

CURRENT CLINICAL IMAGING • RICHARD C. SEMELKA, SERIES EDITOR

# WOMEN'S IMAGING

**MRI with Multimodality Correlation**

EDITED BY

MICHELE A. BROWN

HAYDEE OJEDA-FOURNIER

DRAGANA DJILAS

MOHAMED EL-AZZAZI

RICHARD C. SEMELKA



**WILEY** Blackwell

# Women's Imaging MRI with Multimodality Correlation

Edited by

**Michele A. Brown, MD**

Professor of Clinical Radiology  
UC San Diego Health System  
San Diego, CA, USA

**Haydee Ojeda-Fournier, MD**

Associate Professor of Clinical Radiology  
Medical Director, Breast Imaging Section  
Director Medical Student Education in Radiology  
Moores Cancer Center  
UC San Diego Health System  
La Jolla, CA, USA

**Dragana Djilas, MD, PhD**

Associate Professor of Radiology  
Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Mohamed El-Azzazi MD, PhD**

Clinical Research Scholar, MRI Section, Department of Radiology  
University of North Carolina Chapel Hill, NC, USA;  
Professor of Radiology, Al-Azhar University

Associate Professor of Radiology  
Dammam University, Saudi Arabia; and  
Consultant in Radiology  
King Fahad University Hospital, Saudi Arabia

**Richard C. Semelka, MD**

Director, Magnetic Resonance Services;  
Professor, Vice Chairman of Clinical Research; and  
Vice Chairman of Quality and Safety  
Department of Radiology  
University of North Carolina at Chapel Hill  
Chapel Hill, NC, USA

**WILEY Blackwell**

Copyright © 2014 by John Wiley & Sons, Inc. All rights reserved

Published by John Wiley & Sons, Inc., Hoboken, New Jersey  
Published simultaneously in Canada

No part of this publication may be reproduced, stored in a retrieval system, or transmitted in any form or by any means, electronic, mechanical, photocopying, recording, scanning, or otherwise, except as permitted under Section 107 or 108 of the 1976 United States Copyright Act, without either the prior written permission of the Publisher, or authorization through payment of the appropriate per-copy fee to the Copyright Clearance Center, Inc., 222 Rosewood Drive, Danvers, MA 01923, (978) 750-8400, fax (978) 750-4470, or on the web at [www.copyright.com](http://www.copyright.com). Requests to the Publisher for permission should be addressed to the Permissions Department, John Wiley & Sons, Inc., 111 River Street, Hoboken, NJ 07030, (201) 748-6011, fax (201) 748-6008, or online at <http://www.wiley.com/go/permissions>.

The contents of this work are intended to further general scientific research, understanding, and discussion only and are not intended and should not be relied upon as recommending or promoting a specific method, diagnosis, or treatment by health science practitioners for any particular patient. The publisher and the author make no representations or warranties with respect to the accuracy or completeness of the contents of this work and specifically disclaim all warranties, including without limitation any implied warranties of fitness for a particular purpose. In view of ongoing research, equipment modifications, changes in governmental regulations, and the constant flow of information relating to the use of medicines, equipment, and devices, the reader is urged to review and evaluate the information provided in the package insert or instructions for each medicine, equipment, or device for, among other things, any changes in the instructions or indication of usage and for added warnings and precautions. Readers should consult with a specialist where appropriate. The fact that an organization or Website is referred to in this work as a citation and/or a potential source of further information does not mean that the author or the publisher endorses the information the organization or Website may provide or recommendations it may make. Further, readers should be aware that Internet Websites listed in this work may have changed or disappeared between when this work was written and when it is read. No warranty may be created or extended by any promotional statements for this work. Neither the publisher nor the author shall be liable for any damages arising herefrom.

For general information on our other products and services or for technical support, please contact our Customer Care Department within the United States at (800) 762-2974, outside the United States at (317) 572-3993 or fax (317) 572-4002.

Wiley also publishes its books in a variety of electronic formats. Some content that appears in print may not be available in electronic formats. For more information about Wiley products, visit our web site at [www.wiley.com](http://www.wiley.com).

Library of Congress Cataloging-in-Publication Data:

Women's imaging : MRI with multimodality correlation / edited by Michele A. Brown, Haydee Ojeda-Fournier, Dragana Djilas, Mohamed El-Azzazi, Richard C. Semelka.

p. ; cm.

Includes bibliographical references and index.

ISBN 978-1-118-48284-1 (cloth)

I. Brown, Michele A., editor of compilation. II. Ojeda-Fournier, Haydee, editor of compilation. III. Djilas, Dragana, editor of compilation. IV. El-Azzazi, Mohamed, editor of compilation. V. Semelka, Richard C., editor of compilation.

[DNLM: 1. Magnetic Resonance Imaging—methods. 2. Women's Health. WN 185]

RC386.6.M34

616.07'548—dc23

2013042717

Cover design by Wiley

Printed in Singapore

10 9 8 7 6 5 4 3 2 1

## Women's Imaging



# Contributors

**Chayanin Angthong, MD**

Department of Radiology  
University of North Carolina at Chapel Hill  
Chapel Hill, NC, USA

**Jasmina Boban, MD**

Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Dragana Bogdanovic-Stojanovic, MD, PhD**

Assistant Professor of Radiology  
Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Michele A. Brown, MD**

Professor of Clinical Radiology  
UC San Diego Health System  
San Diego, CA, USA

**Julie Bykowski, MD**

Assistant Professor of Clinical Radiology  
Moores Cancer Center  
UC San Diego Health System  
La Jolla, CA, USA

**Jade de Guzman, MD**

Assistant Professor of Clinical Radiology  
Moores Cancer Center  
UC San Diego Health System  
La Jolla, CA, USA

**Dragana Djilas, MD, PhD**

Associate Professor of Radiology  
Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Mohamed El-Azzazi MD, PhD**

Clinical Research Scholar, MRI Section, Department of  
Radiology  
University of North Carolina, Chapel Hill, NC, USA;  
Professor of Radiology, Al-Azhar University  
Cairo, Egypt;  
Associate Professor of Radiology  
Dammam University, Saudi Arabia;  
Consultant in Radiology  
King Fahad University Hospital, Saudi Arabia

**Randy Fanous, MD, BHSc**

Department of Radiology  
UC San Diego Health System  
San Diego, CA, USA

**Michael J. Gabe, MD**

Department of Radiology  
UC San Diego Health System  
San Diego, CA, USA

**Vladimir Ivanovic, BSCEE**

Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Reena Malhotra**

Department of Radiology  
North Shore-LIJ Health System  
New Hyde Park, NY, USA

**Mary K. O'Boyle, MD**

Clinical Professor of Radiology  
Chief of Ultrasound  
UC San Diego Health System  
San Diego, CA, USA

**Haydee Ojeda-Fournier, MD**

Associate Professor of Clinical Radiology  
Medical Director, Breast Imaging Section  
Director Medical Student Education in Radiology  
Moore Cancer Center  
UC San Diego Health System  
La Jolla, CA, USA

**Dag Pavic, MD**

Associate Professor of Radiology  
Department of Radiology  
Medical University of South Carolina  
Charleston, SC, USA

**Natasa Prvulovic Bunovic, MD**

Consultant in Radiology  
Center for Diagnostic Imaging  
Oncology Institute of Vojvodina  
Novi Sad, Serbia

**Steven S. Raman, MD**

Professor of Radiology, Surgery and Urology  
David Geffen School of Medicine at UCLA  
Los Angeles, CA, USA

**Katherine M. Richman, MD**

Clinical Professor of Radiology  
UC San Diego Health System  
La Jolla, CA, USA

**Ryan C. Rockhill, MD**

Body and Breast Imaging  
Naval Medical Center San Diego  
San Diego, CA, USA

**Lorene E. Romine, MD**

Assistant Clinical Professor of Radiology  
UC San Diego Health System  
San Diego, CA, USA

**Laura E. Rueff, MD, MPH**

Department of Radiology  
University of California Los Angeles  
Los Angeles, CA, USA

**Richard C. Semelka, MD**

Director, Magnetic Resonance Services  
Professor, Vice Chairman of Clinical Research and  
Vice Chairman of Quality and Safety  
Department of Radiology  
University of North Carolina at Chapel Hill  
Chapel Hill, NC, USA

**Shannon St Clair, MD**

Department of Radiology  
UC San Diego Health System  
San Diego, CA, USA

# Preface

Women's health issues consume a large portion of medical resources and healthcare dollars. Proper management requires a team of physicians from various specialties. Within the field of Radiology, there has been a trend toward developing a subspecialty dedicated to comprehensive imaging of women's healthcare needs, including gynecological, obstetric, genitourinary, and breast conditions. The term "Women's Imaging" is used differently in different contexts; for the purpose of this textbook, the term is used to describe imaging of the female reproductive system, including the pelvis and breast. An effective women's imager must work closely with clinical colleagues of various specialties and maintain a current understanding of diagnostic strategies, clinical implications of imaging findings, and the appropriate use of imaging tests to detect and monitor treatment.

The use of magnetic resonance imaging (MRI) for evaluation of gynecological, obstetric, and breast conditions has increased in recent years. MRI provides excellent tissue contrast resolution in the female pelvis and breast without ionizing radiation. Used together with complementary modalities, such as ultrasound and mammography, MRI has been shown to add important information to help guide patient care. The current text aims to provide the essentials of MRI in Women's Imaging, including indications, technique, and interpretation. For a number of entities, we illustrate the companion imaging studies of computed tomography, ultrasound, or mammography.

Hopefully this text serves to redress the considerable underutilization of MRI in these settings. Used appropriately, MRI is cost-effective and singularly informative. There are other textbooks on the separate topics of pelvic and breast MRI; the goal of this text is to combine and update the essentials of Women's Imaging MRI into a comprehensive and succinct overview.

The present volume is separated into two main sections: female pelvis (chapters 1–7) and breast (chapters 8–12). The first chapter presents current common indications and sample protocols for female pelvis MRI. Chapters 2–5 address pathology and respective imaging findings of the vagina and female urethra, pelvic floor, uterus, and adnexa. Chapters 6 and 7 focus on issues specific to pregnancy. Chapter 8 discusses rationale and technique for MRI of the breast. Chapters 9–12 are dedicated to the imaging features of breast disease and the role of MRI-guided intervention in the care of women with abnormal breast imaging findings.

This text is the collective effort of many individuals. I would like to thank the co-editors and contributors for their hard work. In addition, I am indebted to my radiology colleagues at the University of California San Diego for their help and support, with special thanks to every member of the body imaging and breast imaging divisions.

Michele A. Brown, MD

# Contents

Contributors, vii

Preface, ix

- 1 Pelvis MRI: introduction and technique, 1**  
Michele A. Brown & Richard C. Semelka
- 2 Imaging the vagina and urethra, 8**  
Shannon St. Clair, Randy Fanous, Mohamed El-Azzazi, Richard C. Semelka, & Michele A. Brown
- 3 Pelvic floor imaging, 27**  
Laura E. Rueff & Steven S. Raman
- 4 Imaging the uterus, 49**  
Randy Fanous, Katherine M. Richman, Chayanin Angthong, Mohamed El-Azzazi, & Michele A. Brown
- 5 Imaging the adnexa, 88**  
Michele A. Brown, Mary K. O'Boyle, Chayanin Angthong, Mohamed El-Azzazi, & Richard C. Semelka
- 6 Imaging maternal conditions in pregnancy, 131**  
Lorene E. Romine, Randy Fanous, Michael J. Gabe, Richard C. Semelka, & Michele A. Brown
- 7 Fetal imaging, 180**  
Lorene E. Romine, Ryan C. Rockhill, Michael J. Gabe, Reena Malhotra, Richard C. Semelka, & Michele A. Brown
- 8 Breast MRI: introduction and technique, 239**  
Michael J. Gabe, Jasmina Boban, Dragana Djilas, Vladimir Ivanovic, & Haydee Ojeda-Fournier
- 9 ACR breast MRI lexicon and interpretation, 264**  
Julie Bykowski, Natasa Prvulovic Bunovic, Dragana Djilas, & Haydee Ojeda-Fournier
- 10 Preoperative breast cancer evaluation and advanced breast cancer imaging, 296**  
Jade de Guzman, Dragana Bogdanovic-Stojanovic, Dragana Djilas, & Haydee Ojeda-Fournier
- 11 Postsurgical breast and implant imaging, 322**  
Julie Bykowski, Dag Pavic, Dragana Djilas, & Haydee Ojeda-Fournier
- 12 MR-guided breast interventions, 346**  
Michael J. Gabe, Dragana Djilas, Dag Pavic, & Haydee Ojeda-Fournier

Index, 363



# Chapter 1

## Pelvis MRI: introduction and technique

Michele A. Brown & Richard C. Semelka

### Imaging evaluation of the female pelvis

- Imaging plays an important role in the management of gynecological disease
- Ultrasound is often the initial imaging test
- Poor tissue contrast of CT limits gynecologic applications
- MRI benefits from excellent tissue contrast and lack of ionizing radiation
- Increased experience and availability have led to increased role of MRI
- MRI deemed appropriate by American College of Radiology for gynecological conditions, especially pre-treatment assessment of endometrial and cervical cancer, work-up of suspected adnexal mass, and evaluation of acute pelvic pain in reproductive-aged women in the setting of indeterminate ultrasound [1–4]
- Numerous gynecological and obstetric conditions are depicted by MRI, which may provide initial imaging (e.g., suspected urethral diverticulum) or problem-solving after ultrasound

### Indications for MRI

(Table 1.1)

#### • Benign uterine conditions

- Anomalies
  - MRI considered imaging modality of choice
  - Informs management decisions (e.g., septate versus bicornuate uterus)
- Acquired disease
  - Problem solving for indeterminate ultrasound
  - MRI allows definitive diagnosis for conditions such as urethral diverticulum, leiomyoma, adenomyosis, endometriosis, and dermoid

#### • Uterine malignancy

- Endometrial cancer
  - Preoperative staging: deep myometrial invasion correlated with lymph node invasion [5, 6]
  - MRI shown to aid management for advanced and high grade cancer [7]
- Cervical carcinoma
  - Depth of stromal and parametrial invasion [8, 9]
  - MRI particularly aids management for
    - Tumors larger than 2 cm
    - Endocervical tumors [10]
    - Biopsy-proved adenocarcinoma (cervical versus endometrial origin)
    - Coexistent pelvic mass(es)
    - New diagnosis of cervical cancer during pregnancy
    - Prior radiation therapy [11–15]

#### • Adnexal mass

- Determine origin of mass
- Tissue characterization aids specific diagnosis (e.g., endometrioma, dermoid)
- MRI helps predict likelihood of malignancy to direct proper management and limit surgical intervention for benign disease [16, 17]
- For known ovarian cancer, CT typically used for staging; MRI if CT contraindicated
- MRI may yield definitive diagnosis for adnexal disease that is indeterminate on ultrasound, obviating need for follow-up imaging

#### • Abdominal pain in pregnancy

- Accurate evaluation for appendicitis (and other acute diseases) without ionizing radiation [18, 19]
- Increasing availability of MRI in acute setting

#### • Fetal anomalies

- Problem solving for indeterminate ultrasound
- Usefulness of MRI has increased with ultrafast sequences

**Table 1.1.** Indications for MRI of the female pelvis

Indication	Protocol	Notes
<b>Pelvic pain</b>	General	FS T1WI for endometriosis
<b>Urethral diverticulum</b>	Urethra	Contrast if known/visualized mass
<b>Vaginal mass</b>	Urethra	Contrast if known/visualized mass
<b>Pelvic floor symptoms</b>	Pelvic floor	Sagittal images with Valsalva
<b>Uterine anomaly</b>	Uterine anomaly	True coronal to uterine fundus
<b>Adenomyosis</b>	General	Bright myometrial foci on T2WI
<b>Fibroids</b>	General	Add contrast if pre-embolization
<b>Fibroid versus adnexal mass</b>	General	Vessels extending from uterus to mass suggest uterine origin
<b>Endometrial cancer</b>	Uterine malignancy	High resolution T2WI and T1WI + contrast oblique to endometrium for tumor invasion
<b>Cervical cancer</b>	Uterine malignancy	High resolution T2WI oblique to cervix for parametrial invasion
<b>Adnexal mass characterization</b>	General	FS T1WI for dermoid, endometrioma
<b>Abdominal pain in pregnancy</b>	Maternal abdominal pain	SS-ETSE (+ FS), and steady-state GE for appendix, monitor if possible
<b>Fetal anomaly</b>	Fetal	SS-ETSE oriented to region of interest, monitor if possible

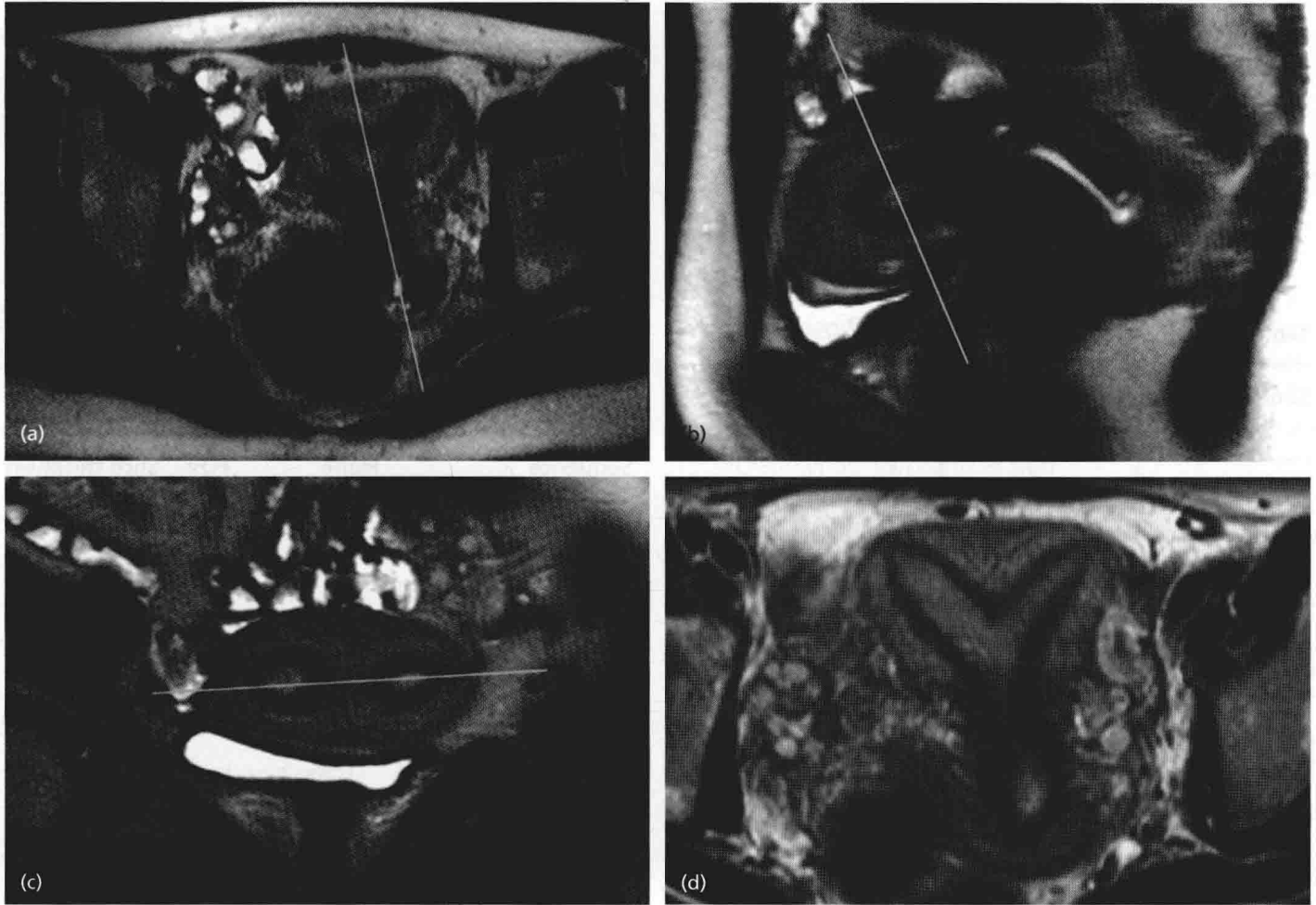
FS = fat saturated; T1WI = T1-weighted images; T2WI = T2-weighted images; SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo

## Patient preparation for MRI

- Empty bladder
- Fasting 4 hours
- Optional
  - Antispasmodic (e.g., glucagon 1 mg)
  - Intra-vaginal gel [20]
- Supine position, or decubitus in late pregnancy
- Phased-array coil positioned over pelvis
- To reduce artifact, may utilize
  - Saturation band over anterior abdominal wall for non-fat-saturated sagittal
  - Supplemental anteroposterior frequency-encoding direction for axial images
- Intrauterine contraceptive devices are safely imaged [21]

## Sequence protocols

- Many protocol options
- Appropriate choice depends on
  - Specific clinical question
  - Available equipment and expertise
- For known or suspected uterine disease/anomalies, T2-weighted sequences are obtained in an oblique plane oriented to uterus (Figure 1.1)
- Individual sequence parameters may vary based on manufacturer, etc.
- Sequences may include
  - Single-shot (SS) echo-train spin echo (ETSE)
    - For example, HASTE or SSFSE
    - Sensitive to fluid, resistant to motion and susceptibility
    - Large field of view
      - Localization, evaluation of coil position
      - Coronal: evaluation of renal anomalies/obstruction
      - Axial: prescribe true sagittal view of uterus
  - T2-weighted
    - Breathhold may be sufficient for benign disease
    - Non-breathhold (high-resolution) for uterine malignancy
    - With or without fat saturation
    - May be done as 3D ETSE
    - Best sequence for uterine zonal anatomy
  - T1-weighted



**Figure 1.1.** Imaging planes oriented to the uterus. Multiple T2-weighted images in a patient with septate uterus. Large field-of-view single-shot sequence **(a)** is obtained first and is used to plan an oblique sagittal T2-weighted sequence **(b)** obtained parallel to the endometrium (line, **a**). The oblique sagittal is used to plan an oblique axial **(c)** obtained perpendicular to the endometrium (line, **b**). The oblique axial may then be used to plan a true coronal of the uterus **(d)** obtained parallel to the endometrium (line, **c**). In the absence of 3D T2-weighted imaging, this process assures appropriate imaging planes regardless of angle/tilt of the uterus.

- Breathhold in- and out-of-phase dual echo
  - Differentiates fat- and blood-containing lesions
  - Sensitive to small foci of fat within adnexal mass
- Non-breathhold (high-resolution) for uterine cancer
- Chemically selective fat saturation for endometriosis
- T2/T1-weighted steady-state free precession gradient echo (GE)
  - For example, TruFISP or FIESTA
  - Rapid, resistant to motion
  - Differentiates vessels from bowel (e.g., appendix)
  - Useful for fetal and maternal imaging
- T1-weighted 3D GE pre- and post-contrast
  - Fat-suppressed GE, repeated for dynamic imaging
  - Provides enhancement information
  - May use MRA parameters (e.g., vascular malformation)
- Diffusion-weighted imaging (DWI) (optional)
  - B values of 0 and at least one other value up to 1000
  - Apparent diffusion coefficient (ADC) map created
  - DWI sequence and ADC map interpreted together
  - Aids detection of tumor, inflammation

- Additional functional techniques may have increasing role [7]
- Oblique planes oriented to the endometrium or cervix important for cancer [22]
- Protocol tailored to clinical question (Table 1.2, Table 1.3, Table 1.4, Table 1.5, Table 1.6, Table 1.7, Table 1.8)

**Table 1.2.** General female pelvis

Sequence	Plane	FOV (cm)	Slice thickness (mm)
SS-ETSE	Coronal	32	8
SS-ETSE	Axial	32	8
T2 ETSE	Sagittal	24	5
T2 ETSE	Axial	24	5
T1 GE in/out- of-phase	Axial	24	5
T1 GE FS	Axial	24	5
DWI (optional)	Axial	28	6
T1 3D GE FS (pre)	Axial or sagittal	24	3
<i>Contrast</i>			
T1 3D GE FS (post × 3)	Axial or sagittal	24	3
T1 GE FS (delayed)	Axial	24	5

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo; FS = fat saturated; DWI = diffusion-weighted imaging

**Table 1.3.** Urethra

Sequence	Plane	FOV (cm)	Slice thickness (mm)
SS-ETSE	Coronal	32	8
T2 ETSE	Coronal	16	4
T2 ETSE	Axial	16	4
T1 GE	Axial	16	4
T1 3D GE FS (pre)	Axial	24	3
<i>Contrast (if known or visualized lesion)</i>			
T1 GE FS (delayed)	Axial	24	3

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo; FS = fat saturated

**Table 1.4.** Pelvic floor

Sequence	Plane	FOV (cm)	Slice thickness (mm)
SS-ETSE	Coronal	32	8
SS-ETSE	Axial	32	8
SS-ETSE	Sagittal	32	5
SS-ETSE (Valsalva, repeat × 3)	Sagittal	32	5 (midline slice)

SS = single shot; ETSE = echo-train spin-echo

**Table 1.5.** Uterine anomaly

Sequence	Plane	FOV (cm)	Slice thickness (mm)
SS-ETSE	Coronal	32	8
SS-ETSE	Axial	32	8
T2 ETSE	Sagittal (to uterus)	24	5
T2 ETSE	Axial (to uterus)	24	5
T2 ETSE	Coronal (to uterus)	24	5
T1 GE in/out-of-phase	Coronal (to uterus)	24	5
T1 GE FS	Axial	24	5

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo; FS = fat saturated

**Table 1.6.** Uterine malignancy

Sequence	Plane	FOV (cm)	Slice thickness (mm)
SS-ETSE	Coronal	32	8
SS-ETSE	Axial	32	8
T2 ETSE	Sagittal	24	5
T2 ETSE	Axial	24	5
T1 GE in/out-of-phase	Axial	24	5
T1 GE FS	Axial	24	5
DWI (optional)	Axial	28	6
T1 3D GE FS (pre)	Axial or sagittal	24	3
<i>Contrast</i>			
T1 3D GE FS (post × 3)	Axial or sagittal	24	3
T1 GE FS (delayed)	Axial	24	5

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo; FS = fat saturated

**Table 1.7.** Maternal abdominal pain

Sequence	Plane	FOV (cm)	Slice thickness (mm)
<b>SS-ETSE</b>	Coronal	32–40	8
<b>SS-ETSE</b>	Axial	32	5
<b>SS-ETSE FS</b>	Axial	32	5
<b>Steady-state GE</b>	Coronal	32	5
<b>Steady-state GE</b>	Axial	32	5
<b>T1 GE in/out-of-phase</b>	Axial	32	5

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo;  
FS = fat saturated

**Table 1.8.** Fetal

Sequence	Plane	FOV (cm)	Slice thickness (mm)
<b>SS-ETSE</b>	Coronal	40	8
<b>SS-ETSE</b>	Axial	40	8
<b>SS-ETSE</b>	Sagittal	40	8
<b>SS-ETSE (repeat as needed)</b>	Directed	24–32	4–6
<b>Steady-state GE (optional)</b>	Directed	24–32	4–6
<b>T1 GE in/out-of-phase (optional)</b>	Directed	24–32	4–6

SS = single shot; ETSE = echo-train spin-echo; GE = gradient echo;  
FS = fat saturated

## Image optimization at 3T

- Potential advantages
  - Increase in signal-to-noise ratio (SNR), or
  - Similar SNR at a faster speed
- Challenges
  - Signal shading magnified by dielectric effects
  - Increased specific absorption rates (SARs)
  - Changes in optimal TR and TE
  - Increased signal inhomogeneities
    - Greater shimming challenge for extrinsic magnetic field
    - Intrinsic field distortion due to increased susceptibility/chemical shift
- Solutions [23–28]
  - Dielectric effect: dielectric pad (= radiofrequency cushion) placed between patient and surface coil
  - Susceptibility: use shorter TE/higher receiver bandwidth, higher spatial resolution
  - 3D GE and ETSE sequences may benefit from higher field strength
  - Consider individual patient
    - Pregnant patients less suitable for 3T due to standing wave effects from amniotic fluid and safety concerns [26]
    - Non-pregnant patients may be imaged safely and effectively at 3T using optimized parameters [28]

## Image interpretation

- Large volume data acquisition
- May be useful to employ a systematic checklist (Table 1.9)
- Several gynecological conditions have MRI features that allow definitive diagnosis



**Table 1.9.** Diagnostic checklist for female pelvis MRI

Structure	MRI features evaluated
Gynecological	
<b>Uterine corpus</b>	Size and position Presence of myometrial mass Endometrium thickness Junctional zone thickness
<b>Cervix</b>	Presence of cystic mass Presence of solid tumor Size of lesion Parametrial involvement
<b>Vagina</b>	Presence of cystic mass Presence of wall thickening/solid tumor
<b>Adnexa</b>	Ovarian size Presence of ovarian mass Cystic or solid Fat containing Blood containing Enhancement features Unilateral or bilateral Paraovarian cystic or solid mass
Non-gynecological	
<b>Bladder</b>	Presence of solid mass Presence of cystocele
<b>Urethra</b>	Presence of diverticulum Size and configuration Solid/enhancing components Presence of hypermobility
<b>Bowel</b>	Caliber Presence of rectocele
<b>Musculoskeletal</b>	Bone marrow signal Degenerative changes Traumatic injury
<b>Lymphatic</b>	Enlarged lymph nodes

## References

1. Lee, J.H., Dubinsky, T., Andreotti, R.F., et al. ACR Appropriateness Criteria(R) pretreatment evaluation and follow-up of endometrial cancer of the uterus. *Ultrasound Quarterly* 2011; 27(2):139–45.
2. Siegel, C.L., Andreotti, R.F., Cardenes, H.R., et al. ACR Appropriateness Criteria(R) pretreatment planning of invasive cancer of the cervix. *Journal of the American College of Radiology* 2012; 9(6):395–402.
3. Harris, R.D., Javitt, M.C., Glanc, P., et al. ACR Appropriateness Criteria(R) clinically suspected adnexal mass. *Ultrasound Quarterly* 2013; 29(1):79–86.
4. Andreotti, R.F., Lee, S.I., Dejesus Allison, S.O., et al. ACR Appropriateness Criteria(R) acute pelvic pain in the reproductive age group. *Ultrasound Quarterly* 2011; 27(3):205–10.
5. Kinkel, K., Kaji, Y., Yu, K.K., et al. Radiologic staging in patients with endometrial cancer: a meta-analysis. *Radiology* 1999; 212(3):711–18.
6. Wakefield, J.C., Downey, K., Kyriazi, S., deSouza, N.M. New MR techniques in gynecologic cancer. *AJR. American Journal of Roentgenology* 2013; 200(2):249–60.
7. Frei, K.A., Kinkel, K., Bonél, H.M., et al. Prediction of deep myometrial invasion in patients with endometrial cancer: clinical utility of contrast-enhanced MR imaging – a meta-analysis and Bayesian analysis. *Radiology* 2000; 216(2):444–9.
8. Sironi, S., De Cobelli, F., Scarfone, G., et al. Carcinoma of the cervix: value of plain and gadolinium-enhanced MR imaging in assessing degree of invasiveness. *Radiology* 1993; 188(3):780–97.
9. Subak, L.L., Hricak, H., Powell, C.B., Azizi, L., Stern, J.L. Cervical carcinoma: computed tomography and magnetic resonance imaging for preoperative staging. *Obstetrics and Gynecology* 1995; 86(1):43–50.
10. Hricak, H., Powell, C.B., Yu, K.K., et al. Invasive cervical carcinoma: role of MR imaging in pretreatment work-up – cost minimization and diagnostic efficacy analysis. *Radiology* 1996; 198(2):403–9.
11. Flueckiger, F., Ebner, F., Poschauko, H., et al. Cervical cancer: serial MR imaging before and after primary radiation therapy – a 2-year follow-up study. *Radiology* 1992; 184(1):89–93.
12. Hricak, H., Swift, P.S., Campos, Z., et al. Irradiation of the cervix uteri: value of unenhanced and contrast-enhanced MR imaging. *Radiology* 1993; 189(2):381–8.
13. Weber, T.M., Sostman, H.D., Spritzer, C.E., et al. Cervical carcinoma: determination of recurrent tumor extent versus radiation changes with MR imaging. *Radiology* 1995; 194(1):135–9.
14. Yamashita, Y., Harada, M., Torashima, M., et al. Dynamic MR imaging of recurrent postoperative cervical cancer. *Journal of Magnetic Resonance Imaging* 1996; 6(1):167–71.
15. Hertel, H., Köhler, C., Grund, D., et al. Radical vaginal trachelectomy (RVT) combined with laparoscopic pelvic lymphadenectomy: prospective multicenter study of 100 patients with early cervical cancer. *Gynecologic Oncology* 2006; 103(2): 506–11.

16. Hricak, H., Chen, M., Coakley, F.V., et al. Complex adnexal masses: detection and characterization with MR imaging – multivariate analysis. *Radiology* 2000; 214(1):39–46.
17. Sohaib, S.A., Sahdev, A., Van Trappen, P., Jacobs, I.J., Reznick, R.H. Characterization of adnexal mass lesions on MR imaging. *AJR. American Journal of Roentgenology* 2003; 180(5):1297–304.
18. Birchard, K.R., Brown, M.A., Hyslop, W.B., Firat, Z., Semelka, R.C. MRI of acute abdominal and pelvic pain in pregnant patients. *AJR. American Journal of Roentgenology* 2005; 184(2):452–8.
19. Oto, A., Ernst, R.D., Shah, R., et al. Right-lower-quadrant pain and suspected appendicitis in pregnant women: evaluation with MR imaging – initial experience. *Radiology* 2005; 234(2):445–51.
20. Brown, M.A., Mattrey, R.F., Stamato, S., Sirlin, C.B. MRI of the female pelvis using vaginal gel. *AJR. American Journal of Roentgenology* 2005; 185(5):1221–7.
21. Pasquale, S.A., Russer, T.J., Foldes, R., Mezrich, R.S. Lack of interaction between magnetic resonance imaging and the copper-T380A IUD. *Contraception* 1997; 55(3): 169–73.
22. Shiraiwa, M., Joja, I., Asakawa, T., et al. Cervical carcinoma: efficacy of thin-section oblique axial T2-weighted images for evaluating parametrial invasion. *Abdominal Imaging* 1999; 24(5): 514–19.
23. Kataoka, M., Kido, A., Koyama, T., et al. MRI of the female pelvis at 3T compared to 1.5T: evaluation on high-resolution T2-weighted and HASTE images. *Journal of Magnetic Resonance Imaging* 2007; 25(3): 527–34.
24. Martin, D.R., Friel, H.T., Danrad, R., De Becker, J., Hussain, S.M. Approach to abdominal imaging at 1.5 Tesla and optimization at 3 Tesla. *Magnetic Resonance Imaging Clinics of North America* 2005; 13(2):241–54.
25. Hussain, S.M., van den Bos, I.C., Oliveto, J.M., Martin, D.R. MR imaging of the female pelvis at 3T. *Magnetic Resonance Imaging Clinics of North America* 2006; 14(4):537–44.
26. Merkle, E.M., Dale, B.M. Abdominal MRI at 3.0 T: the basics revisited. *AJR. American Journal of Roentgenology* 2006; 186(6):1524–32.
27. Cornfeld, D., Weinreb, J. Simple changes to 1.5-T MRI abdomen and pelvis protocols to optimize results at 3T. *AJR. American Journal of Roentgenology* 2008; 190(2): W140–50.
28. Morakkabati-Spitz, N., Schild, H.H., Kuhl, C.K., et al. Female pelvis: MR imaging at 3.0 T with sensitivity encoding and flip-angle sweep technique. *Radiology* 2006; 241(2): 538–45.

## Chapter 2

# Imaging the vagina and urethra

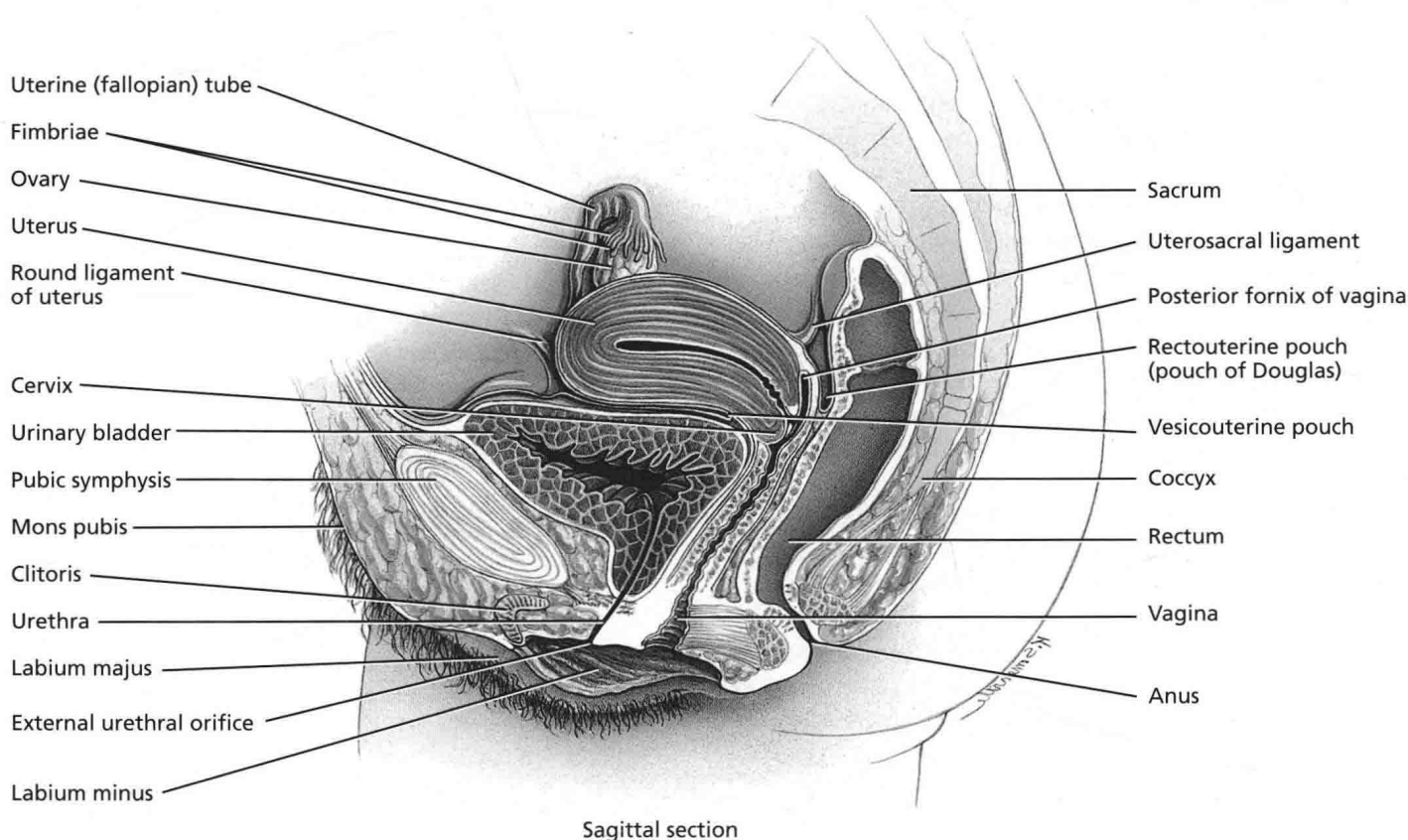
Shannon St. Clair, Randy Fanous, Mohamed El-Azzazi, Richard C. Semelka, & Michele A. Brown

### Vagina

#### Normal anatomy

##### Key facts

- Fibromuscular tube between bladder and rectum, 7–9cm long (Figure 2.1) Embryological origin [1]
  - Upper one-third = Müllerian duct
  - Lower two-thirds = urogenital sinus
- Layers [2]
  - Inner = mucosa
  - Middle = submucosa and muscularis
  - Outer = adventitia, containing vaginal venous plexus
- Fornices: anterior, posterior, lateral
  - Portion of vagina that surrounds the cervix
  - Best visualized on sagittal and transverse images



**Figure 2.1.** Normal female pelvic anatomy in the sagittal plane. (Source: Tortora & Derrickson (Eds), Principles of Anatomy and Physiology, 13th edn. Hoboken, NJ: Wiley, 2012.)

- For descriptive purposes, vagina may be divided into thirds
  - Upper third = level of the lateral fornices
  - Middle third = level of the bladder base
  - Lower third = level of the urethra

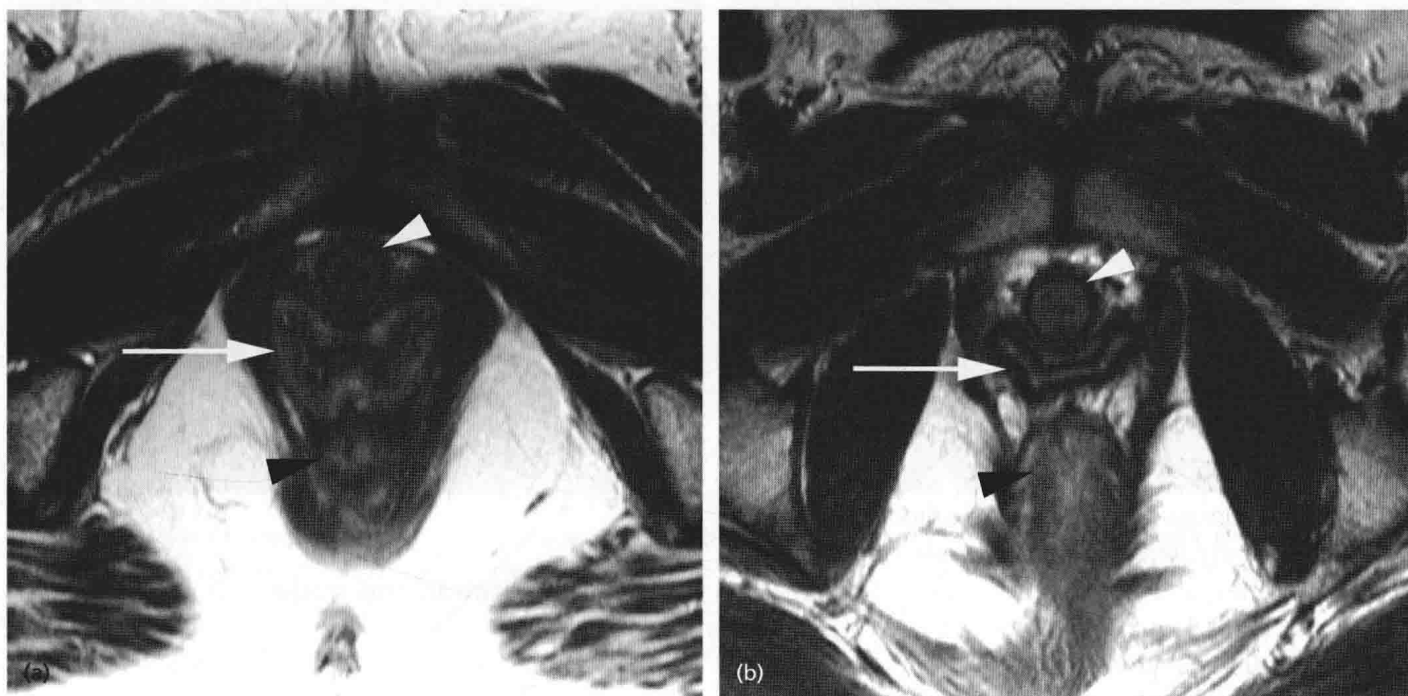
### **MRI features**

- T1WI – low signal intensity
- T2WI
  - Stratified: outer high signal intensity, middle low signal intensity, inner high signal intensity
  - Thickness correlates with estrogen level; most prominent during late proliferative and early secretory phases of menstrual cycle (Figure 2.2)
  - Loss of normal stratification
    - Pregnant = outer and middle intermediate to high signal intensity, inner high signal intensity
    - Premenarchal/ postmenopausal = outer and middle low signal intensity, markedly thin inner high signal intensity
- T1WI + contrast – avid early enhancement of outer and middle layers only

### **Vaginal agenesis/atresia**

#### **Key facts**

- Rare Müllerian duct anomaly ranging from complete to partial agenesis
  - Incidence of all Müllerian duct anomalies in women is 1–15%
  - 1 in 4000–5000 women have vaginal agenesis [3, 4]
- Typically normal ovaries and external genitalia, however associated abnormalities of the uterus, cervix, upper urinary tract, and skeleton may occur
- Presentation depends on presence of functioning endometrium
  - If no functioning endometrium = primary amenorrhea
  - If functioning endometrium present = pain and mass effect at the age of menarche secondary to hematometra (Figure 2.3)
- Untreated patients with functional endometrium may develop endometriosis
- Surgical management depends on presence of functioning endometrium and cervix
  - Complete agenesis + small rudimentary uterine bulb with no functioning endometrium = vaginoplasty



**Figure 2.2.** Normal vagina in two patients. Axial T2-weighted image in two patients (a, b) show variable thickness in the vagina depending on estrogen levels. Note vagina (arrow), urethra (white arrowhead), and rectum (black arrowhead).