

Uterine Artery Embolization (UAE) for Fibroid Treatment: Results of the 6th Radiological Gynecological Expert Meeting

Uterusarterienembolisation (UAE) zur Myombehandlung – Ergebnisse des 6. radiologisch-gynäkologischen Expertentreffens

Authors

Thomas Kröncke^{1*}, Matthias David^{2*}

Affiliations

1 Radiologie, Klinikum Augsburg, Germany

2 Gynäkologie, Charité, Berlin, Germany

Bibliography

DOI <https://doi.org/10.1055/s-0043-106259>

Geburtsh Frauenheilk 2017; 77: 689–692 © Georg Thieme

Verlag KG Stuttgart · New York | ISSN 0016-5751

Correspondence

Prof. Dr. med. Thomas Kröncke, MBA

Klinik für Diagnostische und Interventionelle Radiologie und Neuroradiologie, Klinikum Augsburg

Stenglinstraße 2, 86156 Augsburg, Germany

thomas.kroencke@klinikum-augsburg.de

Correspondence

Prof. Dr. med. Matthias David

Charité/Universitätsmedizin Berlin, Klinik für Gynäkologie,

Campus Virchow-Klinikum

Augustenburger Platz 1, 13353 Berlin, Germany

Introduction

Uterine artery embolization (UAE) is an organ-preserving, established, safe, and effective method in the spectrum of procedures for treating fibroid-related symptoms.

The aim of UAE is the reduction or elimination of fibroid-related symptoms, not the removal of the fibroid. At the same time the size of the fibroid is reduced.

There is consensus between the disciplines of gynecology and interventional radiology that determination of required treatment of uterine fibroids should be based on an examination and advice by a gynecologist. Comprehensive advice regarding treatment options for symptomatic uterine fibroids encompasses not only medication-based and surgical options but also UAE. The decision for or against an alternative therapy should be made taking into account the patient's desire for, and knowledge of, therapeutic alternatives, their chances of success and limitations, as well as typical side effects and possible complications (informed consent).

In Germany, Austria and Switzerland, UAE treatment offers the possibility of a therapeutic procedure for patients with fibroid-related symptoms which provides further individualization of therapy in cases of uterine fibroids.

Aim of the consensus meeting

The intention of the consensus meeting was to evaluate UAE. The participants in the meeting of radiological-gynecological experts, after taking into account the current literature, internationally published recommendations¹ and their own experience, and after extensive discussion, came to a consensus between the two disciplines.

The panel of experts was aware that this was an assessment of the possibilities and limits of a radiological therapy held in conjunction with specialists in gynecology who do not perform the procedure themselves, but who possess expertise and experience in the diagnosis as well as treatment diseases of female reproductive organs.

The group of experts composed of 11 radiologists and 8 gynecologists met on January 14, 2017 in Berlin for the sixth radiological-gynecological consensus meeting included radiologists and gynecologists from Switzerland and Austria. After extensive – and somewhat controversial-discussion, the group came to a consensus regarding the following recommendations. The consensus statement is supported by the gynecologists and radiologists listed at the end of this work.

It reflects the current state of knowledge.

* For the participants of the consensus meeting.

¹ The appendix contains references to selected relevant publications.

Structural prerequisites and quality assurance for performing UAE

UAE should be performed only at clinics possessing the requisite gynecological and radiological expertise regarding the performing of UAE, adequate and structured pain management after the intervention, management of side effects, and the conservative and surgical treatment of fibroids.

Particularly due to the necessity for postinterventional pain management, UAE should be performed on an inpatient basis at a suitable clinic.

Prior to introducing UAE, theoretical and practical training at a center with extensive UAE experience is recommended. In addition to the legally required documentation, the calculated key radiation exposure figures (dose area product, fluoroscopy time) for UAE should be critically reviewed and optimized for quality assurance.

Participation in suitable quality assurance as defined by the professional associations is recommended.

Examinations required prior to UAE

The choice of therapy should be based on an examination performed by a gynecologist including vaginal and/or abdominal ultrasound (as a function of the size of the uterine fibroid). If ultrasound does not allow definitive diagnosis, MRI examination is indicated.

Prior to fibroid embolization, the indication for hysteroscopy and fractionated curettage must be reviewed. Unremarkable cytological smear findings of the uterine cervix must have been obtained within the previous 12 months.

Pregnancy test results as well as the following laboratory results must be available: creatinine, coagulation status, thyroid values (in the case of a history of thyroid disease), blood count, and CRP. Acute inflammation must be ruled out in the case history and clinically.

According to the current state of knowledge, it is not necessary to remove an implanted IUD prior to UAE.

Within the context of the informed consent discussion prior to UAE, the patient should be informed regarding the absence of preinterventional histological confirmation of the presumed uterine fibroids, as is the case with all other organ-preserving fibroid therapies.

The total risk of an undetected uterine malignancy (including uterine sarcoma) in patients undergoing surgery for a presumed fibroid is specified between 0.09% and 0.18% in the current literature. Symptoms and imaging do not allow exclusion of a uterine sarcoma in particular.

The decision for an organ-preserving medication-based, surgical, or interventional-radiological treatment option should therefore include explanation of the risks of delayed diagnosis of a sarcoma. The spreading of tumor cells after UAE has not been observed. In the case of a lack of response to treatment or a lack of a reduction in the size of the leiomyoma(s), an insufficient embolization result and the presence of a uterine sarcoma must be considered in the differential diagnosis.

Indications for UAE

A symptomatic uterine fibroid is an indication for uterine artery embolization. UAE represents an alternative to surgical and medication-based procedures and to fibroid treatment with focused ultrasound regardless of the size and number of fibroids or previous surgeries. The choice of therapy should be based on the objective of the treatment as well as the wishes of the patient.

Success criteria for UAE

UAE treatment success is primarily defined as improvement or complete elimination of the (fibroid-related) symptoms specified by the patient and to a lesser extent as a reduction in the volume of the dominant fibroid or the entire uterus after treatment.

Contraindications for UAE

Technical

Relative

- GnRH-analog pretreatment in the previous 3 months (increased risk of vasospasm of the uterine artery)

According to the current state of knowledge, the administration of ulipristal acetate does not play a role in the indication for UAE or the performing of the procedure and does not affect the results.

Anatomical

Relative

- Isolated, submucosal fibroids type 0 and I of the FIGO classification (Fédération Internationale de Gynécologie et d'Obstétrique) that are accessible for hysteroscopic ablation
- Isolated pedunculated subserosal fibroids
- Supply of the fibroid(s) via an ovarian artery; the benefits and risks of additive embolization of the relevant ovarian artery should be considered.

Clinical

Absolute

- Primarily Malignancy
- Pregnancy
- Acute genital infection
- Clinically manifest hyperthyroidism/acute thyroiditis in hyperthyroidism and planned or ongoing radioiodine therapy if iodine-containing contrast agents are used

Relative

- Documented allergic reaction to contrast agents containing iodine
- Postmenopausal patient
- Allergy to local anesthesia
- Latent hyperthyroidism
- Renal insufficiency
- Desire to become pregnant

UAE in patients wishing to become pregnant

UAE is to be considered a last resort in patients wishing to become pregnant.

UAE in patients with a latent desire for children

For patients with a symptomatic uterine fibroid and a latent desire for children, the role of UAE as a treatment option is still not sufficiently defined in the current literature.

Pregnancy after UAE is possible. The risk of miscarriage may be increased.²

The preservation of fertility and a latent desire for children should be discussed with every patient on an interdisciplinary basis in connection with age, previous interventions, prior pregnancies, and imaging findings prior to UAE.

Before a hysterectomy is considered in a patient with a pronounced uterine fibroid who wishes to become pregnant, the possibility of performing UAE should be investigated.

Pregnancy after UAE

A minimum wait time of approximately 6 months between fibroid treatment with UAE and conception is recommended.

Special case: preoperative uterine artery embolization (PUAE)

PUAE, embolization as preparation immediately before surgical myoma enucleation, can be considered and offered in individual cases for patients who absolutely want to preserve their uterus but in whom a significantly increased bleeding risk can already be assumed preoperatively and/or in whom the risk of the ultimate need for a hysterectomy is estimated to be very high “for technical reasons” (e.g. very large fibroid and/or multiple fibroids, large fibroid that is difficult to remove, fibroid with unfavorable location).

Radiation protection

Radiation protection is particularly important in UAE. Pulsed fluoroscopy should be used. Serial angiography and oblique projections should be kept to a minimum. A scan frequency of 1 frame/second is typically sufficient. Under normal conditions, the average dose area product should be less than $50 \text{ Gy} \times \text{cm}^2$ (corresponding to $5000 \text{ cGy} \times \text{cm}^2$ or $5000 \mu\text{Gy m}^2$) for pulsed systems. The radiation exposure in this case would correspond approximately to 2 to 3 abdominal CT scans.

² Minority opinion (representatives of the DGGG and the AGR and AGE of the DGGG): In addition to miscarriage, placental problems and antepartum hemorrhage may be more common after fibroid embolization (insufficient reliable data).

Side effects

The following are described as relevant side effects and complications of UAE: Post-embolization syndrome, amenorrhea as a consequence of disruption or failure of ovarian function, pain, discharge, angiography-related complications (e.g. groin hematoma), vaginal discharge of fibroid material, hot flashes, endometritis/myometritis, deep vein thrombosis/pulmonary embolus.

Uterine discharge can be normal in the first weeks after UAE. In the case of abnormal vaginal discharge, the patient should be diagnosed and treated for infection. Menorrhagia, cramping of the lower abdomen, discharge of tissue components can occur in the case of submucosal fibroids in particular. Depending on the clinical symptoms and the findings of diagnostic imaging, hysteroscopic myomectomy or vaginal myomectomy as in a fibroid in statu nascendi can be indicated. Hysterectomy is not indicated a priori. In cases of doubt, the center conducting the UAE procedure should be contacted.

Post-treatment examination after UAE

Post-treatment examination by a specialist is recommended approx. 6 months after UAE.

Imaging procedures are useful (e.g., sonography in conjunction with Doppler sonography, MRI). Further clarification is required if the therapy is unsuccessful (no improvement of symptoms and/or size progression of the fibroid), or there are abnormalities in the imaging (increase in the size of the fibroid(s) or uterus) and/or there is a lack of devascularization of the fibroid(s).

Outlook

These recommendations regarding uterine artery embolization in the case of fibroid-related symptoms are to be revised again in 2019 based on the available data and experience.

Appendix

Consensus meeting participants

PD Dr. med. Ralf Adamus/Nuremberg
Dr. med. Robert Armbrust/Berlin
Dr. med. Michael Bartsch/Hamburg
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)
Prof. Dr. med. Thomas Kröncke/Augsburg
Prof. Dr. med. Peter Landwehr/Hannover
Dr. med. Matthias Matzko/Dachau
Prof. Dr. med. Thomas Pfammatter/Zürich (CH)
Dr. med. Gernot Rott/Duisburg
PD Dr. med. Dirk Schnapauff/Berlin
Prof. Dr. med. Uwe Ulrich/Berlin
Prof. Dr. med. Dierk Vorwerk/Ingolstadt
PD Dr. Peter Waldenberger/Salzburg (AT)

Participating professional associations and working groups

AGE, Arbeitsgemeinschaft Gynäkologische Endoskopie der DGGG [Gynecological Endoscopy Working Group of the German Society of Gynecology and Obstetrics]

AGR, Arbeitsgemeinschaft gynäkologischer Radiologie der DGGG [Gynecological Radiology Working Group of the German Society of Gynecology and Obstetrics]

AG URZ, Arbeitsgemeinschaft Universitärer Reproduktionsmedizinischer Zentren der DGGG [Working Group of University Reproductive Medicine Centers of the German Society of Gynecology and Obstetrics]

BVF, Berufsverband der Frauenärzte [Professional Association of Gynecologists]

DeGIR, Deutsche Gesellschaft für Interventionelle Radiologie und minimal-invasive Therapie [German Society for Interventional Radiology and Minimally Invasive Therapy]

DGGEF, Arbeitsgemeinschaft Gynäkologische Endokrinologie und Fortpflanzungsmedizin e.V. [Working Group for Gynecological Endocrinology and Reproductive Medicine]

DGGG, Deutsche Gesellschaft für Gynäkologie und Geburtshilfe [German Society of Gynecology and Obstetrics]

NOGGO, Nordostdeutsche Gesellschaft für Gynäkologische Onkologie [Northeastern German Society of Gynecological Oncology]

ÖGIR, Österreichische Gesellschaft für Interventionelle Radiologie [Austrian Society of Interventional Radiology]

SGGG, Schweizerische Gesellschaft für Gynäkologie und Geburtshilfe [Swiss Society of Gynecology and Obstetrics]

SSVIR, Swiss Society of Cardiovascular and Interventional Radiology

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Conflict of Interest

The authors declare that they have no conflict of interest.

Published simultaneously

DOI: 10.1055/s-0043-108551
 Fortschr Röntgenstr 2017; 189: 511–514
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 ISSN 1438–9029

Information

This statement is equivalent to statement no. 213 of the German Society of Gynecology and Obstetrics (DGGG).