

MR-Guided Focused Ultrasound* in Fibroid Treatment – Results of the 3rd Radiological-Gynecological Expert Meeting

Magnetresonanzgeführter fokussierter Ultraschall zur Myombehandlung – Ergebnisse des 3. radiologisch-gynäkologischen Expertentreffens

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DOI <https://doi.org/10.1055/s-0043-108994>

Published online: 2017 | Fortschr Röntgenstr 2017; 189: 515–518 © Georg Thieme Verlag KG, Stuttgart · New York, ISSN 1438-9029

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Introduction

Fibroid treatment with MR-guided focused ultrasound (MRgFUS; syn.: HIFU = high-intensity focused ultrasound) is a thermoablative method in which the tissue to be treated is heated by focused ultrasound in single small volume increments (sonifications, syn: sonications) under constant MRI control until complete denaturation of the planned fibroid volume is achieved. After thermoablation, imaging showed a lack of contrast enhancement of the treated tissue (NPV = non-perfused volume).

MRgFUS is organ-preserving and noninvasive and can be performed on an outpatient basis.

The treatment method is offered only by a few specialized centers.

The goal of MRgFUS treatment is to reduce or eliminate fibroid-related symptoms in affected women. A reduction in fibroid size can be achieved with ultrasound treatment. Complete fibroid regression is not to be expected and is also not the goal of the treatment.

The disciplines of gynecology and radiology agree that the indication for the treatment of uterine fibroids should be determined by a gynecologist following examination and counseling of the patient. Comprehensive patient counseling regarding the

treatment options in symptomatic uterine fibroids should include medication, surgery, and the two non-surgical treatment options uterine artery embolization (UAE) and MRgFUS. The decision for or against a treatment alternative should be made under consideration of the patient's wishes and with knowledge of other treatment options, the chance of success, limitations, typical side effects, and possible complications (informed consent).

MRgFUS treatment provides a treatment method for patients with fibroid-related symptoms and allows further treatment individualization for uterine fibroids in Germany, Austria, and Switzerland.

Goal of the consensus meeting

The intention of this third consensus meeting was to evaluate and categorize MRgFUS in the fibroid treatment spectrum. The 12 participants of the radiological-gynecological expert meeting came to a consensus following thorough discussion with evaluation of the current literature¹ and their own experiences.

The group of experts was aware that the possibilities and limitations of a radiological treatment method was being discussed together with gynecology specialists who do not actually perform

* To be differentiated from non-MR-guided focused ultrasound.

1 The appendix contains references to select relevant publications.

the procedure but have expertise and experience with the diagnosis and medication-based and surgical treatment of diseases of the female genitals.

The expert group comprised of 4 radiologists and 8 gynecologists that met on January 14, 2017 in Berlin for the third radiological-gynecological consensus meeting regarding MRgFUS treatment also included gynecologists from Switzerland.

Following extensive and at times also controversial discussion, the group agreed in consensus upon the following recommendations. The consensus paper is supported by the gynecologists and radiologists listed at the end of the article. The paper reflects the current state of knowledge.

Structural requirements for performing MRgFUS treatment

MRgFUS should only be performed at hospitals with the necessary expertise and experience. This also includes the conservative and surgical management of side effects and complications. Moreover, there should be options for introducing postinterventional pain therapy.

Examinations required prior to MRgFUS treatment

Treatment decisions are based on gynecological examination including vaginal and/or abdominal ultrasound (depending on the size of the uterine fibroid). An MRI scan with contrast agent ideally in prone position must be acquired for planning purposes. The contrast-enhanced image helps to assess whether and to what degree the fibroid is perfused.

Prior to every MRgFUS procedure, the indication for hysteroscopy and fractionated abrasion should be examined as a function of bleeding pattern and endometrium thickness and structure. A cytological smear of the cervix uteri with a normal result needs to have been performed within the last 12 months.

During the informed consent discussion prior to MRgFUS, the patient should also be informed of the lack of preinterventional histological confirmation, as in all other organ-preserving fibroid treatment methods.

Indications for MRgFUS treatment

A symptomatic uterine fibroid with an anatomical position allowing safe access for MRgFUS is an indication for MRgFUS treatment. Treatment is complicated by the presence of more than five fibroids. In the case of fibroids with a diameter of more than 10 cm, the indication for MRgFUS treatment should be carefully considered due to the large fibroid volume and the associated long treatment time.

MRgFUS represents an alternative to surgical and medication-based methods such as UAE. Treatment decisions should be based on the treatment objective and the wishes of the patient. If tech-

nically feasible, MRgFUS is a good option for patients desiring the least invasive treatment possible.

Success criteria for MRgFUS treatment

The goal of treatment with focused ultrasound is to achieve the greatest possible NPV (= non-perfused volume).

The improvement or elimination of fibroid-related symptoms is viewed as treatment success following MRgFUS treatment. A volume reduction is desired but is considered a secondary treatment goal.

Contraindications for MRgFUS treatment

- Primarily Malignancy (absolute)
- Pregnancy (absolute)
- Acute inflammatory process (absolute)
- Subserosal pedunculated fibroids (absolute)
- Submucosal fibroids type 0 and I (relative; absolute in case of a desire to have children)
- Insufficient acoustic window for treatment (e. g. bowel overlying the fibroid) (absolute)
- More than 5 fibroids (relative, decided on a case-by-case basis)
- Uterine fibroids with a diameter greater than 10 cm (relative, decided on a case-by-case basis)
- Large scars in the acoustic window (relative)
- Fibroid positioned near the os sacrum (relative)
- General contraindications to MR contrast agents (relative)
- Relative and absolute MRI contraindications

Ulipristal acetate can result in increased perfusion of fibroids; consequently the evaluation of the feasibility of MRgFUS treatment and the actual treatment can be negatively affected.

In the case of suspicion of a malignancy of the uterus, MRgFUS is absolutely contraindicated.

MRgFUS treatment in patients with a desire to have children

There is no published prospective data regarding women who wish to have children and have been treated with MRgFUS/HIFU. Therefore, MRgFUS/HIFU treatment cannot be recommended prior to a planned pregnancy. However, if a patient wants to become pregnant after MRgFUS/HIFU treatment, a minimum wait time of approximately 6 months between fibroid treatment with MRgFUS and conception should probably be observed.

Side effects/complications of MRgFUS treatment

Relevant side effects and complications during and after MRgFUS treatment are rare:

- Pain

- Skin burns
- Inflammation of the subcutaneous fatty tissue and the musculature of the abdominal wall
- Paresthesia of the leg due to nerve irritation or damage
- Deep vein thrombosis (very rare)
- Intestinal perforation (extremely rare)

In addition to increased and/or irregular bleeding within three months after treatment, fibroid treatment with MRgFUS can result in discharge of (necrotic) fibroid material in terms of a “fibroid in the nascent state” that is unpleasant and painful for the patient. Uterus-preserving ablation performed via the vagina possibly also in combination with surgical hysteroscopy is also possible here. Perioperative antibiotic prophylaxis is recommended in these cases.

Follow-up after MRgFUS treatment

Follow-up by a specialist after MRgFUS is recommended. Imaging methods are helpful (e. g. ultrasound in connection with Doppler ultrasound, MRI). If treatment is not successful (no improvement in symptoms and/or increase in fibroid size) or in the case of pathologies on imaging (increase in size of fibroid(s) or uterus), further diagnostic workup is required.

Future

The recommendations regarding MRgFUS treatment of fibroids are to be revised in approximately two years under consideration of the data and experiences available at that time.

Appendix

Participants in the consensus meeting 2017

Prof. Dr. med. Michael Bohlmann/ Mannheim
 Dr. med. Alexander Burges/ Munich
 Prof. Dr. med. Matthias David/ Berlin
 Prof. Dr. med. Markus Düx/ Frankfurt a.M.
 Prof. Dr. med. Dr. phil. Dr. h. c. mult. Andreas D. Ebert/ Berlin
 Prof. Dr. med. Peyman Hadji/ Frankfurt a.M.
 Dr. med. Thomas Hess/ Winterthur (CH)
 PD Dr. med. Peter Hunold/ Lübeck
 Dr. med. Hans-Christian Kolberg/ Bottrop
 Dr. med. Matthias Matzko/ Dachau
 PD Dr. med. Vera Schreiter/ Berlin
 Prof. Dr. med. Uwe Ulrich/ Berlin

Participating societies and working groups:

AGE, Arbeitsgemeinschaft Gynäkologische Endoskopie der DGGG
 AG URZ, Arbeitsgemeinschaft Universitärer Reproduktionsmedizinischer Zentren der DGGG
 Berufsverband der Frauenärzte (BVF)
 DeGIR, Deutsche Gesellschaft für Interventionelle Radiologie und minimal-invasive Therapie

DGGEF, Arbeitsgemeinschaft Gynäkologische Endokrinologie und Fortpflanzungsmedizin e. V.

DGGG, Deutsche Gesellschaft für Gynäkologie und Geburtshilfe
 DRG, Deutsche Röntgengesellschaft

SGGG, Schweizerische Gesellschaft für Gynäkologie und Geburtshilfe

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<http://dx.doi.org/10.1016/j.jogc.2016.02.006>

Published simultaneously

Published simultaneously in *Geburtshilfe und Frauenheilkunde* 6-2017:
Geburtshilfe Frauenheilkd 2017; DOI: 10.1055/s-0043-106257

Conflict of Interest

The authors declare that they have no conflict of interest.