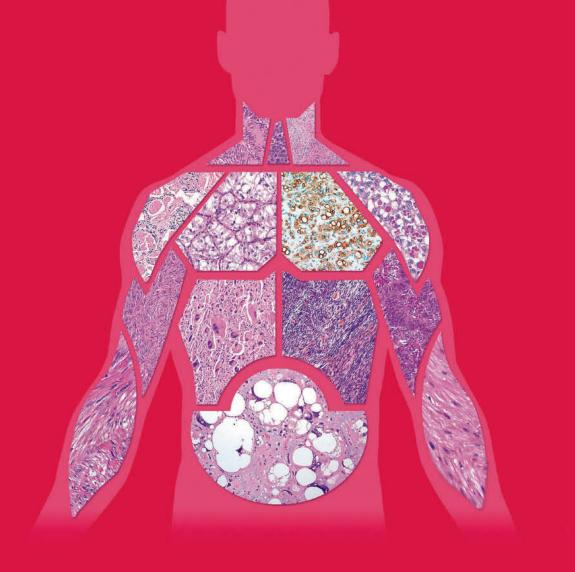


# Soft Tissue Tumors

Lindberg

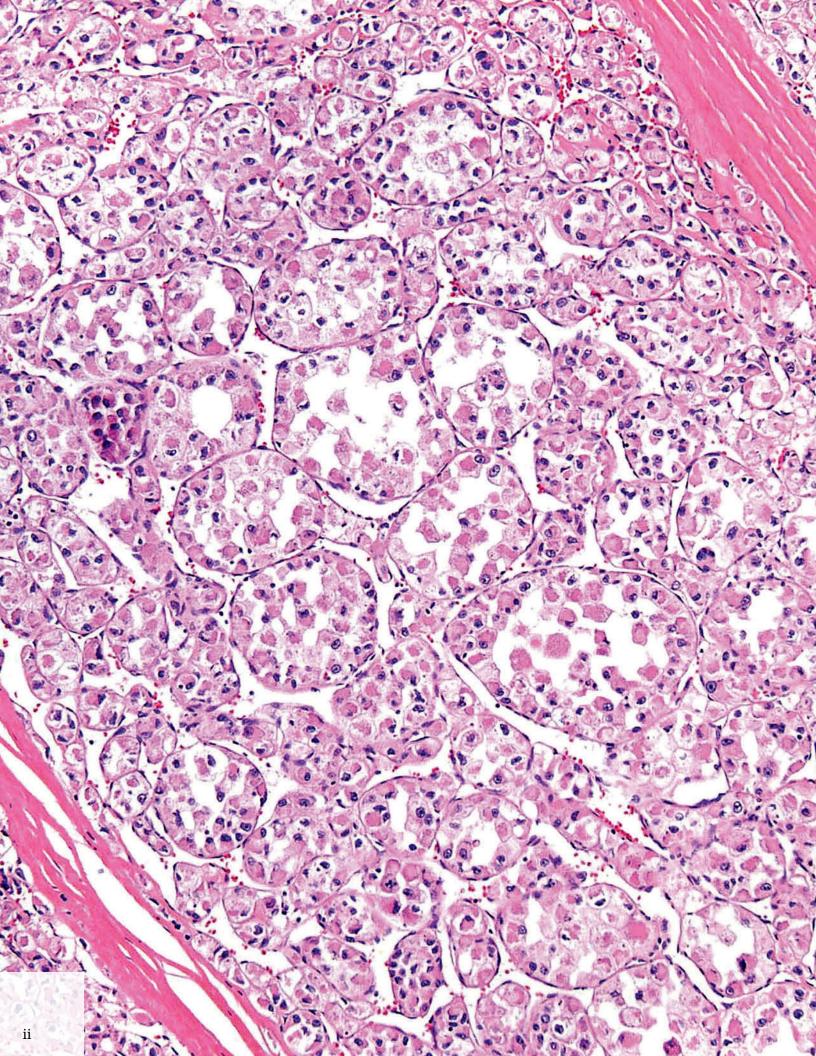




# Soft Tissue Tumors

Lindberg





# Soft Tissue Tumors

Third Edition

# Matthew R. Lindberg, MD

Associate Professor of Pathology University of Arkansas for Medical Sciences Little Rock, Arkansas

#### DIAGNOSTIC PATHOLOGY: SOFT TISSUE TUMORS, THIRD EDITION

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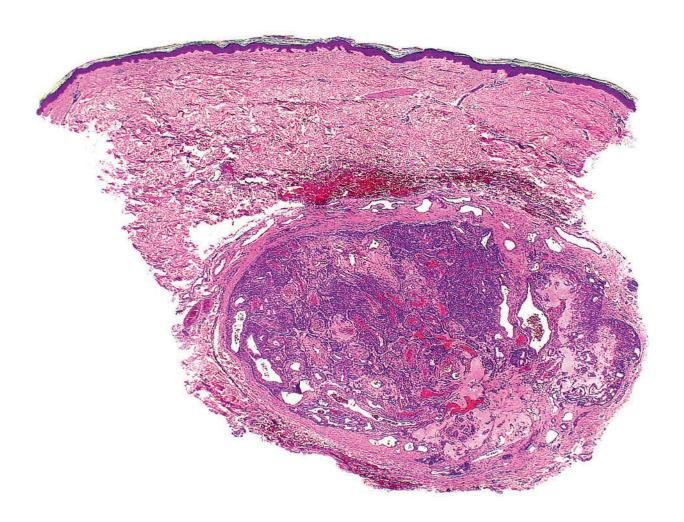
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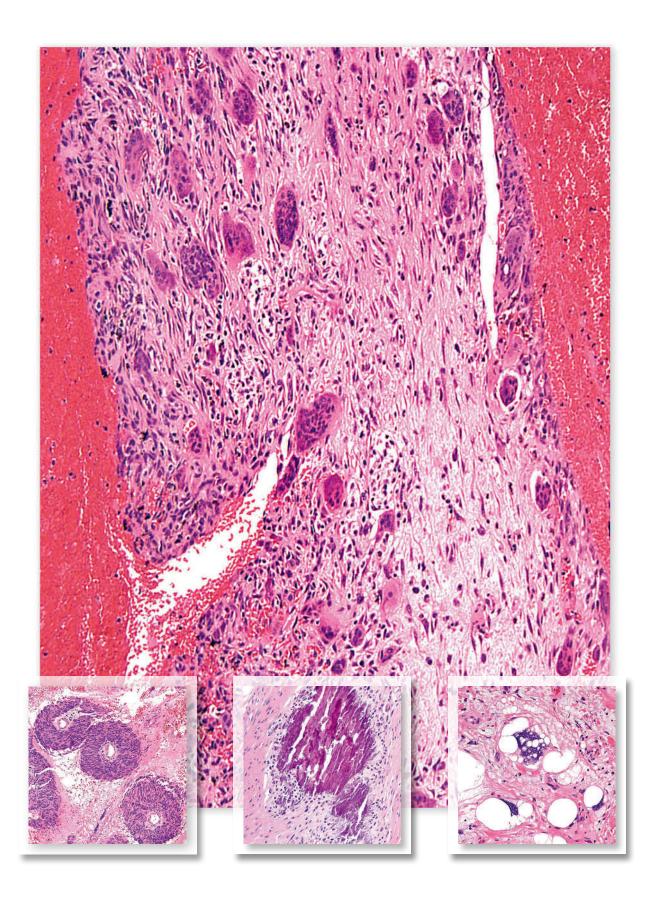


# Dedication

To my wonderful wife, Rani, for her love, support, and infinite patience during the course of putting together this book. Also, to the excellent team of editors and coauthors with whom I have been very privileged to work. Many, many thanks for your hard work and high standards.

MRL





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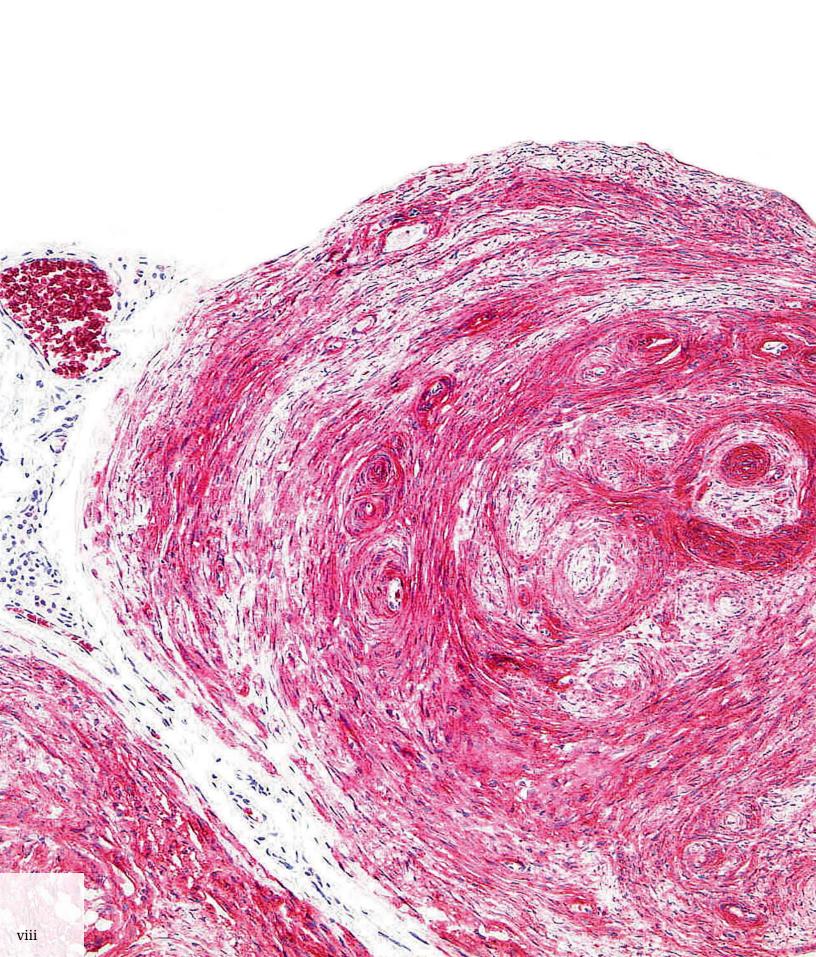
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# Preface

The field of soft tissue pathology continues to grow and change, with new diagnostic entities, immunohistochemical antibodies, and molecular tests introduced seemingly every month. The 3rd edition of *Diagnostic Pathology: Soft Tissue Tumors* strives to incorporate this new knowledge in the form of new chapters, updated text, and additional high-quality histologic images.

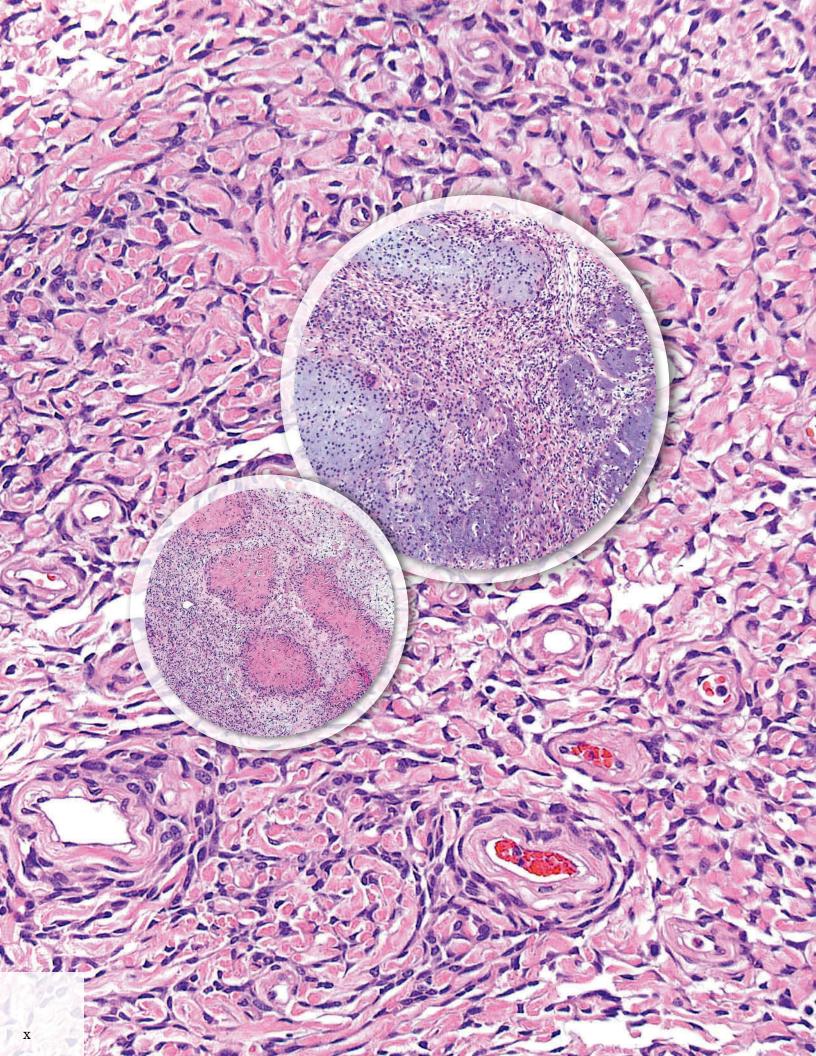
Since the publication of the 2nd edition, several new entities have been described in the scientific literature, including atypical spindle cell lipomatous tumor, superficial CD34-positive fibroblastic tumor, and *BCOR*-rearranged sarcoma. Chapters dedicated to each of these entities (as well as others) have been added in this new edition. Existing chapters devoted entirely to immunohistochemistry and molecular testing have also been updated to reflect recent discoveries in this span, and many individual chapters have seen a variety of text and gallery improvements. In particular, several galleries have been modified by swapping in new images that better reflect the extensive morphologic spectrum of soft tissue pathology.

As before, the 3 innovative "Approach to Diagnosis" chapters are still included in this new edition. Using a combination of clinical information, overall histologic pattern, and specific histologic findings, these chapters can aid the struggling pathologist in developing a thoughtful differential diagnosis for even some of the more challenging or unusual soft tissue cases. I hope you find these unique additions helpful in your own practice, both now and for many years to come.

Lastly, as always, owners of *Diagnostic Pathology: Soft Tissue Tumors*, 3rd edition, receive online access to all information and images contained in this text, plus much more. Enjoy!

#### Matthew R. Lindberg, MD

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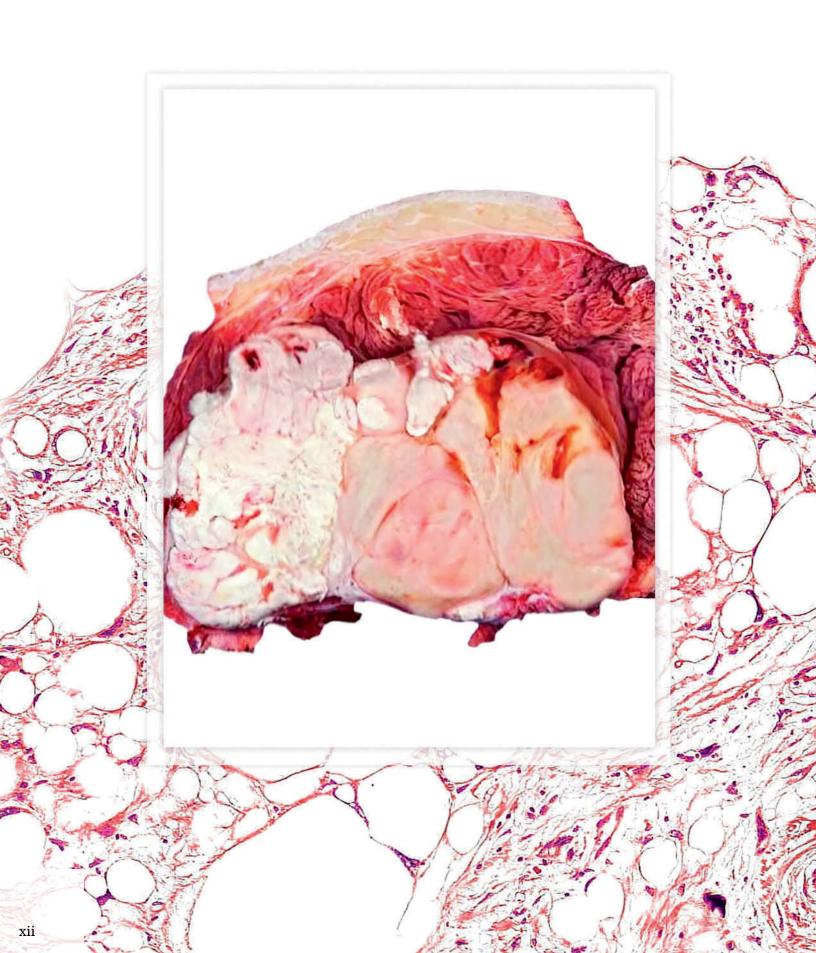
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# Sections

**SECTION 1: Soft Tissue Introduction** 

SECTION 2: Diagnostic Approach to Soft Tissue Tumors

**SECTION 3: Tumors of Adipose Tissue** 

SECTION 4: Fibroblastic/Myofibroblastic Lesions

SECTION 5: Pediatric Fibroblastic/Myofibroblastic Tumors

SECTION 6: Fibrohistiocytic, Histiocytic, and Dendritic Cell Tumors

**SECTION 7: Smooth Muscle Tumors** 

SECTION 8: Pericytic (Perivascular) Tumors

**SECTION 9: Tumors of Skeletal Muscle** 

SECTION 10: Vascular Tumors (Including Lymphatics)

**SECTION 11: Chondroosseous Tumors** 

**SECTION 12: Peripheral Nerve Sheath Tumors** 

**SECTION 13: Genital Stromal Tumors** 

SECTION 14: Tumors of Mesothelial Cells

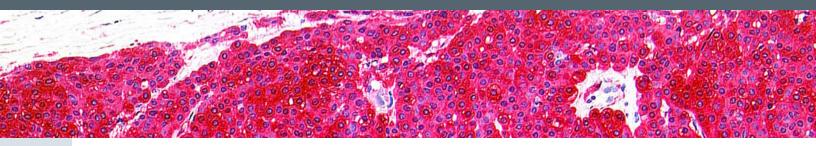
SECTION 15: Hematopoietic Tumors in Soft Tissue

SECTION 16: Tumors of Uncertain Differentiation

SECTION 17: Undifferentiated/Unclassified Sarcomas

SECTION 18: Mesenchymal Tumors of Gastrointestinal Tract

**SECTION 19: Other Entities** 



#### **SECTION 1: SOFT TISSUE INTRODUCTION**

#### INTRODUCTION AND OVERVIEW

- 4 Gross Examination Matthew R. Lindberg, MD
- 6 Grading and Staging
  Matthew R. Lindberg, MD

#### **ANCILLARY TECHNIQUES**

- 12 Soft Tissue Immunohistochemistry Matthew R. Lindberg, MD
- **18** Molecular Features of Soft Tissue Tumors Matthew R. Lindberg, MD



# SECTION 2: DIAGNOSTIC APPROACH TO SOFT TISSUE TUMORS

#### **OVERVIEW**

22 Biopsy and Resection of Soft Tissue Tumors
Matthew R. Lindberg, MD

#### **CLINICAL APPROACH**

26 Age- and Location-Based Approach to Diagnosis
Matthew R. Lindberg, MD

#### HISTOLOGIC APPROACH

- 28 Pattern-Based Approach to Diagnosis Matthew R. Lindberg, MD
- **36** Feature-Based Approach to Diagnosis Matthew R. Lindberg, MD



#### **SECTION 3: TUMORS OF ADIPOSE TISSUE**

#### **BENIGN**

- 46 Lipoma
  - . Matthew R. Lindberg, MD
- **52 Lipomatosis of Nerve** *Jerad M. Gardner, MD*
- 54 Synovial Lipomatosis Matthew R. Lindberg, MD
- **56** Angiolipoma Jerad M. Gardner, MD
- 60 Spindle Cell/Pleomorphic Lipoma Matthew R. Lindberg, MD
- 66 Chondroid Lipoma Matthew R. Lindberg, MD
- **70 Myolipoma** *Jerad M. Gardner, MD*

- 74 Hibernoma
  - Matthew R. Lindberg, MD
- 78 Myelolipoma
- Matthew R. Lindberg, MD

  80 Lipoblastoma
  - Matthew R. Lindberg, MD
- 84 Atypical Spindle Cell Lipomatous Tumor Matthew R. Lindberg, MD

#### INTERMEDIATE, LOCALLY AGGRESSIVE

88 Atypical Lipomatous Tumor/Well-Differentiated Liposarcoma

Matthew R. Lindberg, MD

#### **MALIGNANT**

- **94 Dedifferentiated Liposarcoma** *Matthew R. Lindberg, MD*
- 100 Myxoid Liposarcoma

  Matthew R. Lindberg, MD
- 106 Pleomorphic Liposarcoma Matthew R. Lindberg, MD

#### SECTION 4: FIBROBLASTIC/MYOFIBROBLASTIC LESIONS

#### **BENIGN**

- **114 Nodular Fasciitis** *Matthew R. Lindberg, MD*
- 120 Proliferative Fasciitis/Myositis
  Matthew R. Lindberg, MD
- 124 Ischemic Fasciitis

  Matthew R. Lindberg, MD
- Matthew R. Lindberg, MD

  126 Myositis Ossificans

  Matthew R. Lindberg, MD
- 130 Fibroosseous Pseudotumor of Digit
  David Lucas, MD and Elizabeth A. Montaomery, MD
- **134 Fibroma of Tendon Sheath** *David Lucas. MD*
- 140 Desmoplastic Fibroblastoma
  David Cassarino, MD, PhD and Khin Thway, MD, FRCPath
- Matthew R. Lindberg, MD

  148 Mammary-Type Myofibroblastoma
  Matthew R. Lindberg, MD
- 152 Intranodal Palisaded Myofibroblastoma Matthew R. Lindberg, MD

**156** Pleomorphic Fibroma David Cassarino, MD, PhD

**158 Dermatomyofibroma** *David Cassarino, MD, PhD* 

**Storiform Collagenoma** *David Cassarino, MD, PhD* 

**162 Keloid** *David Cassarino, MD, PhD* 

**164** Nuchal-Type Fibroma *Matthew R. Lindberg, MD* 

#### **INTERMEDIATE (LOCALLY AGGRESSIVE)**

**168 Desmoid-Type Fibromatosis** *Matthew R. Lindberg, MD* 

#### **INTERMEDIATE (RARELY METASTASIZING)**

**174 Dermatofibrosarcoma Protuberans** *Matthew R. Lindberg, MD* 

**184 Solitary Fibrous Tumor** *Matthew R. Lindberg, MD* 

**192 Low-Grade Myofibroblastic Sarcoma** *Matthew R. Lindberg, MD* 

**196** Inflammatory Myofibroblastic Tumor Matthew R. Lindberg, MD

**202 Myxoinflammatory Fibroblastic Sarcoma** *Matthew R. Lindberg, MD* 

210 Superficial CD34(+) Fibroblastic Tumor Matthew R. Lindberg, MD

#### **MALIGNANT**

**214** Adult-Type Fibrosarcoma *Jerad M. Gardner, MD* 

**216 Myxofibrosarcoma** *Matthew R. Lindberg, MD* 

**222 Low-Grade Fibromyxoid Sarcoma** *Matthew R. Lindberg, MD* 

**232 Sclerosing Epithelioid Fibrosarcoma** *Matthew R. Lindberg, MD* 

SECTION 5: PEDIATRIC
FIBROBLASTIC/MYOFIBROBLASTIC
TUMORS

#### **BENIGN**

**240 Fibrous Hamartoma of Infancy** *Matthew R. Lindberg, MD* 

**244** Calcifying Aponeurotic Fibroma *Matthew R. Lindberg, MD* 

248 Calcifying Fibrous Tumor

Kandi Stallings-Archer, MD and Elizabeth A. Montgomery, MD

**250** Inclusion Body Fibromatosis *Matthew R. Lindberg, MD* 

**Hyaline Fibromatosis Syndrome**Kandi Stallings-Archer, MD, Elizabeth A. Montgomery,

MD, and Cyril Fisher, MD, DSc, FRCPath

254 Fibromatosis Colli

Kandi Stallings-Archer, MD and Cyril Fisher, MD, DSc, FRCPath

**256 Gardner Fibroma** *Jerad M. Gardner, MD* 

#### **INTERMEDIATE (LOCALLY AGGRESSIVE)**

258 Lipofibromatosis

Kandi Stallings-Archer, MD and Elizabeth A. Montgomery, MD

**260** Giant Cell Fibroblastoma Matthew R. Lindberg, MD

#### **INTERMEDIATE (RARELY METASTASIZING)**

**264 Infantile Fibrosarcoma** *Matthew R. Lindberg, MD* 

SECTION 6: FIBROHISTIOCYTIC, HISTIOCYTIC, AND DENDRITIC CELL TUMORS

#### **BENIGN**

**270 Dermatofibroma and Fibrous Histiocytoma** *David Cassarino, MD, PhD* 

**276 Deep Benign Fibrous Histiocytoma** *Matthew R. Lindberg, MD* 

278 Localized-Type Tenosynovial Giant Cell Tumor David Lucas, MD

284 Diffuse-Type Tenosynovial Giant Cell Tumor David Lucas, MD

290 Cellular Neurothekeoma Jerad M. Gardner, MD and Cyril Fisher, MD, DSc, FRCPath

**294 Xanthomas** *Matthew R. Lindberg, MD* 

298 Solitary (Juvenile) Xanthogranuloma
David Cassarino, MD, PhD and Elizabeth A. Montgomery,
MD

300 Reticulohistiocytoma David Cassarino, MD, PhD

**304** Deep Granuloma Annulare *Jerad M. Gardner, MD* 

**306** Rheumatoid Nodule *Jerad M. Gardner. MD* 

308 Langerhans Cell Histiocytosis Matthew R. Lindberg, MD

310 Extranodal Rosai-Dorfman Disease Matthew R. Lindberg, MD

314 Crystal-Storing Histiocytosis
Matthew R. Lindberg, MD and Elizabeth A. Montgomery,
MD

#### **INTERMEDIATE (RARELY METASTASIZING)**

316 Plexiform Fibrohistiocytic Tumor Matthew R. Lindberg, MD

**320** Giant Cell Tumor of Soft Tissue David Lucas, MD

#### **MALIGNANT**

- **324 Histiocytic Sarcoma** *Matthew R. Lindberg, MD*
- 326 Follicular Dendritic Cell Sarcoma Matthew R. Lindberg, MD
- 328 Interdigitating Dendritic Cell Sarcoma
  Matthew R. Lindberg, MD, L. Jeffrey Medeiros, MD, and
  Cvril Fisher, MD, DSc, FRCPath

#### SE

#### **SECTION 7: SMOOTH MUSCLE TUMORS**

#### **BENIGN**

- 332 Smooth Muscle Hamartoma Jerad M. Gardner, MD
- **334** Superficial Leiomyoma

  Jerad M. Gardner, MD and Jonathan B. McHugh, MD
- 338 Deep Leiomyoma
  Matthew R. Lindberg, MD

#### **INTERMEDIATE**

342 Epstein-Barr Virus-Associated Smooth Muscle Tumor David Lucas, MD

#### **MALIGNANT**

**344 Leiomyosarcoma** *Matthew R. Lindberg, MD* 



SECTION 8: PERICYTIC (PERIVASCULAR)
TUMORS

#### **BENIGN**

- 352 Glomus Tumors (and Variants)
  Thomas Mentzel, MD and Matthew R. Lindberg, MD
- 358 Myopericytoma Matthew R. Lindberg, MD and Thomas Mentzel, MD
- 362 Myofibroma and Myofibromatosis
  Matthew R. Lindberg, MD
- **366** Angioleiomyoma Matthew R. Lindberg, MD



## SECTION 9: TUMORS OF SKELETAL MUSCLE

#### **BENIGN**

- 370 Focal Myositis
  - Matthew R. Lindberg, MD and Cyril Fisher, MD, DSc, FRCPath
- **372** Adult Rhabdomyoma *Matthew R. Lindberg, MD*
- 374 Fetal Rhabdomyoma

  Matthew R. Lindberg, MD
- 376 Genital Rhabdomyoma Matthew R. Lindberg, MD
- 378 Cardiac Rhabdomyoma Matthew R. Lindberg, MD

#### **MALIGNANT**

- 380 Embryonal Rhabdomyosarcoma Matthew R. Lindberg, MD
- 386 Alveolar Rhabdomyosarcoma Matthew R. Lindberg, MD
- 392 Spindle Cell Rhabdomyosarcoma Matthew R. Lindberg, MD
- 396 Sclerosing Rhabdomyosarcoma Matthew R. Lindberg, MD
- 400 Pleomorphic Rhabdomyosarcoma Jerad M. Gardner, MD
- **404 Epithelioid Rhabdomyosarcoma** *Matthew R. Lindberg, MD*



# SECTION 10: VASCULAR TUMORS (INCLUDING LYMPHATICS)

#### **BENIGN**

- 408 Papillary Endothelial Hyperplasia
  David Cassarino, MD, PhD and Amitabh Srivastava, MD
- **410** Bacillary Angiomatosis

  David Cassarino, MD, PhD
- **412 Congenital Hemangioma** *Kandi Stallings-Archer, MD*
- 416 Infantile Hemangioma Kandi Stallings-Archer, MD
- **420 Lobular Capillary Hemangioma** *Matthew R. Lindberg, MD*
- 422 Epithelioid Hemangioma Matthew R. Lindberg, MD
- **426 Spindle Cell Hemangioma** *Jerad M. Gardner, MD*
- 430 Intramuscular Hemangioma
  Matthew R. Lindberg, MD and Jonathan B. McHugh, MD
- **434** Hobnail Hemangioma David Cassarino, MD, PhD
- **436** Acquired Tufted Angioma David Cassarino, MD, PhD
- **438** Microvenular Hemangioma David Cassarino, MD, PhD
- **440 Sinusoidal Hemangioma** *David Cassarino, MD, PhD*
- **442** Glomeruloid Hemangioma David Cassarino, MD, PhD
- **444** Angiomatosis

  David Lucas, MD
- **446** Lymphangioma
  David Cassarino, MD, PhD
- **450** Massive Localized Lymphedema *Matthew R. Lindberg, MD*
- **452** Atypical Vascular Lesion David Lucas, MD

#### **INTERMEDIATE (LOCALLY AGGRESSIVE)**

**454** Kaposiform Hemangioendothelioma David Lucas, MD

#### **INTERMEDIATE (RARELY METASTASIZING)**

- **456** Papillary Intralymphatic Angioendothelioma David Cassarino, MD, PhD
- **458** Retiform Hemangioendothelioma David Cassarino, MD, PhD
- **460** Composite Hemangioendothelioma David Lucas, MD
- **462** Pseudomyogenic Hemangioendothelioma Matthew R. Lindbera, MD

#### **MALIGNANT**

- **466** Epithelioid Hemangioendothelioma
  Matthew R. Lindberg, MD and Thomas Mentzel, MD
- 470 Angiosarcoma
- Matthew R. Lindberg, MD
- **476 Kaposi Sarcoma** *Jerad M. Gardner, MD and Thomas Mentzel, MD*

## SECTION 11: CHONDROOSSEOUS TUMORS

#### **BENIGN**

- 486 Soft Tissue Chondroma
  - David Lucas, MD
- **492** Synovial Chondromatosis David Lucas, MD

#### **MALIGNANT**

- 496 Extraskeletal Osteosarcoma
  - David Lucas, MD
- 500 Extraskeletal Mesenchymal Chondrosarcoma David Lucas, MD



#### **BENIGN**

- 506 Solitary Circumscribed Neuroma
- Jerad M. Gardner, MD
  508 Schwannoma

Matthew R. Lindberg, MD

- 522 Neurofibroma
  - Matthew R. Lindberg, MD
- 530 Perineurioma
  - Matthew R. Lindberg, MD
- **536 Hybrid Nerve Sheath Tumor** *Matthew R. Lindberg, MD*
- **540 Granular Cell Tumor** *Matthew R. Lindberg, MD*
- 546 Dermal Nerve Sheath Myxoma
- David Lucas, MD

  550 Ganglioneuroma
  - Matthew R. Lindberg, MD
- **Neuromuscular Choristoma** *David Lucas, MD*

#### **INTERMEDIATE**

556 Melanotic Schwannoma Matthew R. Lindberg, MD

#### **MALIGNANT**

- 558 Malignant Peripheral Nerve Sheath Tumor Matthew R. Lindberg, MD
- 566 Epithelioid Malignant Peripheral Nerve Sheath Tumor
  - Matthew R. Lindberg, MD

    Compared to the Matthew R. Lindberg, MD

    Compared to the Matthew R. Lindberg, MD
    - Matthew R. Lindberg, MD and Cyril Fisher, MD, DSc, FRCPath

# SECTION 13: GENITAL STROMAL TUMORS

- **574 Fibroepithelial Stromal Polyp** *Matthew R. Lindberg, MD*
- **576** Angiomyofibroblastoma *Matthew R. Lindberg, MD*
- 580 Cellular Angiofibroma Matthew R. Lindberg, MD
- 584 Deep (Aggressive) Angiomyxoma Matthew R. Lindbera. MD

## SECTION 14: TUMORS OF MESOTHELIAL CELLS

#### **BENIGN**

- **590** Adenomatoid Tumor Matthew R. Lindberg, MD
- 592 Multicystic Peritoneal Mesothelioma
  Matthew R. Lindbera. MD
- 594 Well-Differentiated Papillary Mesothelioma

  David Lucas, MD and Cyril Fisher, MD, DSc, FRCPath

#### **MALIGNANT**

- 598 Malignant Mesothelioma
  - Matthew R. Lindberg, MD and Cyril Fisher, MD, DSc, FRCPath

# SECTION 15: HEMATOPOIETIC TUMORS IN SOFT TISSUE

- **Solitary Extramedullary Plasmacytoma** *Matthew R. Lindberg, MD*
- 608 Myeloid Sarcoma
  Matthew R. Lindberg, MD
- 610 Lymphoma of Soft Tissue Matthew R. Lindberg, MD and Khin Thway, MD, FRCPath



#### **BENIGN**

**614 Intramuscular Myxoma** *Matthew R. Lindberg, MD* 

618	<b>Juxtaarticular Myxoma</b> Matthew R. Lindberg, MD and Khin Thway, MD, FRCPath	728	Undifferentiated Round Cell Sarcoma With <i>CIC-DUX</i> Translocation
620	Superficial Angiomyxoma  Jerad M. Gardner, MD	734	David Lucas, MD  BCOR-CCNB3 Fusion-Positive Sarcoma
624	Acral Fibromyxoma Matthew R. Lindberg, MD	_	David Lucas, MD
626	Pleomorphic Hyalinizing Angiectatic Tumor Matthew R. Lindberg, MD and Cyril Fisher, MD, DSc,		SECTION 18: MESENCHYMAL TUMORS OF GASTROINTESTINAL TRACT
630	FRCPath Aneurysmal Bone Cyst of Soft Tissue	740	<b>Benign Neural Gastrointestinal Polyps</b> <i>Matthew R. Lindberg, MD</i>
632	Matthew R. Lindberg, MD  Ectopic Hamartomatous Thymoma	744	Gastrointestinal Stromal Tumor Matthew R. Lindberg, MD
	Matthew R. Lindberg, MD	760	Gastrointestinal Schwannoma Matthew R. Lindberg, MD
	ERMEDIATE (LOCALLY AGGRESSIVE)	762	Gastrointestinal Smooth Muscle Neoplasms
634	Hemosiderotic Fibrolipomatous Tumor Matthew R. Lindberg, MD	766	Matthew R. Lindberg, MD Inflammatory Fibroid Polyp
INT	ERMEDIATE (RARELY METASTASIZING)	770	Matthew R. Lindberg, MD Gangliocytic Paraganglioma
636	Atypical Fibroxanthoma	772	Matthew R. Lindberg, MD  Plexiform Fibromyxoma
642	David Cassarino, MD, PhD  Angiomatoid Fibrous Histiocytoma		Matthew R. Lindberg, MD
650	Matthew R. Lindberg, MD Ossifying Fibromyxoid Tumor	776	Malignant Gastrointestinal Neuroectodermal Tumo Matthew R. Lindberg, MD
656	Matthew R. Lindberg, MD  Myoepithelioma of Soft Tissue		SECTION 19: OTHER ENTITIES
	Matthew R. Lindberg, MD	BE	
664		<b>BE</b> 782	NIGN Amyloidoma
664	Matthew R. Lindberg, MD  Phosphaturic Mesenchymal Tumor		NIGN Amyloidoma Matthew R. Lindberg, MD
664	Matthew R. Lindberg, MD  Phosphaturic Mesenchymal Tumor  Matthew R. Lindberg, MD  LLIGNANT  Synovial Sarcoma	782 784	NIGN Amyloidoma Matthew R. Lindberg, MD Ganglion Cyst Matthew R. Lindberg, MD
664 MA	Matthew R. Lindberg, MD  Phosphaturic Mesenchymal Tumor  Matthew R. Lindberg, MD  LLIGNANT	782 784 786	NIGN  Amyloidoma  Matthew R. Lindberg, MD  Ganglion Cyst  Matthew R. Lindberg, MD  Tumoral Calcinosis  Matthew R. Lindberg, MD
664 MA 666 678	Matthew R. Lindberg, MD Phosphaturic Mesenchymal Tumor Matthew R. Lindberg, MD  LIGNANT Synovial Sarcoma Matthew R. Lindberg, MD Epithelioid Sarcoma Matthew R. Lindberg, MD Matthew R. Lindberg, MD	782 784	NIGN Amyloidoma Matthew R. Lindberg, MD Ganglion Cyst Matthew R. Lindberg, MD Tumoral Calcinosis
MA 666 678 684	Matthew R. Lindberg, MD Phosphaturic Mesenchymal Tumor Matthew R. Lindberg, MD  LIGNANT  Synovial Sarcoma Matthew R. Lindberg, MD  Epithelioid Sarcoma Matthew R. Lindberg, MD  Alveolar Soft Part Sarcoma Matthew R. Lindberg, MD	782 784 786	NIGN  Amyloidoma Matthew R. Lindberg, MD  Ganglion Cyst Matthew R. Lindberg, MD  Tumoral Calcinosis Matthew R. Lindberg, MD  Idiopathic Tumefactive Fibroinflammatory Lesions Matthew R. Lindberg, MD  Cardiac Myxoma
664 MA 666 678	Matthew R. Lindberg, MD Phosphaturic Mesenchymal Tumor Matthew R. Lindberg, MD  LIGNANT Synovial Sarcoma Matthew R. Lindberg, MD Epithelioid Sarcoma Matthew R. Lindberg, MD Alveolar Soft Part Sarcoma	782 784 786 788	NIGN  Amyloidoma Matthew R. Lindberg, MD  Ganglion Cyst Matthew R. Lindberg, MD  Tumoral Calcinosis Matthew R. Lindberg, MD  Idiopathic Tumefactive Fibroinflammatory Lesions Matthew R. Lindberg, MD  Cardiac Myxoma Matthew R. Lindberg, MD  Cardiac Fibroma
MA 666 678 684	Matthew R. Lindberg, MD Phosphaturic Mesenchymal Tumor Matthew R. Lindberg, MD  LIGNANT  Synovial Sarcoma Matthew R. Lindberg, MD Epithelioid Sarcoma Matthew R. Lindberg, MD Alveolar Soft Part Sarcoma Matthew R. Lindberg, MD Clear Cell Sarcoma Jerad M. Gardner, MD Perivascular Epithelioid Cell Tumor (PEComa)	782 784 786 788 792 796	NIGN  Amyloidoma Matthew R. Lindberg, MD  Ganglion Cyst Matthew R. Lindberg, MD  Tumoral Calcinosis Matthew R. Lindberg, MD  Idiopathic Tumefactive Fibroinflammatory Lesions Matthew R. Lindberg, MD  Cardiac Myxoma Matthew R. Lindberg, MD
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#### SECTION 17: UNDIFFERENTIATED/UNCLASSIFIED SARCOMAS

**724 Undifferentiated Pleomorphic Sarcoma** *Matthew R. Lindberg, MD* 

Matthew R. Lindberg, MD
 Peripheral Hemangioblastoma
 Matthew R. Lindberg, MD
 Melanotic Neuroectodermal Tumor of Infancy
 Matthew R. Lindberg, MD

Paraganglioma

# **826** Ependymoma of Soft Tissue Matthew R. Lindberg, MD

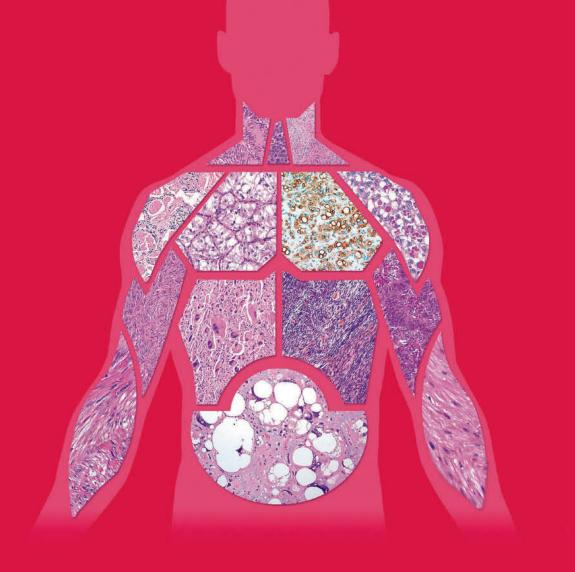
#### MALIGNANT

**FRCPath** 

1 - 17		
828	Metastatic Tumors to Soft Tissue Sites Matthew R. Lindberg, MD	
832	Neuroblastoma and Ganglioneuroblastoma Kandi Stallings-Archer, MD, Jessica M. Comstock, MD, and Cyril Fisher, MD, DSc, FRCPath	
842	Extraaxial Soft Tissue Chordoma Jerad M. Gardner, MD	
844	Undifferentiated Embryonal Sarcoma of Liver Matthew R. Lindberg, MD	
846	Primary Pulmonary Myxoid Sarcoma David Lucas, MD	
850	Biphenotypic Sinonasal Sarcoma Matthew R. Lindberg, MD	
854	Spindle Epithelial Tumor With Thymus-Like Differentiation Matthew R. Lindberg, MD	
856	Low-Grade Endometrial Stromal Sarcoma	

Charles Matthew Quick, MD and Khin Thway, MD,

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# Soft Tissue Tumors

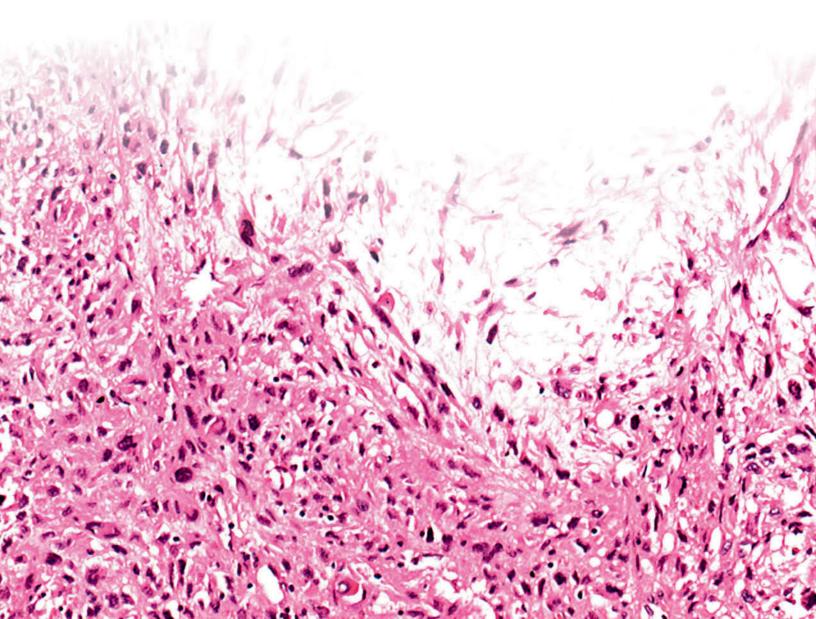
Lindberg



# Section 1 Soft Tissue Introduction

### Introduction and Overview

Gross Examination Grading and Staging	4 6
Ancillary Techniques	
Soft Tissue Immunohistochemistry	
Molecular Features of Soft Tissue Tumors	18



#### INTRODUCTION

#### Overview

- Thorough but focused gross examination is vital component of overall evaluation and diagnostic work-up of soft tissue tumors
- Common errors (e.g., undersampling, inappropriate sampling, not inking margins) can severely hamper or preclude accurate diagnosis, pathologic staging, and subsequent clinical planning
- Review of patient clinical history and information prior to gross examination is strongly recommended

#### **CLINICAL FINDINGS**

#### History

- Review any available clinical notes or operative report
  - Note age of patient and clinical presentation of tumor
  - o Note if patient has prior history of tumor in same anatomic location, nearby, or elsewhere
- Determine whether tumor has been previously biopsied or treated or if there is established diagnosis

#### **Imaging**

- Review any pertinent radiographs, CT, or MR scans
  - o Determine if tumor is homogeneous or heterogeneous
  - o Identify any notable structures involved (e.g., large nerve
  - o Determine whether radiologist favors benign or malignant process
- Note anatomic location (e.g., thigh, neck, retroperitoneum,
- Note tissue plane (i.e., superficial/subcutaneous vs. deep/intramuscular)

#### **GROSSING PROCEDURE**

#### **External Examination**

a soft tissue tumor specimen varies depending on the type of surgery, but many tumors (especially sarcomas) are removed with at least a thin rim of surrounding soft tissue. If the tumor has been previously sampled by core

well. (Right) Although known benian soft tissue tumors are often excised without orientation by the surgeon,

by stitches and require inking, as depicted, for satisfactory margin evaluation.

- Specimen should be weighed and measured in 3 dimensions
- Describe external appearance and shape of mass

- Note any orientation provided by surgeon (e.g., stitches, strip of overlying skin, large nerves)
  - Skin is often excised to remove previous biopsy tract with rest of specimen
- Ink peripheral margins of specimen
  - o Oriented tumors often require inking with up to 6 different colors
  - o Unoriented tumors may be inked in 1 color

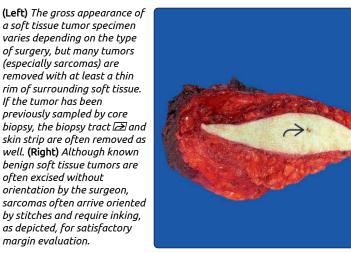
#### Sectioning and Internal Evaluation

- Serially section ("bread loaf") mass in 1-cm thick sections, perpendicular to long axis of specimen
- Lay out all slabs of tumor and examine cut surfaces
- Describe appearance of cut surface and note areas of different coloration &/or texture
  - o Common colors: Tan, white, gray, red, brown
  - Common textures: Firm/fibrous, fleshy, gelatinous/glistening, fatty
- Note any areas of hemorrhage &/or necrosis
  - o Quantify necrosis (none, ≤ 50%, or > 50%)
  - o Highly necrotic tumors should be placed in formalin to fix and to minimize fragmentation
- Take representative fresh tissue for possible ancillary techniques or treatment protocols (may be snap frozen)

#### Sampling

- Standard approach is to take 1 section per cm of greatest tumor dimension (margin sections counted separately)
  - o e.g., 16-cm tumor gets 16 blocks with 1 tissue piece in each or 8 blocks with 2 tissue pieces in each
  - o Fewer sections may be submitted for large tumors with diffuse homogeneous appearance
- Inked margins, specifically close (< 2 cm) margins, should be sampled with perpendicular sections
- Sections should be taken from all distinctive areas (e.g., fibrous, gelatinous, fleshy, etc.)
  - o Sections taken at interface between different areas can provide very useful histologic information
- Obviously necrotic areas should be minimally sampled
  - o These areas often represent high-grade morphologies and are generally less useful diagnostically

#### **External Examination**



#### Specimen Inking

