

Transcervical Radiofrequency Ablation of Uterine Fibroids Global Registry (SAGE): Study Protocol and Preliminary Results

This article was published in the following Dove Press journal:
Medical Devices: Evidence and Research

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Background: Transcervical fibroid ablation (TFA) is a minimally invasive, effective treatment of symptomatic uterine fibroids that utilizes intrauterine ultrasound for imaging and radiofrequency energy for ablation. Outcomes reported with TFA have been positive, with significant reductions in fibroid volume, improvements in symptom severity and health-related quality of life, and low complication and surgical reintervention rates. The SAGE registry characterizes the long-term (5-year) outcomes of TFA when used to treat symptomatic uterine fibroids in real-world usage.

Methods/Design: SAGE is an ongoing postmarket global registry involving up to 50 sites and up to 500 women who select TFA with the Sonata system for treatment of symptomatic uterine fibroids. Patients are followed for 5 years. Main outcomes include symptom severity score and health-related quality of life subscales of the UFS-QoL, general health status on the EQ-5D, perceived treatment benefit, treatment satisfaction, work and activity patterns, overall patient treatment outcome, adverse events, pregnancy incidence and outcomes, and surgical reinterventions for heavy menstrual bleeding.

Discussion: The SAGE registry represents the largest known study of TFA for uterine fibroids and will generate up to 2500 patient-years of outcome data. Preliminary results from the first 160 treated women suggest broad applicability of TFA to a wide range of fibroid types and sizes and an excellent safety profile, with a device-related adverse event rate of 0.6% and a serious procedure-related adverse event rate of 0.6%. Of the 241 fibroids treated, 10% were submucous, 52% transmural, 28% intramural, and 10% subserous. Ablated fibroid diameters ranged from <1 cm to >10 cm, with 27% of fibroids having maximum diameters >5 cm. The real-world experience from SAGE will strengthen the existing evidence on the durability of TFA in providing meaningful relief from uterine fibroid symptoms and will have important clinical and economic implications for patients, physicians, and healthcare payers.

Trial Registration: <https://clinicaltrials.gov>, NCT03118037. Registered on 18 April 2017.

Keywords: leiomyoma, myoma, radiofrequency ablation, SAGE, Sonata, transcervical fibroid ablation, uterine fibroid

Background

Uterine fibroids are highly prevalent benign soft tissue tumors of the female genital tract, detectable in approximately 70% of White women and over 80% of Black women by the age of 50 in the United States.¹ In Germany, it has been estimated that over 60% of women have fibroids by age 50.^{2,3} Approximately 20–50% of women with fibroids in the United States and nearly 55% of women in Germany

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report associated symptoms that may include heavy menstrual bleeding (HMB) and/or bulk symptoms that are sufficiently bothersome to interfere with activities of daily living and work productivity.^{2,4} Self-management of symptoms with medication or lifestyle modification is commonly attempted prior to seeking medical care, but these initial symptom management techniques are often unsuccessful.⁵ Medical therapy for fibroids, most notably the GnRH analogues, is not a long-term solution, is costly, and associated with side effects relating to pseudomenopause such as osteoporosis.

Hysterectomy provides definitive resolution of fibroid symptoms; yet there is growing concern that this surgery for fibroid treatment is overutilized⁶ and patients are increasingly interested in less invasive, uterus-preserving treatment options associated with shorter recovery times and less disruption to daily life.⁵ Myomectomy and uterine artery embolization are uterus-preserving alternatives to hysterectomy that are appropriate for many patients. Indeed, hysterectomy is rarely surgically or clinically required for fibroid treatment, as myomectomy is nearly always technically feasible although the risks associated with occult sarcoma and the attendant risks of morcellation may influence decision-making with regard to management. However, depending on the surgical route, myomectomy can be associated with a significant recovery period and potential morbidity, while generally requiring Cesarean section for future deliveries. Uterine artery embolization is effective but can be associated with significant morbidity due to pain, uterine inflammation (post-embolization syndrome), chronic leukorrhea secondary to fibroid sloughing, and rarely, uterine necrosis and mortality. It has also been associated with concerns about ovarian senescence and decreased fertility potential, as well as intrauterine adhesions. Additionally, these treatments are associated with surgical reintervention rates as high as 24% at 2 years,^{7–10} which limits their usefulness as a long-term solution to fibroid management. There remains a clinical need for a less invasive, uterus-conserving treatment for fibroids that provides durable symptom resolution.

To better address the needs of this underserved patient population, an incisionless, uterus-preserving outpatient procedure has been developed that uses intrauterine ultrasound to locate the fibroids and radiofrequency energy to treat them. The technique, transcervical fibroid ablation (TFA) with the Sonata system, has been shown to be effective and safe in several prospective, multicenter

studies. One-, two-, three- and five-year outcome data have been published, demonstrating durable symptom relief and improvements in quality of life along with high rates of patient satisfaction.^{11,12} Among these studies was an FDA prospective, multicenter, controlled pivotal trial of 147 patients treated with TFA who were followed for 3 years,¹¹ and the VITALITY study that followed patients for a mean of 5.4 years.¹² The cumulative evidence demonstrating the durability of this procedure in providing meaningful relief from uterine fibroid symptoms will be further strengthened by long-term follow-up data after treatment with TFA in the real-world clinical practice setting, hence the impetus for the Transcervical Radiofrequency Ablation of Uterine Fibroids Global Registry (SAGE registry).

Methods/Design

Objective

The purpose of the SAGE registry is to collect long-term outcome data in a large cohort of women who received TFA for uterine fibroids in real-world clinical practice.

Study Design

The SAGE registry was registered at ClinicalTrials.gov (NCT03118037) before first patient enrollment. SAGE is a prospective, multicenter, observational postmarket registry that will enroll up to 500 patients treated with TFA at up to 50 sites in Europe. Eligible patients have selected TFA with the Sonata system for treatment of symptomatic uterine fibroids. Patient recruitment began in June 2017. The total study duration is expected to be 7.5 years, with 2.5 years of patient recruitment and 5 years of follow-up. The protocol for this study was developed in accordance with the SPIRIT 2013 guidance for protocols of clinical trials.¹³

Participants and Eligibility Criteria

The eligibility criteria for registry enrollment are purposely broad in order to reflect real-world device usage in a heterogenous sample of women. Eligible patients are adult women (age ≥ 18 years) with symptomatic uterine fibroids who select TFA for treatment of their fibroids and demonstrate a willingness to participate in the registry and to return for specified follow-up visits. No limitations are imposed on maximum patient age, fibroid type, fibroid size, prior surgical history, baseline symptom severity, or desire for future fertility. Following protocol approval from the

ethics committee at each investigative site, investigators prospectively screen and enroll eligible patients in consecutive fashion. Patient recruitment is mainly facilitated by gynecologist investigators at each center who inform women about the study and refer potentially eligible women to dedicated research staff to receive additional information. All patients are informed of the potential risks and benefits of study participation and provided written informed consent before any study procedures are initiated. Pre-treatment patient assessments include demographics, medical and surgical history, imaging of the uterus to characterize the number, size, and type of fibroids, and questionnaires including the EuroQol 5-Dimension (EQ-5D) questionnaire, Uterine Fibroid Symptom and Quality-of-Life (UFS-QoL) questionnaire, and the Work Productivity and Activity Impairment Questionnaire: Specific Health Problem (WPAI:SHP).

Transcervical Fibroid Ablation Procedure

The sonography-guided TFA procedure treatment device (Sonata[®] system, Gynesonics, Inc., Redwood City, CA, United States) includes an integrated intrauterine sonography probe and radiofrequency ablation handpiece that allows the gynecologist to identify, target, and ablate uterine fibroids (Figure 1). The integration of real-time ultrasound imaging enables the physician to visualize, target, and ablate a greater range of fibroids than could be approached via operative hysteroscopy. A graphical interface is displayed on the live ultrasound image that identifies the target ablation area and the extent of sub-ablative thermal heating. The gynecologist utilizes this information to confirm the ablation is within the fibroid while confining the thermal safety border to within the uterine serosa. A single ablation may suffice to treat a fibroid, but



Figure 1 The Sonata transcervical fibroid ablation device. The device consists of an integrated intrauterine sonography probe with a radiofrequency ablation handpiece. Image courtesy from Gynesonics, Inc with permission.

additional ablations may be necessary depending on fibroid size, location, and geometry. TFA is an outpatient procedure, and has been performed in operating rooms, ambulatory surgery centers, and physician offices. General anesthesia is not required; the type of anesthesia may thus be individualized and has included regional anesthesia, conscious sedation and multimodal local anesthesia. As reported in the SONATA clinical trial,¹⁴ TFA is associated with very high patient tolerance and low pain scores.

Outcome Measures

Follow-up visits occur at 4 weeks (± 2 weeks) post-treatment, and annually thereafter (± 2 months) for 5 years post-treatment. A schedule of patient assessments during the study is provided in Table 1. Symptom severity and quality of life are assessed with the symptom severity score (SSS) and health-related quality of life (HRQL) subscales of the UFS-QoL questionnaire.¹⁵ Scores are reported on a 0 to 100 scale where higher SSS scores indicate more severe symptoms and lower HRQL scores indicate worse quality of life. Changes in general health status are assessed with the EQ-5D questionnaire, which consists of five questions that provide a description of the patient's health state with scores ranging from 0 (indicating death) to 1 (indicating perfect health). At each follow-up visit, patients self-report their perceived treatment benefit as improved, no change, or worsened relative to baseline. Treatment satisfaction is measured on a 6-item scale ranging from very satisfied to very dissatisfied. The WPAI:SHP questionnaire¹⁶ assesses change in work and activity patterns following treatment and is designed with a 28-day recall period to capture the impact of uterine fibroids over a full menstrual cycle. Overall patient treatment outcome is assessed using the Overall Treatment Effect (OTE) scale. Safety outcomes are reported by adverse events (AE) and the classifications of seriousness and relatedness to device or procedure per the ISO 14155 standard. All AEs related to device or procedure, including those considered to be serious, are reported. A serious AE results in a) a life-threatening illness or injury, b) a permanent impairment of a body structure or a body function, c) an in-patient hospitalization or prolongation of an existing hospitalization of more than 24 hours, or d) a medical or surgical intervention to prevent life threatening illness or injury or permanent impairment to a body structure or a body function. The incidence and outcome of pregnancies are collected. Lastly, occurrence

Table I Schedule of Study Assessments

Assessment	Screening/Baseline	Treatment	4 Weeks	1 Year	2 Years	3 Years	4 Years	5 Years
Informed consent	X							
Demographics; medical history	X							
Uterus/fibroid imaging	X							
TFA treatment		X						
Treatment recovery questionnaire			X					
EQ-5D	X			X	X	X	X	X
UFS-QOL	X			X	X	X	X	X
WPAI:SHP	X			X	X	X	X	X
Subject satisfaction				X	X	X	X	X
OTE				X	X	X	X	X
Pregnancy status and outcome			X	X	X	X	X	X
Adverse events		X	X	X	X	X	X	X
Surgical reintervention for HMB			X	X	X	X	X	X

Abbreviations: EQ-5D, EuroQol 5-Dimension; HMB, heavy menstrual bleeding; OTE, overall treatment effect; TFA, transcervical fibroid ablation; UFS-QoL, Uterine Fibroid Symptom and Quality-of-Life; WPAI:SHP, Work Productivity and Activity Impairment Questionnaire: Specific Health Problem.

of surgical reintervention for HMB due to fibroids are collected.

Data Quality Assurance

All research procedures planned in this study are in accordance with a protocol that is approved by appropriate institutional or regional ethics committees. Each investigative site is provided with a didactic review of the protocol and procedures. Study training emphasizes the protocol and study compliance, compliance with applicable regulations, patient selection criteria, principals of good clinical practice, and data quality and monitoring. Investigators and coordinators enter clinical study data into an electronic data capture system. Each system user is trained and assigned a unique username and password to ensure data protection and security. Each patient is assigned a unique identification number to ensure confidentiality prior to data entry into the electronic database. Monitoring visits to the clinical sites are made periodically to ensure that the study is conducted in accordance with the protocol and that the clinical data are validated against source documentation at the investigative site. Original source documents are reviewed for verification of data recorded on the electronic case report forms and in the electronic database. Site investigators report safety outcomes to their respective ethics committee according to national regulations and local ethics committee requirements.

Additionally, all serious AEs are reported to the study sponsor at SAGE.safety@gynesonics.com immediately after the site becomes aware of the event.

Statistical Methods

A power analysis was not performed for this observational registry; the sample size was determined based on pragmatic factors including the number of participating sites and the anticipated patient enrollment at each site. The safety analysis population consists of all patients who provide informed consent and receive treatment with TFA. Baseline patient characteristics are summarized with standard descriptive statistics. Categorical variables are described with percents and counts. Continuous variables are described with means and standard deviations, or medians and ranges. Missing data imputation will not be performed.

Ethics and Dissemination

The research methods adhered to the guidelines set forth in the Declaration of Helsinki. The protocol for this registry was approved by the ethics committees at each participating site. Study coordinators explained the research procedures to each participant prior to enrollment and all patients provided informed consent before study participation. Important protocol modifications will be posted to <https://clinicaltrials>.

gov, NCT03118037. All personal information about registry participants remain confidential and will only be reported in aggregate format such that patient confidentiality is protected before, during, and after the study. There are no provisions in the protocol for post-trial care. Following participation in the study, patients may seek medical care with any provider they choose, and clinical results will not be collected following patient withdrawal. The study results will be submitted for conference presentations and for publication in peer-reviewed journals.

Preliminary Results

Between June 2017 and February 2020, 160 women with symptomatic uterine fibroids received TFA treatment in the SAGE registry at 7 sites in Germany and Switzerland (Table 2). Additional sites are currently in start-up phase in the United Kingdom. The mean patient age was 42 ± 7 years, mean body mass index was 24 ± 5 kg/m², and 19% of patients previously underwent an intrauterine procedure (most commonly myomectomy). The mean follow-up duration thus far is 5.3 ± 6.0 months, with the maximum follow-up currently at 25

months. A total of 271 fibroids were identified sonographically, but not ablated. The reasons for lack of ablation included close proximity of the serosa precluding effective/safe ablation, physician choice to use hysteroscopic resection as a concomitant therapy, fibroids that were not deemed material to the patient's symptomatology per the treating physician (eg, small subserosal myoma), and fibroids that were not appropriate for TFA (eg, pedunculated fibroids). Characteristics of the 241 treated uterine fibroids are reported in Figure 2 and included a broad range of fibroid types and sizes. Specifically, 10% of treated fibroids were submucosal, 52% transmural, 28% intramural, and 10% subserosal. Ablated fibroid diameters ranged from <1 cm in maximum diameter to >10 cm, with more than 27% of the treated fibroids >5 cm. No serious device-related AE has been reported. Non-serious procedure-related AEs were reported in 8 (5.0%) patients and were attributable to pain or vaginal bleeding. One (0.6%) device related AE was reported as a second-degree skin burn at the site of the dispersive electrodes due to human error/misuse by the site staff. One (0.6%) serious procedure-related AE was reported; non-infectious endometritis that resolved with

Table 2 List of Participating Sites and Principal Investigators in the SAGE Registry*

Site Name	EC Name/Ref. Number	Site Location	Principal Investigator
Frauenklinik Universitätsklinikum Jena	Ethik-Kommission der Friedrich-Schiller-Universität Jena Reference #: 12/16/4996	Jena, Germany	Ingo Runnebaun
MarienKrankenhaus Schwerte	Ethikkommission der Ärztekammer Westfalen-Lippe Reference #: 2017-1899-b-S	Schwerte, Germany	Michael Hartmann
Evangelisches Krankenhaus Köln-Weyertal gemeinnützige GmbH	Ärztekammer Nordrhein Reference #: 2,017,188	Köln, Germany	Thomas Römer
Klinikverbund Kempten-Oberallgäu gGmbH	Ethikkommission der Universität Ulm Reference #: 132/17 - CL/bal.	Kempten, Germany	Ricardo Felderbaum
Marien Hospital Witten	Ethikkommission der Universität Witten/Herdecke Reference #: 115/2017	Witten, Germany	Sven Schiermeier
Josephs-Hospital Warendorf	Ethikkommission der Ärztekammer Westfalen-Lippe Reference #: 2017-1899-b-S	Warendorf, Germany	Matthias Engelhardt
Spital Oberengadin	Kantonale Ethikkommission Zürich Reference #: Gesuch BASEC-Nr.: 2018-01461	Samedan, Switzerland	Ladina Christoffel
Liverpool Women's NHS Foundation Trust**	Liverpool Women's Foundation Trust Fund Reference #: not assigned	Liverpool, UK	Adel Soltan
Addenbrooke's Hospital**	Pending- Reference #: not assigned	Cambridge, UK	Mariam Baumgarten

Notes: *List current through February 2020. **Site in study start-up phase.

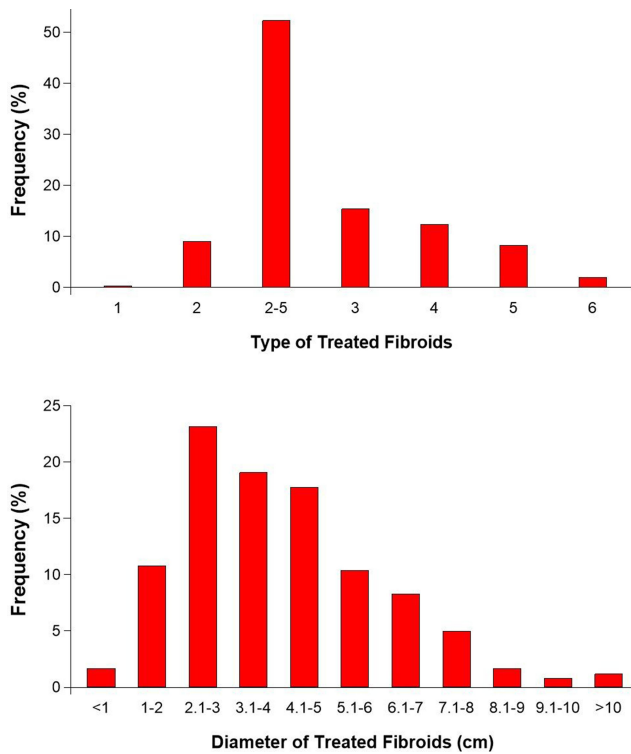


Figure 2 Uterine fibroid characteristics among the first 160 women enrolled in the SAGE registry. Histograms represent the frequency of fibroids types (top panel) and diameters (bottom panel) that have been treated with transcervical fibroid ablation.

intravenous antibiotics and analgesia. As the event involved an overnight hospital stay, it was classified as a serious AE (Table 3).

Discussion

The collective clinical experience with TFA demonstrates that the procedure is a safe and effective option for women seeking a minimally invasive treatment for symptomatic uterine fibroids. In the SONATA trial of TFA, Miller et al reported favorable outcomes among 147 women treated with TFA and followed for 2 years.¹¹ Compared to baseline, the SSS decreased from 55±19 to 23±19 (p<0.001), HRQL increased from 40±21 to 83±19 (p<0.001), and EQ-5D scores increased from 0.72±0.21 to 0.89±0.14 (p<0.001). Overall treatment satisfaction at 2 years was 94%. The mean percentage of missed work time, overall work impairment, and activity impairment significantly decreased at follow-up. Through 2 years, surgical reintervention for HMB was performed in 5.6% of patients. In the VITALITY study by Garza-Leal et al that followed patients over a mean of 5.4 years,¹² the SSS decreased from 65±17 to 28±36, HRQL improved from 27±22 to 76 ±33, and the surgical reintervention rate was 11.8%. For

Table 3 Preliminary Safety Results in the SAGE Registry

Variable	Value
Patient characteristics	
No. patients	160
No. fibroids treated	241
Age, yr	42±7 (159)*
Fibroid type	
Transmural (FIGO 2–5)	52% (126/241)
Intramural (FIGO 3 or 4)	28% (67/241)
Submucosal (FIGO 1 or 2)	10% (23/241)
Subserosal (FIGO 5 or 6)	10% (25/241)
Procedural data	
Procedure time, min	25±14 (160)*
Ablated fibroid diameter, cm	4.2±2.2 (241)*
Ablated fibroid/patient	1.5±0.9 (160)*
Adverse events	
Follow-up duration, mo*	1.3 (0.1, 25.0)*
Any procedure-related AE	5.0% (8/160)
Any serious procedure-related AE	0.6% (1/160)
Non-infectious endometritis	1
Any nonserious procedure-related AE	5.0% (8/160)**
Cramping/pain	5
Vaginal bleeding	4
Fibroid shedding	1
Flu-like symptoms	1
Nausea/vomiting	1
Other	1
Any nonserious device-related AE	0.6% (1/160)
Skin burn***	1
Any serious device-related AE	0% (0/160)

Notes: *Data reported as mean±standard deviation (n) or median (min, max). **13 events reported in 8 patients. ***Second degree skin burn due to human error/misuse by the site staff.

Abbreviation: FIGO, International Federation of Gynecology and Obstetrics.

comparison, a study using a large commercial payor database reported rates of reintervention over 2 years of follow-up as 23.5% for uterine artery embolization, 18% for hysteroscopic myomectomy, 19% for endometrial ablation, and 8% for laparoscopic myomectomy.^{8,17} Thus,

TFA provides a viable treatment alternative to the aforementioned procedures that may provide more durable fibroid-related symptom relief.

The preliminary results from the first 160 treated women in SAGE suggest broad applicability of TFA to a wide range of fibroid types and sizes and an excellent safety profile. These initial results corroborate those from previous clinical trials, as well as over 600 commercial cases worldwide. The wide repertoire of fibroid types treated in SAGE, including transmural, intramural, and subserosal fibroids, are not generally accessible to operative hysteroscopy. The registry will follow women for 5 years, after which a comprehensive characterization of long-term outcomes of TFA will be possible. SAGE represents the largest known study of TFA for uterine fibroids and will generate up to 2500 patient-years of long-term outcomes data. Final results from the SAGE registry are anticipated to be published in 2026. The outcomes derived from SAGE will further strengthen the existing evidence on the durability of TFA in providing meaningful relief from uterine fibroid symptoms and will have important clinical and economic implications for patients, physicians and healthcare payers.

Data Sharing Statement

Data derived from this research will not be made available as these are preliminary results from an ongoing registry.

Acknowledgments

The authors acknowledge David B. Toub, MD, MBA, and Taraneh G. Farazi, PhD for their review of this article, and Larry Miller, PhD for editorial assistance.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Funding

Funding for the study was provided by Gynesonics, Inc. (Redwood City, CA, United States). Gynesonics, Inc. provided research support to the participating institutions in the SAGE registry.

Disclosure

The authors report no competing interests.

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