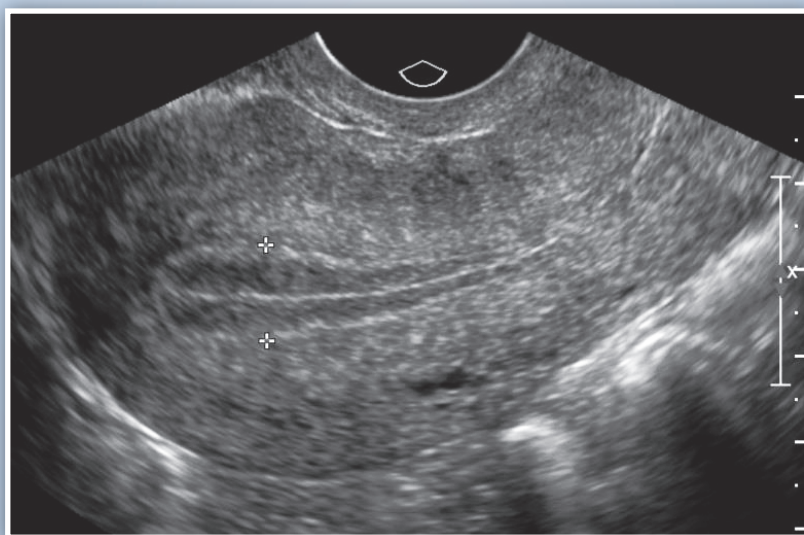


AIUM Practice Parameter for the Performance of

# Ultrasound of the Female Pelvis

*Parameter developed in collaboration with the American College of Radiology (ACR),  
the American College of Obstetricians and Gynecologists (ACOG), the Society for Pediatric  
Radiology (SPR), and the Society of Radiologists in Ultrasound (SRU).*



The American Institute of Ultrasound in Medicine (AIUM) is a multidisciplinary association dedicated to advancing the safe and effective use of ultrasound in medicine through professional and public education, research, development of parameters, and accreditation. To promote this mission, the AIUM is pleased to publish, in conjunction with the American College of Radiology (ACR), the American College of Obstetricians and Gynecologists (ACOG), the Society for Pediatric Radiology (SPR), and the Society of Radiologists in Ultrasound (SRU), this *AIUM Practice Parameter for the Performance of Ultrasound of the Female Pelvis*. We are indebted to the many volunteers who contributed their time, knowledge, and energy to bringing this document to completion.

The AIUM represents the entire range of clinical and basic science interests in medical diagnostic ultrasound, and, with hundreds of volunteers, the AIUM has promoted the safe and effective use of ultrasound in clinical medicine for more than 50 years. This document and others like it will continue to advance this mission.

Practice parameters of the AIUM are intended to provide the medical ultrasound community with parameters for the performance and recording of high-quality ultrasound examinations. The parameters reflect what the AIUM considers the minimum criteria for a complete examination in each area but are not intended to establish a legal standard of care. AIUM-accredited practices are expected to generally follow the parameters with recognition that deviations from these parameters will be needed in some cases, depending on patient needs and available equipment. Practices are encouraged to go beyond the parameters to provide additional service and information as needed.



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## I. Introduction

The clinical aspects contained in specific sections of this parameter (Introduction, Indications, Specifications of the Examination, and Equipment Specifications) were developed collaboratively by the American Institute of Ultrasound in Medicine (AIUM), the American College of Radiology (ACR), the American College of Obstetricians and Gynecologists (ACOG), the Society for Pediatric Radiology (SPR), and the Society of Radiologists in Ultrasound (SRU). Recommendations for physician requirements, written request for the examination, documentation, and quality control vary among the organizations and are addressed by each separately.

This parameter has been developed to assist physicians performing sonographic studies of the female pelvis. Ultrasound examinations of the female pelvis should be performed only when there is a valid medical reason, and the lowest possible ultrasonic exposure settings should be used to gain the necessary diagnostic information. In some cases, additional or specialized examinations may be necessary. Although it is not possible to detect every abnormality, adherence to the following parameter will maximize the probability of detecting most abnormalities. For ultrasound examinations of the urinary bladder, see the *AIUM Practice Parameter for the Performance of an Ultrasound Examination of the Abdomen and/or Retroperitoneum*.

## II. Indications

Indications for pelvic sonography include but are not limited to:

1. Evaluation of pelvic pain;
2. Evaluation of pelvic masses;
3. Evaluation of endocrine abnormalities, including polycystic ovaries;
4. Evaluation of dysmenorrhea (painful menses);
5. Evaluation of amenorrhea;
6. Evaluation of abnormal bleeding;
7. Evaluation of delayed menses;
8. Follow-up of a previously detected abnormality;
9. Evaluation, monitoring, and/or treatment of infertility patients;
10. Evaluation in the presence of a limited clinical examination of the pelvis;
11. Evaluation for signs or symptoms of pelvic infection;
12. Further characterization of a pelvic abnormality noted on another imaging study;
13. Evaluation of congenital uterine and lower genital tract anomalies;
14. Evaluation of excessive bleeding, pain, or signs of infection after pelvic surgery, delivery, or abortion;
15. Localization of an intrauterine contraceptive device;
16. Screening for malignancy in high-risk patients;
17. Evaluation of incontinence or pelvic organ prolapse;
18. Guidance for interventional or surgical procedures; and
19. Preoperative and postoperative evaluation of pelvic structures.

### III. Qualifications of Personnel

See [www.aium.org](http://www.aium.org) for AIUM Official Statements including *Standards and Guidelines for the Accreditation of Ultrasound Practices* and relevant Physician Training Guidelines.

### IV. Written Request for the Examination

The written or electronic request for an ultrasound examination should provide sufficient information to allow for the appropriate performance and interpretation of the examination.

The request for the examination must be originated by a physician or other appropriately licensed health care provider or under the provider's direction. The accompanying clinical information should be provided by a physician or other appropriate health care provider familiar with the patient's clinical situation and should be consistent with relevant legal and local health care facility requirements.

### V. Specifications of the Examination

The following sections detail the examination to be performed for each organ and anatomic region in the female pelvis. All relevant structures should be identified by the transabdominal and/or transvaginal approach. A transrectal or transperineal approach may be useful in patients who are not candidates for introduction of a vaginal probe and in assessing the patient with pelvic organ prolapse. More than 1 approach may be necessary.<sup>1,2</sup>

#### A. General Pelvic Preparation

For a complete transabdominal pelvic sonogram, the patient's bladder can be distended if necessary to displace the small bowel from the field of view. Occasionally, overdistention of the bladder may compromise the evaluation. When this occurs, imaging may be repeated after partial bladder emptying. If an abnormality of the urinary bladder is detected, it should be documented and reported.

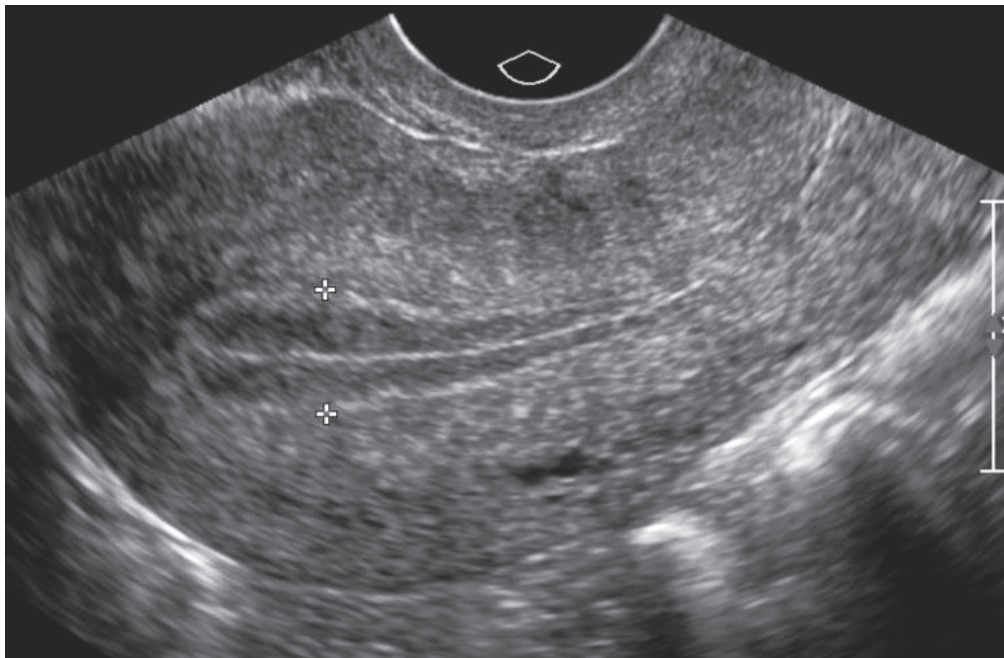
For a transvaginal sonogram, the urinary bladder is preferably empty. The patient, the sonographer, or the physician may introduce the vaginal transducer, preferably under real-time monitoring. Consideration of having a chaperone present should be in accordance with local policy.<sup>3</sup>

#### B. Uterus

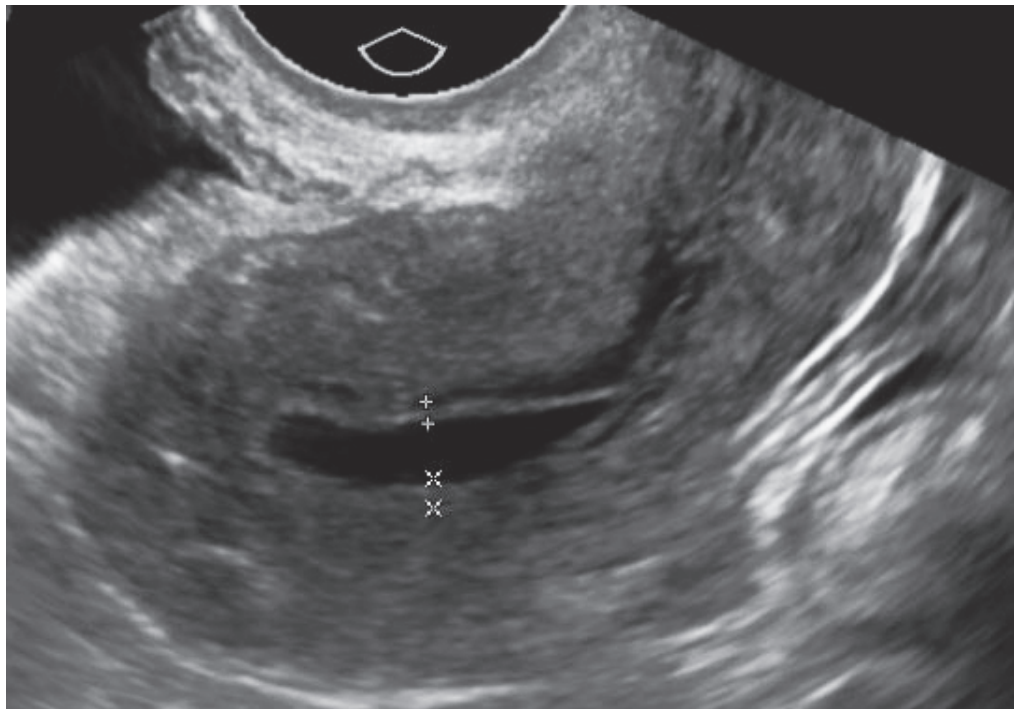
The vagina and uterus provide anatomic landmarks that can be used as reference points for the other pelvic structures, whether normal or abnormal. In examining the uterus, the following should be evaluated: (1) the uterine size, shape, and orientation; (2) the endometrium; (3) the myometrium; and (4) the cervix. The vagina may be imaged as a landmark for the cervix. The overall uterine length is evaluated in a sagittal view from the fundus to the cervix (to the external os, if it can be identified). The depth of the uterus (anteroposterior dimension) is measured in the same sagittal view from its anterior to posterior walls, perpendicular to the length. The maximum width is measured in the transverse or coronal view. If volume measurements of the uterine corpus are performed, the cervical component should be excluded from the uterine length measurement.

Abnormalities of the uterus should be documented.<sup>4</sup> The myometrium and cervix should be evaluated for contour changes, echogenicity, masses, and cysts. Masses that may require follow-up or intervention should be measured in at least 2 dimensions, acknowledging that it is not usually necessary to measure all uterine fibroids. The size and location of clinically relevant fibroids should be documented.

The endometrium should be analyzed for thickness, focal abnormalities, echogenicity, and the presence of fluid or masses in the cavity. The thickest part of the endometrium should be measured perpendicular to its longitudinal plane in the anteroposterior diameter from echogenic to echogenic border (Figure 1). The adjacent hypoechoic myometrium and fluid in the cavity should be excluded (Figure 2). Assessment of the endometrium should allow for variations expected with phases of the menstrual cycle and with hormonal supplementation.<sup>5-7</sup> It should be reported if the endometrium is not adequately seen in its entirety or is poorly defined. Sonohysterography may be a useful adjunct to evaluate the patient with abnormal uterine bleeding or to further clarify an abnormally thickened endometrium. (See the *AIUM Practice Parameter for the Performance of Sonohysterography*.) If the patient has an intrauterine contraceptive device, its location should be documented.



**Figure 1.** Measurement of endometrial thickness. The endometrial thickness is measured in its thickest portion from echogenic to echogenic border (calipers) perpendicular to the midline longitudinal plane of the uterus.



**Figure 2.** Measurement of endometrium with fluid in the cavity. In the presence of endometrial fluid, measurements of the 2 separate layers of the endometrium (calipers), excluding the fluid, are added to determine the endometrial thickness.

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The addition of 3-dimensional to 2-dimensional ultrasound (transabdominal, transvaginal, transperineal, and/or transrectal) can be helpful in many circumstances, including but not limited to evaluating the relationship of masses with the endometrial cavity, identifying uterine congenital anomalies and a thickened and/or heterogeneous endometrium, and evaluating the location of an intrauterine device and the integrity of the pelvic floor.<sup>8,9</sup>

### C. Adnexa, Including Ovaries and Fallopian Tubes

When evaluating the adnexa, an attempt should be made to identify the ovaries first, since they can serve as a major point of reference for assessing the presence of adnexal pathology. The ovarian size may be determined by measuring the ovary in 3 dimensions (width, length, and depth) on views obtained in 2 orthogonal planes.<sup>10</sup> Any ovarian abnormalities should be documented.<sup>11-15</sup>

The ovaries may not be identifiable in some patients. This occurs most frequently before puberty, after menopause, or in the presence of a large leiomyomatous uterus. The adnexal region should be surveyed for abnormalities, particularly masses and dilated tubular structures.

If an adnexal abnormality is noted, its relationship with the ovaries and uterus should be assessed. The size and sonographic characteristics of adnexal masses should be documented.

Spectral, color, and/or power Doppler ultrasound may be useful to evaluate the vascular characteristics of pelvic lesions.<sup>15-19</sup>

## D. Cul-de-Sac

The cul-de-sac and bowel posterior to the uterus may not be clearly defined. This area should be evaluated for the presence of free fluid or a mass. If a mass is detected, its size, position, shape, sonographic characteristics, and relationship to the ovaries and uterus should be documented. Differentiation of normal loops of bowel from a mass may be difficult if only a trans-abdominal examination is performed. A transvaginal examination may be helpful to distinguish a suspected mass from fluid and feces within the normal rectosigmoid colon.

## VI. Documentation

Adequate documentation is essential for high-quality patient care. There should be a permanent record of the ultrasound examination and its interpretation. Images of all appropriate areas, both normal and abnormal, should be recorded. Variations from normal size should be accompanied by measurements. Images should be labeled with the patient identification, facility identification, examination date, and side (right or left) of the anatomic site imaged. An official interpretation (final report) of the ultrasound findings should be included in the patient's medical record. Retention of the ultrasound examination should be consistent both with clinical needs and with relevant legal and local health care facility requirements.

Reporting should be in accordance with the *AIUM Practice Parameter for Documentation of an Ultrasound Examination*.

## VII. Equipment Specifications

A sonographic examination of the female pelvis should be conducted with a real-time scanner, preferably using sector, curved linear, and/or endovaginal transducers. The transducer or scanner should be adjusted to operate at the highest clinically appropriate frequency, realizing that there is a trade-off between resolution and beam penetration.<sup>3</sup>

## VIII. Quality Control and Improvement, Safety, Infection Control, and Patient Education

Policies and procedures related to quality control, patient education, infection control, and safety should be developed and implemented in accordance with the *AIUM Standards and Guidelines for the Accreditation of Ultrasound Practices*.

Equipment performance monitoring should be in accordance with the *AIUM Standards and Guidelines for the Accreditation of Ultrasound Practices*.

## IX. ALARA Principle

The potential benefits and risks of each examination should be considered. The ALARA (as low as reasonably achievable) principle should be observed when adjusting controls that affect the acoustic output and by considering transducer dwell times. Further details on ALARA may be found in the AIUM publication *Medical Ultrasound Safety, Third Edition*.

## Acknowledgments

This parameter was revised by the AIUM in collaboration with the American College of Radiology (ACR), the American College of Obstetricians and Gynecologists (ACOG), the Society for Pediatric Radiology (SPR), and the Society of Radiologists in Ultrasound (SRU) according to the process described in the AIUM *Clinical Standards Committee Manual*.

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