy to remove while providing the utmost safety for the patient and the surgeon.

For that, we propose the additional use of a small and very flexible holding arm that supports the surgeon and also holds
the therapy devices in place for additional MRI scanning during the procedure. This holding arm can be attached and removed in <1 minute. The US system can be moved to the end of the MRI bed (and back) in <1 minute, and the attached navigation camera on the US probe is permanently available. The time for the marker setup and the registration procedure is still a somewhat time-consuming process, but can certainly be automated and sped up in the very near future.

Since most MRI systems are almost exclusively used for diagnostic purposes, such an easy to set up and remove system would even be attractive for sites that only do few interventional or therapeutic MRI procedures. And, it would particularly be interesting for some of the easier and some of the more complex interventional MRI procedures (Figure 7 – green and orange exclamation marks).

Difficult cardiac or neurosurgical procedures would require a dedicated surgery room with an MRI installed or available (Figure 1).

We evaluated the presented interventional MRI setups, conventional closed-bore high-field MRI (Figure 3), dedicated interventional MRI procedure room (Figure 1), and vertical-field MRI (Figure 2), with the proposed Interventional US/MRI fusion procedure setup (Figures 4–6) based on patient access, comfort to the surgeon, real-time imaging and therapy guidance, as well as total cost, for both easier and more advanced procedures (Figure 7).

The findings are presented in Table 1 and show that the Interventional US/MRI fusion procedure setup has potential advantages for all but the more complex procedures.

With MRI-compatible and approved US systems and easy to use and inexpensive accessories, interventional MRI could become an attractive option for other clinical applications as well.

If the interventional procedure is guided by the US system, an MRI-compatible in-room monitor is not needed anymore. As the US and the MRI are never used together, there will most likely not be any US-related MRI imaging artifacts. The cost of the in-room monitor alone is likely almost the same as the total cost of the devices mentioned in this paper.

This paper cannot present a thorough patient- and procedure-based evaluation research on the possible advantages of the setup as there are no MRI-compatible and MRI-safe US systems commercially available and as the holding arm and tracking method as presented are not clinically certified products. The intention of the presented perspective should help researchers rethink the current technical component research of interventional MRI.

We hope that many more research and industry teams will agree with this rethinking and help further develop procedures, tools, and devices that can be used for upcoming clinical validation.

Figure 6 3D printed fully MRI-compatible modular holding arm individually adjustable for procedure needs and patient shape. Notes: The arm with 7 degrees of freedom is directly attached to the rail of the bed and entirely manufactured from plastic materials. Abbreviation: MRI, magnetic resonance imaging.
**Table 1** Comparison of different interventional MRI setups for easy and more complex procedures based on patient access and surgeon comfort, as well as on real-time therapy guidance capabilities in combination with high-quality imaging, and the ability to use for different procedures as highlighted in Figure 7.

<table>
<thead>
<tr>
<th>Intervventional MRI issue</th>
<th>Standard diagnostic MRI – horizontal field (Figure 3)</th>
<th>Standard diagnostic MRI – vertical field (Figure 2)</th>
<th>In-room US fusion added (Figure 4/5)</th>
<th>Dedicated interventional MRI suite (Figure 1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient access for the therapist: How accessible is the therapy region for the surgeon?</td>
<td>-- Typically 70 cm diameter and &gt;60 cm to the MRI center – very difficult for the surgeon to reach the surgical target.</td>
<td>0 Typically 45 cm vertical opening and around 50 cm to the MRI center, patient access &gt;240 degrees.</td>
<td>+ Full patient access for therapy.</td>
<td>+ Full patient access for therapy.</td>
</tr>
<tr>
<td>Surgeon comfort: Is the surgeon able to perform the surgery in a comfortable position?</td>
<td>-- Surgeon has to reach into the magnet bore in a sitting or bent position.</td>
<td>0 Surgeon typically can sit next to the magnet bore but still has to reach in the bore.</td>
<td>+ Surgeon can stand next to the patient in a normal standing position.</td>
<td>+ Surgeon can stand next to the patient in a normal standing position.</td>
</tr>
<tr>
<td>Real-time therapy guidance: Focus is here on whether real-time imaging for tool and device tracking is available – not for the actual diagnostic imaging process!</td>
<td>0 Only MRI is useable for guidance – fast imaging is low resolution; high-quality imaging is slow.</td>
<td>0 Only MRI is useable for guidance – fast imaging is low resolution; high-quality imaging is slow.</td>
<td>+ The US provides real-time imaging to the previously acquired high-quality MRI.</td>
<td>+ Interventional MRI suites come with additional real-time imaging in form of X-ray and/or US.</td>
</tr>
<tr>
<td>Interstitial therapies (green “!” in Figure 7): Is the system setup suitable for the green therapies listed in Figure 7?</td>
<td>0 Setup can be used with limitations for easy procedures. Very limited space and reduced view of surgical site.</td>
<td>0 Setup can be used with limitations for easy procedures. Very limited space and reduced view of surgical site.</td>
<td>+ Improved due to better patient access and better surgical site view.</td>
<td>++ Excellent patient access in combination with real-time image guidance and no space restrictions in the magnet bore.</td>
</tr>
</tbody>
</table>

**Figure 7** Possible therapies for the Interventional US/MRI fusion procedure setup.

**Notes:** The easier procedures are marked with a green exclamation mark. Medium/complex procedures (orange exclamation mark) could potentially be done with the proposed setup, while the difficult ones should be reserved to dedicated surgery rooms with advanced imaging modalities (eg, MRI) installed. The procedures marked with a red exclamation mark are reserved for dedicated interventional MRI suites as shown in Figure 1.

(Continued)
Table 1 (Continued)

<table>
<thead>
<tr>
<th>Interventional MRI issue</th>
<th>Standard diagnostic MRI – horizontal field (Figure 3)</th>
<th>Standard diagnostic MRI – vertical field (Figure 2)</th>
<th>In-room US fusion added (Figure 4/5)</th>
<th>Dedicated interventional MRI suite (Figure 1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>More advanced therapies (orange and red “+” in Figure 7): Is the system setup suitable for the red! and orange therapies listed in Figure 7?</td>
<td>– MRI is not installed in a real surgical suite. Sterility and airflow is not acceptable for complicated open surgeries and there is only limited monitoring and support equipment available. Not feasible for complicated surgical procedures.</td>
<td>– MRI is not installed in a real surgical suite. Sterility and airflow is not acceptable for complicated open surgeries and there is only limited monitoring and support equipment available. Not feasible for complicated surgical procedures.</td>
<td>0 MRI is not installed in a real surgical suite. Sterility and airflow is not acceptable for complicated open surgeries and there is only limited monitoring and support equipment available. Not feasible for complicated surgical procedures.</td>
<td>+ MRI is installed in a real surgical suite and comes with all equipment found in an advanced surgical setup.</td>
</tr>
</tbody>
</table>

Cost of Setup for iMRI procedures: Baseline is a conventional 1.5T diagnostic MRI with a horizontal field

| Cost of MRI plus €50k to €150k for the Interventional MRI Package consisting of MRI-compatible in-room monitor(s), dedicated nonmagnetic therapy tools, anesthesia and monitoring equipment, and manipulation devices. | Cost of a high-field vertical-field MRI is about €100k to €500k above the price for a standard diagnostic MRI with horizontal field plus the cost of the Interventional MRI Package. | Cost of MRI plus Interventional Package plus €50,000 for the US and tracking hardware and software. | The cost for the building/construction and the additional equipment are very high (>$1.000k) – also a reason why there are only few such systems installed worldwide. |

Notes: The additional cost for such a setup was also coarsely evaluated. The table evaluation is always based upon the best possible option for the particular point in question (eg, the most open patient access for therapeutic interaction is a patient on a bed freely accessible, which gives a “+” to the “In-Room US Fusion” concept and the “Dedicated Interventional MRI”). A “0” indicates it is possible/OK, “–” not feasible, “−” feasible, and “+-” excellent/prohibitive.

Abbreviations: MRI, magnetic resonance imaging; iMRI, interventional MRI; US, ultrasound.

Disclosure
The authors report no conflicts of interest in this work.

References