Timothy R. Deer Jason E. Pope Tim J. Lamer David Provenzano *Editors*

Deer's Treatment of Pain

An Illustrated Guide for Practitioners



Deer's Treatment of Pain

Timothy R. Deer • Jason E. Pope Tim J. Lamer • David Provenzano Editors

Deer's Treatment of Pain

An Illustrated Guide for Practitioners



Editors Timothy R. Deer The Spine and Nerve Center of the Virginias Charleston, WV USA

Tim J. Lamer Department of Anesthesiology and Perioperative Medicine Mayo Clinic Rochester, MN USA Jason E. Pope Evolve Restorative Center Santa Rosa, CA USA

David Provenzano Pain Diagnostics and Interventional Care Sewickley, PA USA

ISBN 978-3-030-12280-5 ISBN 978-3-030-12281-2 (eBook) https://doi.org/10.1007/978-3-030-12281-2

© Springer Nature Switzerland AG 2019

This work is subject to copyright. All rights are reserved by the Publisher, whether the whole or part of the material is concerned, specifically the rights of translation, reprinting, reuse of illustrations, recitation, broadcasting, reproduction on microfilms or in any other physical way, and transmission or information storage and retrieval, electronic adaptation, computer software, or by similar or dissimilar methodology now known or hereafter developed.

The use of general descriptive names, registered names, trademarks, service marks, etc. in this publication does not imply, even in the absence of a specific statement, that such names are exempt from the relevant protective laws and regulations and therefore free for general use.

The publisher, the authors, and the editors are safe to assume that the advice and information in this book are believed to be true and accurate at the date of publication. Neither the publisher nor the authors or the editors give a warranty, express or implied, with respect to the material contained herein or for any errors or omissions that may have been made. The publisher remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

This Springer imprint is published by the registered company Springer Nature Switzerland AG The registered company address is: Gewerbestrasse 11, 6330 Cham, Switzerland

This book is possible due to the support and love of my wife Missy, who has been by my side for many wonderful years. I also dedicate this book to God, who has blessed me through grace well beyond my greatest expectations. I also thank my children, Morgan, Taylor, Reed, and Bailie, for inspiring me in so many ways. Lastly, to our patients, I am eternally grateful for your trust and am hopeful that my work will impact your life in a way that is meaningful.

Timothy R. Deer, MD, DABPM, FIPP

To Emily, my wife, CHRO, and partner. Thank you for your unwavering support, grace, and the selflessness through the many years we have spent together. I love you more than words can express and humbled by your hard work you put in every day for our family and our practice. We are together in all adventures.

To my daughter Vivienne. Thank you for your kindness. You are growing up into a kind and talented girl and I enjoy all we do together.

To my twins Liam and Olivia. Thank you for your love and adventurous, brave outlook. It reminds me anything is possible.

To my father and mother. Thank you for your unyielding love and advice as I navigate the balance of father, son, husband, and work.

To my twin brother, who always humbled me with his intellect and perspective and leading by example.

To Eric Bruntlett, Steven Falowski, and Michael Fishman, for their friendship, candor, advice, and guidance.

To my entire Evolve Restorative Center family. Thank you for your willingness to help patients and going the extra mile to make sure our patients are cared for and supported. We are building something great.

To Timothy Deer, my brother, thank you for your friendship, support, advice, and partnership. We will continue to work towards globalization of pain care to improve safety and outcomes. We are not victims of circumstance.

To Springer, for their continued work in making this text edition a reality.

Finally, thank you to God, for making all things possible.

Jason E. Pope, MD, DABPM, FIPP

To my wife Halena, my best friend, advisor, and counselor. Your love, support, and companionship are so cherished and appreciated by me. And to our amazing children, Olivia, Tyler, Elizabeth, and Allison. We are so grateful for your love and watching you grow into the wonderful and exceptional young adults that you have all become is the most important and inspirational part of our lives.

To my patients, whom I have had the honor and pleasure of caring for over the past 30 plus years. I have learned more from you than all of my 30 plus years of training and education.

To my Mayo Clinic colleagues. It is an honor to work with such a dedicated and talented group of people in an amazing medical facility.

To all of the students, residents, and fellows that I have been honored to help teach and mentor. Watching your evolution into successful physicians and individuals is both gratifying and inspirational.

Tim J. Lamer, MD

To Dana, Nora, and Marco. Thank you for all your support and understanding (Another project that kept me in the study!).

To all my previous educators. Without your guidance and instruction, I would have never acquired the knowledge required to pursue this project.

Finally, to our patients. It is an honor to help you deal with the challenges of pain. I continue to learn from these experiences daily.

David Provenzano, MD

Foreword

I am honored to write this foreword for *Deer's Treatment of Pain: An Illustrated Guide for Practitioners.* As an interventional pain management specialist, I have devoted my career to the treatment of patients with pain and share Tim Deer's passion for patient care, clinical research, and education of the next generation of pain physicians. I have known Dr. Deer for two decades as a friend and colleague and have watched with admiration and respect as he has become among the most highly regarded figures in the field. His leadership in interventional pain is internationally recognized, and he is directly responsible for much of the sentinel work in many areas of pain therapy, particularly neuromodulation and intrathecal drug delivery.

Dr. Deer is an accomplished triathlete. His charitable work with the Ironman Foundation exemplifies the generosity and integrity he brings to all that he does, including the publication of this important textbook, which establishes a new standard reference for our field. This comprehensive resource includes over 100 chapters written by today's thought leaders in every area of pain treatment. It provides the reader with the basic science, anatomy, physiology, and practical treatment considerations necessary to successfully navigate the ever-increasing complexity of interventional pain management.

A recent study by the National Institutes of Health's National Center for Complementary and Integrative Health indicates that nearly 50 million American adults experience significant chronic pain or severe pain [1]. Because pain is one of the most common reasons patients seek help from healthcare providers, it is critical that physicians are well versed in its diagnosis and management. I commend Timothy R. Deer and his colleagues, Jason E. Pope, Tim J. Lamer, and David Provenzano, for assembling this comprehensive volume which details the current understanding of both pain and its treatment. Sections address both pharmacologic and interventional approaches to pain control, as well as neuromodulation, intrathecal drug delivery systems, regenerative therapies, and minimally invasive structural surgeries. Well-placed illustrations and other graphics serve to enhance the text, further increasing the book's value. In summary, the expertise presented in *Deer's Treatment of Pain* provides us with an authoritative reference work that belongs on the desk of every student, clinician, and scientist involved in the study and treatment of pain.

Aaron Calodney, MD, FIPP, ABIPP, DABPM Director, Clinical Research, Precision Spine Care Clinical Associate Professor of Anesthesiology Louisiana State University Medical School, Shreveport Shreveport, LA USA

Reference

 Nahin RL. Estimates of pain prevalence and severity in adults: United States, 2012. J Pain. 2015;16(8):769–80.

Preface

Chronic pain is an international issue that impacts the well-being of all nations for numerous reasons. In the past decade, we have come to the realization that the opioid epidemic is a tremendous challenge and has taken a huge toll on society. In that same complex setting, we also have the challenge of an aging population, traumatic injuries, disease progression, and sportsrelated ailments all of which cause chronic pain syndromes that require solutions. It is in this complicated time that we feel new guidance is needed for the physician or healthcare provider who is attempting to do an optimal job in treating this patient population. This is the environment that created the need for *Deer's Treatment of Pain*.

This textbook has been created with a number of committed physicians who are striving to improve therapeutic efficacy, patient safety, and ethical medical treatment. This is intended to merge a comprehensive text with an atlas feeling where you can visually learn best treatment options for your patient. The physician who embraces these options of an algorithmic treatment approach based on best evidence, expert opinion, and clinical experience will enhance their ability to provide optimal care to those who suffer and seek a remedy.

In developing this educational material, I had the honor of working with Tim Lamer, MD; David Provenzano, MD; and Jason E. Pope, MD. This team of dedicated physicians has created a group who shaped the contents, edited the materials, suggested additions, and used the highest standards to make a commitment to creating an optimal learning experience for the reader.

I wish each of you the best of success as you raise the standard of care in your community by continuing to strive to learn new options and methods. It is my belief that by working together, we can truly offer solutions that will better the human experience.

Charleston, WV, USA

Timothy R. Deer

Contents

Part I Introduction

1	The Disease-Based Treatment of Pain. 3 Andrew So and Karina Gritsenko 3	
2	Algorithms of Pain Treatment 13 Andrew So and Karina Gritsenko 13	
3	The Opioid Epidemic and the Need for a Pain Strategy	
Part	II Anatomy and Physiology of Pain	
4	Nerve Function and Neurons	
5	Peripheral Nerve Anatomy. 35 Daniel Rothstein and Didier Demesmin 35	
6	Spinal Cord Anatomy 43 Alan Gonzalez Cota 43	
7	Anatomy of the Brain and Brain Stem	
8	Mediators of Pain and Pain Processing	
9	Taxonomy of Pain 75 Nicholas J. Bremer 75	
10	Pain-Relieving Mechanisms in Neuromodulation79Vikram Sengupta, Sascha Qian, Ned Urbiztondo, and Nameer Haider	
Part III Psychology of Pain		
11	The Normal Response to Pain 93 Randall P. Brewer 93	
12	Pain and Suffering	
13	Social Impact of Pain Response	
14	Role of Religion and Spirituality in the Patient Pain Experience	

15	Anxiety and Depression in Patients with Chronic Pain
16	Psychological Treatments to Improve Outcomes
17	Psychological Evaluation for Those Receiving Devicesfor the Treatment of PainIoannis M. Skaribas and Kevin Smith
Part	IV Pharmacological Treatment of Pain
18	Acetaminophen and Nonsteroidal Anti-inflammatory Drugs
19	Anticonvulsants in the Treatment of Pain
20	Botulinum Toxin
21	Sodium Channel Antagonists
22	Antispasmodics and Muscle Relaxants
23	Antidepressants in Pain Management
24	Ketamine and NMDA-Receptor Antagonists
25	Novel Analgesics
26	Injectable Corticosteroids
27	Topical Therapies
28	Opioids for Chronic Non-cancer Pain
29	Opioids for Cancer Pain and Hospice Care
30	Pharmacogenetics and Pharmacogenomics of Pain Treatment
Part	V Interventional Treatment of Pain
31	Radiation Safety
32	Radiofrequency Ablation

33	Basic Science of Radio Frequency 275 Tiffany Lin, Simon Willis, Dost Khan, and Maunak V. Rana 275
34	Cryotherapy
35	Chemodenervation: Neurolytic Blockade and Potent Neurotoxins for the Treatment of Cancer Pain
36	Blockade of the Nerves of the Head and Face
37	Ganglion Blocks of the Head and Face
38	Destructive Procedures of the Head and Face
39	Interlaminar Approach for Cervical Epidural Steroid Injection
40	Cervical Transforaminal Epidural Injections
41	Diagnostic Cervical Nerve Root Blocks
42	Cervical Facet Joint Injection and Medial Branch Blocks
43	Cervical Facet Radiofrequency Neurotomy
44	Intercostal Nerve Block
45	Thoracic Transforaminal Epidural Access
46	Thoracic Interlaminar Epidural Steroid Injection
47	Thoracic Facet and Medial Branch Blocks. 389Rebecca A. Sanders
48	Radiofrequency Ablation in the Thoracic Spine
49	Lumbar Interlaminar Epidural Injection
50	Lumbar Transforaminal Epidural Corticosteroid Injections
51	Lumbar Facet and Medial Branch Block
52	Sacroiliac Joint Injection
53	Sacroiliac Joint Radiofrequency

54	Sacroiliac Joint Fusion
55	Caudal Epidural Injection
56	Stellate Ganglion Block
57	Lumbar Sympathetic Block
58	Celiac Plexus Block and Superior Hypogastric Plexus Block
59	Ganglion Impar Block
60	Large Joint Injections
Part	VI Neuromodulation
61	Deep Brain Stimulation
62	Motor Cortex Stimulation
63	Occipital Nerve Stimulation
64	Sphenopalatine Ganglion Block.531Mark N. Malinowski and Nicholas J. Bremer
65	Percutaneous Cervical Spinal Cord Stimulation
66	Surgical Leads for the Cervical Spine
67	Dorsal Root Ganglion Stimulation: Cervical Spine
68	Percutaneous Thoracic Neurostimulation for Chronic Pain
69	Surgical Lead for the Thoracic Spine
70	Percutaneous Lumbar Stimulation
71	Thoracic and Lumbar Dorsal Root Ganglion Spinal Stimulation
72	Percutaneous Sacral Nerve Stimulation

xvi

73	Sacral Dorsal Root Ganglion Spinal Stimulation
74	Peripheral Nerve Stimulation
75	Stimulation Methods and Device Choices: Dorsal Root Ganglion Stimulation, Spinal Cord Stimulation, and Peripheral Nerve Stimulation 615 Seth Christian, Vafi Salmasi, and Michael S. Leong
76	Patient Selection
77	Wound Closure and Surgical Healing629Stephen D. Coleman, Vafi Salmasi, and Michael S. Leong
78	Complications of Neuromodulation
Part	VII Neuroaxial Therapies
79	Intrathecal Pharmacology
80	Patient Selection
81	Intrathecal Device Considerations
82	Intrathecal Agents and Algorithms:Review of PACC 2012 and 2017 Guidelines and BeyondMichael S. Leong, Lynn K. Ngai, and William A. Stuart
83	Implantable Drug Delivery Systems for Cancer Pain and End of Life Care 675 Ann Cai Shah, Kenneth Ike, Lisa Stearns, and Lawrence R. Poree
84	Intrathecal Drug Delivery Systems for Chronic Non-cancer Pain
85	IDDS for Movement Disorders
86	Ziconotide for Intrathecal Use
Part	VIII Regenerative Therapies for Chronic Pain
87	Scientific Bases of PRP Therapy
88	Scientific Basis for Stem Cell Therapy
89	Stem Cells
90	Regenerative Therapies for Chronic Intradiscal Pain

Contents

91	PRP Therapies (Tendons, Joints, Spine)
Part	IX Intradiscal and Minimally Invasive Structural Surgeries
92	Discography
93	Endoscopic Discectomy
94	Vertebral Augmentation for Painful Vertebral Compression Fractures
95	Vertebroplasty and Other Methods of Vertebral Augmentation
96	Minimally Invasive Lumbar Decompression
97	Interspinous Process Spacers for Indirect Lumbar Decompression
Part	X Rehabilitation and Adjuvant Therapies for Pain
98	Physical Therapy
99	Acupuncture
Part XI Administrative Considerations	
100	Cybersecurity of Medical Devices: Past, Present, and Future
101	Coding and Billing
102	Medicare Payment Quality Measures
Inde	x

Contributors

Rishi Raj Agarwal, MD Department of Anesthesiology, Northwestern University, Feinberg School of Medicine, Chicago, IL, USA

Anuj Aggarwal, MD Department of Anesthesiology, Perioperative and Pain Medicine, Stanford University, Redwood City, CA, USA

Yamah Amiri, MBBS Pain Management, Ochsner Health System, New Orleans, LA, USA

Hamed Asadi, PhD, FRANZCR, CCINR, EBIR Interventional Neuroradiology Service, Department of Radiology, Austin Hospital, Melbourne, Australia

School of Medicine, Faculty of Health, Deakin University, Waurn Ponds, Australia

Interventional Neuroradiology Unit, Monash Imaging, Monash Health, Melbourne, Australia

Javid Baksh, DO, MD Premier Pain Solutions, Asheville, NC, USA

G. Baranidharan, FRCA, FFPMRCA Leeds Pain and Neuromodulation Centre, Leeds, UK

Markus A. Bendel, MD Mayo Clinic, Rochester, MN, USA

Nicholas J. Bremer, MD, D.ABA Center for Pain Relief, Spine and Nerve Centers of the Virginias, Charleston, WV, USA

Randall P. Brewer, MD Department of Anesthesiology, Duke University School of Medicine, Durham, NC, USA

Departments of Neurology and Anesthesiology, Louisiana State University School of Medicine, Shreveport, LA, USA

River Cities Interventional Pain Specialists, Shreveport, LA, USA

Taylor Brittan, BS Neurosurgery Department, Ochsner Health System, New Orleans, LA, USA

Abram H. Burgher The Pain Center of Arizona, Peoria, AZ, USA

Lucas W. Campos, MD, PhD Interventional Pain Physicians, Chico, CA, USA

Alexios G. Carayannopoulos Department of Physical Medicine and Rehabilitation, Rhode Island Hospital, Providence, RI, USA

Comprehensive Spine Center: Rhode Island Hospital, Providence, RI, USA

Newport Hospital, Newport, RI, USA

Warren Alpert Medical School, Brown University, Providence, RI, USA

American Society of Interventional Pain Physicians, Rhode Island Chapter, Paducah, KY, USA

Jonathan D. Carlson, MD Hawaii Pain & Spine, Kailua, HI, USA

Midwestern Medical School, Glendale, AZ, USA

Hawaii Pain & Spine, Arizona Pain, Kailua, HI, USA

Nicole S. Carter Faculty of Medicine, Nursing and Health Sciences, Monash University, Melbourne, Australia

Ronil V. Chandra Faculty of Medicine, Nursing and Health Sciences, Monash University, Melbourne, Australia

Interventional Neuroradiology Unit, Monash Imaging, Monash Health, Melbourne, Australia

Shamard Charles, MD Harvard T.H. Chan School of Public Health, NBC News Group, New York, NY, USA

Tony Y. Chon, MD Division of General Internal Medicine, Mayo Clinic, Rochester, MN, USA

Seth Christian, MD, MBA Paradigm Orthopedics and Sports Medicine, Slidell, LA, USA

Steven P. Cohen, MD Department of Anesthesiology and Critical Care Medicine, Johns Hopkins School of Medicine, Baltimore, MD, USA

Department of Physical Medicine and Rehabilitation, Johns Hopkins School of Medicine, Baltimore, MD, USA

Department of Anesthesiology, Uniformed Services University of the Health Sciences, Bethesda, MD, USA

Department of Physical Medicine and Rehabilitation, Uniformed Services University of the Health Sciences, Bethesda, MD, USA

Stephen D. Coleman, MD Anesthesia, Perioperative and Pain Medicine, Stanford University, Stanford, CA, USA

David Copenhaver, MD, MPH UC Davis Cancer Pain Management and Supportive Care, University of California, Davis Division of Pain Medicine, Davis, CA, USA

Alan Gonzalez Cota, MD, FIPP A.I.M. Interventional Spine Division, Rockville, MD, USA

Legacy Pain Management Rehabilitation, Bethesda, MD, USA

KureSmart Pain Management, Annapolis, MD, USA

Timothy R. Deer, MD, DABPM, FIPP The Spine and Nerve Center of the Virginias, Charleston, WV, USA

Didier Demesmin, MD University Pain Medicine Center, Somerset, NJ, USA

Rutgers Robert Wood Johnson Medical School, Piscataway, NJ, USA

JFK Johnson Rehabilitation Institute, Edison, NJ, USA

Pain Medicine Department, Saint Peter's University Hospital, New Brunswick, NJ, USA

University Clinical Research Center, Somerset, NJ, USA

Daniel R. Denis, MD, FRCS(C) Ochsner Health System, Kenner, LA, USA

Mehul J. Desai, MD, MPH International Spine, Pain & Performance Center, Washington, DC, USA

George Washington University, School of Medicine, Washington, DC, USA

James Dierkes, MD Department of Anesthesiology, Duke University Medical Center, Durham, NC, USA

Dan DuBose, MD

Sam Eldabe, MBChB, FRCA, FFPMRCA Department of Pain and Anaesthesia, James Cook University Hospital, Middlesbrough, UK

Jennifer L. Erian, MD Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Steven Falowski, MD Functional Neurosurgery, Neurosurgical Associates of Lancaster, Lancaster, PA, USA

Kris Ferguson, MD Aspirus Hospital, Wausau, WI, USA

Christina E. Fitch, DO, MPH University of Massachusetts Medical School- Baystate Campus, Springfield, MA, USA

Alexander Fontenot, MD Anesthesiology Department, Ochsner Health system, New Orleans, LA, USA

Rishi Gaiha, MD Department of Anesthesiology, Northwestern University, Feinberg School of Medicine, Chicago, IL, USA

Christopher Gates Velentium LLC, Katy, TX, USA

Halena M. Gazelka, MD Division of Pain Medicine, Department of Anesthesiology and Perioperative Medicine, Mayo Clinic College of Medicine and Science, Rochester, MN, USA

Tony George, DO University Pain Medicine Center, Somerset, NJ, USA

Christopher Gilligan, MD, MBA Brigham and Women's Hospital, Boston, MA, USA

Vitaly Gordin, MD Department of Anesthesiology and Perioperative Medicine, Pennsylvania State University, Hershey, PA, USA

Yuri Gordin, MD Department of Physical Medicine and Rehabilitation, Pennsylvania State University, Hershey, PA, USA

Edna Gouveia, MD Neurosurgery Department, Ochsner Health System, New Orleans, LA, USA

Patrick D. Grace, MD Department of Anesthesiology and Pain Medicine, The University of Kansas Health System, Kansas City, KS, USA

Karina Gritsenko, MD Albert Einstein College of Medicine/Montefiore Medical Center, Montefiore Multidisciplinary Pain Program, The Bronx, NY, USA

Alba Guevara, MD Department of Anesthesiology and Perioperative Medicine, Pennsylvania State University, Hershey, PA, USA

Maged Guirguis, MD Pain Management, Ochsner Health System, New Orleans, LA, USA Interventional Pain Management, Department of Anesthesiology & Critical Care Medicine, Ochsner Health System, New Orleans, LA, USA

University of Queensland Ochsner Medical School, Brisbane, QLD, Australia

Louisiana State University School of Medicine, New Orleans, LA, USA

Ochsner Medical Center, Jefferson, LA, USA

Ashish Gulve, FFPMCAI, FCARCSI, MD Department of Pain Management, The James Cook University Hospital, Middlesbrough, UK

Osama Hafez, MD Department of Anesthesiology, Moffitt Cancer Center, University of South Florida, Tampa, FL, USA

Jonathan Hagedorn, MD Department of Pain Medicine, Mayo Clinic, Rochester, MN, USA

Nameer Haider, MD, FAAPMR-AQPM, DABPM Spinal & Skeletal Pain Medicine, Utica, NY, USA

Ian Hakkinen, MBBS Pain Management, Ochsner Health System, New Orleans, LA, USA

Joshua Hanna, MD Neurosurgery Department, Ochsner Health System, New Orleans, LA, USA

Joshua A. Hirsch, MD Neuroendovascular Program, Massachusetts General Hospital, Boston, MA, USA

Harvard Medical School, Boston, MA, USA

Yasmine Hoydonckx, MD, FIPP Department of Anesthesia, Toronto Western Hospital, Toronto, ON, Canada

Christy Hunt, DO, MS Department of Physical Medicine and Rehabilitation, Mayo Clinic, Rochester, MN, USA

Corey W. Hunter, MD Department of Rehabilitation Medicine, Icahn School of Medicine at Mount Sinai, New York, NY, USA

Ainsworth Institute of Pain Management, New York, NY, USA

Kenneth Ike, MD Department of Anesthesia, Pain Management Clinic, Grady Hospital, Atlanta, GA, USA

Marilyn S. Jacobs, PhD, ABPP Department of Psychiatry and Biobehavioral Sciences, David Geffen School of Medicine at UCLA, Los Angeles, CA, USA

Sameer Jain, MD Department of Pain Medicine, Pain Treatment Centers of America, Little Rock, AR, USA

Piotr K. Janicki, MD, PhD Department of Anesthesiology and Perioperative Medicine, Penn State College of Medicine, MS Hershey Medical Center, Hershey, PA, USA

Gabriel P. Jasper Jasper Spine Institute, Brick, NJ, USA

Jasper Spine Learning Institution, Jersey Shore University, Neptune, NJ, USA

Navdeep Jassal, MD Department of Neurology, University of South Florida, Tampa, FL, USA

Mihir M. Kamdar, MD MGH Divisions of Palliative Care and Anesthesia Pain Medicine, Massachusetts General Hospital, Boston, MA, USA

Dost Khan, MD Department of Anesthesiology, Northwestern University Feinberg School of Medicine, Chicago, IL, USA

Chong H. Kim, MD MetroHealth/Case Western Reserve University, Department of PM&R, Cleveland, OH, USA

Chase A. Kissling, MD Walter Reed National Military Medical Center, Bethesda, MD, USA

Narayan R. Kissoon, MD Division of Headache, Department of Neurology, Mayo Clinic, Rochester, MN, USA

Division of Pain, Department of Anesthesiology, Mayo Clinic, Rochester, MN, USA

Kiran K. Koneti, FRCA, EDRA Department of Anaesthetics, Sunderland Royal Hospitals, Sunderland, UK

Brian Harris Kopell, MD Center for Neuromodulation, Department of Neurosurgery, Mount Sinai Health System, New York, NY, USA

Bryan K. Lai, MD Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Tim J. Lamer, MD Division of Pain Medicine, Department of Anesthesiology and Perioperative Medicine, Mayo Clinic, Rochester, MN, USA

Luke A. Law, MD Department of Anesthesiology, Mayo Clinic, Rochester, MN, USA

Eric T. Lee, MD, MA Musculoskeletal Care and Regenerative Medicine, St Charles Spine Institute, Thousand Oaks, CA, USA

Michael S. Leong, MD Stanford Pain Management Center, Redwood City, CA, USA

Thabele Leslie-Mazwi, MD Neuroendovascular Program, Massachusetts General Hospital, Boston, MA, USA

Harvard Medical School, Boston, MA, USA

Clarence T. Li, MD Departments of Neurology, Mayo Clinic, Rochester, MN, USA

Sean Li, MD Premier Pain Centers, An Affiliate of National Spine and Pain Centers, Shrewsbury, NJ, USA

Susan Lim, MD Brigham and Women's Hospital, Boston, MA, USA

Tiffany Lin, MD Department of Anesthesiology and Critical Care, The University of Chicago, Chicago, IL, USA

Paul J. Lynch, MD, DABA Arizona Pain Specialists, Scottsdale, AZ, USA

Ankit Maheshwari, MD Case Western Reserve University, Cleveland, OH, USA University Hospitals of Cleveland, Cleveland, OH, USA

Julian Maingard, MBBS, BMedSci Interventional Neuroradiology Service, Department of Radiology, Austin Hospital, Melbourne, Australia

School of Medicine - Faculty of Health, Deakin University, Waurn Ponds, Australia

Samia Malik, MD Department of Neurology, University of South Florida, Tampa, FL, USA

Mark N. Malinowski, DO, DABA Adena Spine Center, Chillicothe, OH, USA

Amanda Markow New York Presbyterian Brooklyn Methodist Hospital, New York, NY, USA

Tory L. McJunkin, MD, DABA Arizona Pain Specialists, Scottsdale, AZ, USA

W. Porter McRoberts, MD FAAPMR ABPMR ABA/Pain, University of Miami School of Medicine, Miami, FL, USA

Pankaj Mehta, MD, DABA, DABPM Pain Specialists of Austin, Killeen, TX, USA

Daniel Morgan, PhD Department of Anesthesiology and Perioperative Medicine, Pennsylvania State University, Hershey, PA, USA

Michael Mueller, MD, MPH Department of Anesthesiology and Perioperative Medicine, Pennsylvania State University, Hershey, PA, USA

Geeta Nagpal, MD Department of Anesthesiology, Northwestern University, Feinberg School of Medicine, Chicago, IL, USA

Andrew Ng Department of Anesthesiology, Jefferson Pain Center, Thomas Jefferson University Hospital, Philadelphia, PA, USA

Lynn K. Ngai, MD Stanford University, Stanford, CA, USA

Eric V. Ngo, MD Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Tyler Nix, MD, MPH Anesthesiology Department, Ochsner Health System, New Orleans, LA, USA

Devang Padalia, MD Interventional Pain Department, Moffitt Cancer Center, University of South Florida, Tampa, FL, USA

Dipan Patel, MD, DABA Garden State Pain Control Center, Clifton, NJ, USA

Residents & Fellows Society, North American Neuromodulation Society (NANS), Jersey City, NJ, USA

Vikram B. Patel, MD Phoenix Interventional Center for Advanced Learning, Algonquin, IL, USA

Sanket Pathak, MD Department of Anesthesiology, Thomas Jefferson University Hospital, Philadelphia, PA, USA

Denis G. Patterson, MD Nevada Advanced Pain Specialists, Reno, NV, USA

Christian Peccora, MD Texas Pain Consultant Associates, Houston, TX, USA

Philip Peng, MBBS, FRCPC Department of Anesthesia, Toronto Western Hospital, Toronto, ON, Canada

Jeffrey Peterson, BA COO, Center for Pain Relief, Inc. & The Spine and Nerve Centers of the Virginias, Charleston, WV, USA

Julie G. Pilitsis, MD, PhD Department of Neuroscience and Experimental Therapeutics, Neurosurgery, Albany Medical College, Albany, NY, USA

Thomas P. Pittelkow, DO, MPH Division of Pain Medicine, Department of Anesthesiology and Perioperative Medicine, Mayo Clinic College of Medicine and Science, Rochester, MN, USA

Jason E. Pope, MD, DABPM, FIPP Evolve Restorative Center, Santa Rosa, CA, USA

Lawrence R. Poree, MD, MPH, PhD Department of Anesthesia and Perioperative Care, University of California at San Francisco, San Francisco, CA, USA

Jose Posas III Pain Management, Ochsner Health System, New Orleans, LA, USA

Chane Price, MD Interventional Pain Physician, University of Miami, Coral Gables, FL, USA

David Provenzano, MD Pain Diagnostics and Interventional Care, Sewickley, PA, USA

Sascha Qian, MD Spinal & Skeletal Pain Medicine, Utica, NY, USA

Xiang Qian, MD, PhD Department of Anesthesiology, Perioperative and Pain Medicine, Stanford University, Stanford University Pain Management Center, Redwood City, CA, USA

Wenchun Qu, MD, PhD Department of Physical Medicine and Rehabilitation, Division of Pain Medicine, Spine Center, Mayo Clinic, Rochester, MN, USA

Maunak V. Rana, MD Department of Anesthesiology and Critical Care, The University of Chicago, Chicago, IL, USA

Corey Reeves, MD Department of Physical Medicine and Rehabilitation, University of South Florida, Tampa, FL, USA

Dustin Reynolds, MD Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Tina Rivenbark, MHA, CMPE Advanced Practice Solutions, Wilmington, NC, USA

Mathew Roberts, MS Medical University of South Carolina, Charleston, SC, USA

Daniel Rothstein, MD University Pain Medicine Center, Somerset, NJ, USA

Rutgers Robert Wood Johnson University Hospital, New Brunswick, NJ, USA

Pain Medicine Department, Saint Peter's University Hospital, New Brunswick, NJ, USA

Jeffery Rowe, MD Main Line Spine & Main Line Spine Surgery Center, King of Prussia, PA, USA

Vafi Salmasi, MD Department of Anesthesiology, Perioperative and Pain Medicine, Systems Neuroscience and Pain Laboratory, Stanford University School of Medicine, Redwood City, CA, USA

Rebecca A. Sanders, MD Department of Pain Management, Kansas Spine and Specialty Hospital, Wichita, KS, USA

Puneet Sayal, MD International Spine, Pain & Performance Center, Washington, DC, USA

Dawood Sayed, MD The University of Kansas Health System, Kansas City, KS, USA

Vikram Sengupta, MD Spinal & Skeletal Pain Medicine, Utica, NY, USA

Anish Sethi, DO Chronic Pain Medicine, Department of Anesthesiology, Thomas Jefferson University Hospital, Philadelphia, PA, USA

Ann Cai Shah, MD Department of Anesthesia and Perioperative Care, Anesthesia and Pain Medicine, University of California at San Francisco, San Francisco, CA, USA

Bunty J. Shah, MD Penn State Milton S. Hershey Medical Center, Hershey, PA, USA

Jay M. Shah, MD Department of Pain Medicine, Weill-Cornell Tri-Institutional Pain Program, New York, NY, USA

Neal Shah, MD Interventional Pain Department, Moffitt Cancer Center, University of South Florida, Tampa, FL, USA

Aanchal Sharma, MD Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Jaspreet Singh, MD Department of Neurology, University of South Florida, Tampa, FL, USA

Elena E. Skaribas Southern Methodist University, Dallas, TX, USA

Ioannis M. Skaribas, MD, DABA Expert Pain P.A., Department of Anesthesiology, McGovern Medical School, The University of Texas Health Science Center at Houston, Houston, TX, USA

Lee-Anne Slater, FRANZCR, CCINR Faculty of Medicine, Nursing and Health Sciences, Monash University, Melbourne, Australia

Interventional Neuroradiology Unit, Monash Imaging, Monash Health, Melbourne, Australia

Kevin Smith, PhD Kevin G. Smith, Ph.D. and Associates, P.C., Houston, TX, USA

Sadie E. Smith, MD Department of Anesthesiology and Perioperative Medicine, Penn State Health/Penn State College of Medicine, Hershey, PA, USA

Andrew So, MD Albert Einstein College of Medicine/Montefiore Medical Center, New York City, NY, USA

Lisa Stearns, MD Center for Pain and Supportive Care, Phoenix, AZ, USA

Roger Strachan, MD, FRCSE, FRCS(SN) Department of Neurosurgery, James Cook University Hospital, Middlesbrough, UK

Geoffrey Stricsek, MD Neurological Surgery, Thomas Jefferson University, Philadelphia, PA, USA

William A. Stuart, RPh Hartley Medical Center Pharmacy, Inc., Long Beach, CA, USA

Edward L. Swing, PhD Arizona Pain Specialists, Scottsdale, AZ, USA

Reda Tolba, MD Wake Forest Baptist Medical Center, Winston Salem, NC, USA

Hiep Tran, MD Department of Anesthesiology, Baylor Scott and White Health, Temple, TX, USA

Texas A&M University College of Medicine, Killeen, TX, USA

Katherine D. Travnicek, MD Pain Institute of Nevada, Las Vegas, NV, USA

Ned Urbiztondo, MD Spinal & Skeletal Pain Medicine, Utica, NY, USA

Elias Veizi, MD, PhD Case Western Reserve University, Cleveland, OH, USA

Louis Stokes VA Medical Center, Cleveland, OH, USA

Amy Wachholtz, PhD Department of Psychology, University of Colorado-Denver, Denver, CO, USA

Atul A. Walia, DO Integris Pain Management, University of Oklahoma Health Sciences Center, Oklahoma City, OK, USA

Torin Walters, MD Department of Diagnostic Radiology, Joan C. Edwards School of Medicine, Marshall University, Senior Partner Radiology Inc, Huntington, WV, USA

Dajie Wang, MD Chronic Pain Medicine, Department of Anesthesiology, Thomas Jefferson University Hospital, Philadelphia, PA, USA

James C. Watson, MD Departments of Neurology, Mayo Clinic, Rochester, MN, USA Departments of Anesthesiology, Pain Medicine, Mayo Clinic, Rochester, MN, USA

Brian H. Wetherington, MD Department of Anesthesiology and Pain Medicine, The University of Kansas Health System, Kansas City, KS, USA

Simon Willis, MD Department of Anesthesiology, University of Virginia School of Medicine, Charlottesville, VA, USA

Nicole Wolfgram, RN, FNP-BC Aspirus Hospital, Wausau, WI, USA

Kurt A. Yaeger, MD Department of Neurosurgery, Icahn School of Medicine, Mount Sinai Hospital, New York, NY, USA

Ajax Yang, MD Department of Rehabilitation Medicine, Icahn School of Medicine at Mount Sinai, New York, NY, USA

Part I Introduction

The Disease-Based Treatment of Pain

Andrew So and Karina Gritsenko

Introduction

The concept of pain has been an implicit variable in the equation of human life for many centuries. As one of the oldest universal phenomena, pain and its effects have asserted varying degrees of influence on religion, philosophy, and politics over time. Most notably, however, the phenomenon of pain has challenged physicians in their treatment planning and methodology; despite its ubiquitous existence, pain has never been universally understood due to its differing definitions across societies, beliefs, and contexts. Subsequently, medical professionals have been continuously tasked with redefining the phenomenon of pain.

Today, pain is one of the leading motivations for patients to seek physician treatment, as it is a component of many disease states; some would even argue that pain has evolved into a disease state in and of itself. This designation is the product of centuries of research that have resulted in various theories attempting to define pain as a complex biological process integrating the roles of anatomy, function, and external experience. With each new theory, the scope of pain's definition has expanded, and physicians have created new approaches to treatment.

As medical professionals' definition and understanding of the pain phenomenon has changed over time, treatment methods have evolved into multimodal approaches designed to attack the various components of this complex disease pathway. To better understand today's pain management techniques, it is important to familiarize oneself with the historical theories of pain, the current pain model, and the available treatments that have crafted the current approach.

A. So (🖂)

K. Gritsenko

Background and Historical Perspective

As it is regarded today, pain has not always been considered a biological phenomenon. In the prehistoric times, pain was thought to be a means of punishment that arose from an external experience of the spiritual world. In Ancient Chinese philosophy, pain was believed to be an imbalance between the vital forces of yin and yang, the complementary/opposing forces of life.

Early Greek philosophers also had their own postulations on the sensation of pain, as evident in the text of the *Iliad* by Homer (eight century to mid-fourth century BC), who used terms such as *penthos* and *algos* to describe different forms of physical and moral pain. The philosopher Aristotle designated the heart as the center of pain and sensation and believed that pain stemmed from the sensitivity of human organs.

Religion has also been used to explain the concept of pain, particularly in the Middle Ages. With the spread of Christianity, there seemed to be a preoccupation on the crucifixion of Christ and therefore a greater emphasis on bodily suffering. From this preoccupation emerged the idea that pain was a form of divine retribution, as demonstrated with martyred saints and the concept of purgatory.

The Shift to the Anatomical Basis of Pain

Over the centuries, the concept of pain slowly registered a paradigm shift from a philosophical and theological ideology to a more anatomical explanation. The works of Herophilus and Galen contributed to this pivotal shift. Herophilus studied the anatomy of the brain in the third century and laid early foundations in the understanding of the human nervous system and its anatomy. Approximately 1500 years later, the works of the physician Claudius Galen (130–201 AD) would suggest the existence of two sets of nerves with different roles: motor and sensation. Galen also demonstrated that the complete transection of the spinal cord caused sensory and



Albert Einstein College of Medicine/Montefiore Medical Center, New York City, NY, USA

Albert Einstein College of Medicine/Montefiore Medical Center, Montefiore Multidisciplinary Pain Program, The Bronx, NY, USA

T. R. Deer et al. (eds.), Deer's Treatment of Pain, https://doi.org/10.1007/978-3-030-12281-2_1

motor deficits and theorized that the spinal column was the connection between the brain and other organs. Galen's work demonstrated one of the earliest pathways that suggest pain required an organ to receive information from the outside world, a connecting passageway, and an organizing center to transform outside sensory information to perception.

During the Renaissance, philosopher Rene Descartes would describe a detailed somatosensory pathway. In his 1662 manuscript, *Treatise of Man*, Descartes builds on Galen's theories, describing the existence of nerves in the human body that convey both sensory and motor information. Descartes proposed the existence of gates between the brain and nerves, from which arouse the idea of mechanism; this theory compares the human body to a machine and describes pain as a consequence of damage to said hypothetical machine. Descartes' work showed that pain was an internal process organized by the human brain, rather than an external presence that entered the body, as was historically believed. In many ways, Descartes initiated the proposition of a human physiological process of pain.

In the years that followed these early anatomical studies, several important theories, which will be discussed in the next section of this chapter, would expand on these anatomical findings and be pivotal in constructing our modern-day knowledge of the pain pathway.

The Early Pain Theories

The current working definition of pain, as defined by the International Association for the Study of Pain, is "an unpleasant sensory and emotional experience associated with actual or potential tissue damage." The IASP's definition further describes pain as an integrative experience with both biological and psychosocial components. While there have been numerous theories proposed to explain pain, this chapter will focus on the most prevalent and influential theories of the past.

The Specificity Theory (1811)

The specificity theory was one of the first modern pain theories to suggest a separate anatomical neural pathway for pain perception. Maximilian von Frey postulated that the human body had five sensations (touch, heat, cold, pleasure, and pain), each with their own dedicated receptors and pathways to the brain. He theorized that noxious stimuli activated their own specialized receptors and were relayed by individualized pain fibers. Adding to the specificity theory, Charles Bell later suggested the idea of specialized spinal nerve roots with different functions. He hypothesized that the ventral nerve roots were responsible for motor and sensory activity in the dorsal nerve roots. From the specificity theory stems some of our modern ideas of the pain pathway – the ideas of specialized pain nociceptors, specialized nerves that carry varying pain signals (i.e., alpha-delta and C-nociceptive fibers), and a specialized spinothalamic tract in the spinal cord dedicated to nociception.

Pattern Theory (1894)

The Pattern theory, proposed by Alfred Goldscheider in 1894, suggests that the perception of pain is not the product of just one stimulation site transmitting to the brain, as proposed by the specificity theory, but rather an aggregate of multiple stimulation sites. It also postulates that the frequency of stimulation is responsible for variance in the intensity of pain. Goldscheider theorized that if a pain stimuli was applied repeatedly in higher frequency (temporal summation), it would appear more intense to the individual. Similarly, he suggested that pain could be spread over a large area and that such stimuli applied to a greater body area would convey a more intense pain (spatial summation).

Gate Control Theory of Pain (1965)

In 1965, Ronald Melzack and Charles Patrick would propose the gate control theory, which answered some of the limitations of its predecessor theories and ultimately revolutionized the field of pain medicine. The research behind the gate control theory of pain is rooted in the notion that pain signals are transmitted from the peripheral nervous system via small afferent pain fibers (alpha-delta and c fibers) to the dorsal horn of the spinal cord, which eventually leads to the brain. Along this pathway lie hypothetical "gates" that can be closed by varying mechanisms in order to inhibit the propagation of pain signals.

The gate control theory outlines two inhibitory pathways. First, it describes the ability of large nerve fibers (alpha-beta fibers), which carry non-noxious information and are located in the substantia gelatinosa of the spinal cord, to inhibit smaller afferent nerve fibers carrying nociceptive information. This proposed mechanism explains the theory behind the practice of rubbing or caressing a painful area of the body in order to alleviate pain. This idea is also the fundamental foundation behind the concepts of neuromodulation and transcutaneous electrical nerve stimulation (TENS).

The second inhibitory pathway is a descending inhibitory mechanism from the brain centered in the reticular formation. The theory states that heightened activity of this descending efferent pathway from the reticular system leads to an increased likelihood of gate closure and the subsequent inhibition of pain. This explains the theory behind hypnosis and why patients many times ignore the presence of painful stimuli while hypnotized.

Summarized, the overall perception of pain was said to be a balance between the two inhibitory mechanisms and the activity of the small noxious fibers. The gate control theory was important in showing the role of both the central and peripheral nervous systems in the perception of pain and was the first to explain the ability for non-noxious stimuli to modulate and change one's pain perception. As stated, it would also serve as the foundation of neuromodulation, a technique commonly used today for pain management.

Sensitization Models

Although the previous models laid the foundation for the neural pathway of pain, they failed to explain why pain could persist after the removal of the original stimuli, as demonstrated in chronic pain syndromes. The idea of sensitization (which includes both peripheral and central sensitization) would provide an answer to this phenomenon.

The term peripheral sensitization refers to the situation in which a nerve has increased excitability and therefore increased signal transmission. Peripheral sensitization can be applied to the pain pathway as commonly demonstrated following tissue injury, when inflammatory messengers including prostaglandins, cytokines, leukotrienes, and bradykinin are released. These inflammatory messengers cause changes in the voltage-gated sodium channels of neurons, which produce reduced activation potential and increased excitability. Subsequently, peripheral pain receptors are "sensitized" to have a lower threshold for firing, which results in increased response to noxious stimuli and can even result in a response to non-noxious stimuli.

Central sensitization is the idea that persistent pain transmission over time from the peripheral nervous system can cause the pain pathways in the spinal cord and brain to make new, stronger synapses (known as use-dependent plasticity). These new synapses cause a modulated pain transmission containing prolonged responses, or windup. Furthermore, the central sensitization model theorizes that increased activity from brain areas implicated in pain have the ability to modulate pain perception. Central sensitization can be used to explain hyperalgesia, allodynia, and spontaneous pain. There have been more detailed models proposed to explain its mechanism; the details behind these, however, are not the focus of this chapter.

Biopsychosocial Model

Although these historical theories appropriately suggested a biomedical mechanism to pain, the biopsychosocial model, which was proposed in 1977 by Engel, was the first to suggest that psychosocial factors can affect pain perception and maintenance of pain. This refuted the then-traditional view that pain was merely a sensation. Similarly, the Institute of Medicine in 2011 stated its belief that "pain arises in the nervous system, but represents a complex and evolving interplay of biological, behavioral, environmental, and social factors...."

There is not one singularly accepted scientific explanation behind psychology's role in the pain pathway, but many theories rest on the foundation that psychological factors may sensitize both peripheral and central nerve pathways, causing a heightened sensitivity to pain which ultimately affects the cognitive processing of the pain stimulus in the brain. The biopsychosocial model illustrates that pain is the product of a biomedical pathway (activation and transmission of noxious stimuli) and an integration of psychosocial influences, such as beliefs, cognition, mental state, and society.

This model can be used to explain why chronic pain patients often develop negative coping mechanisms (e.g., catastrophizing perceived helplessness and low selfefficacy), which ultimately influence pain severity and prolong a patient's symptoms. Similarly, depression has been found to be highly associated with chronic pain, with a prevalence rate between 30 and 54% and often serving as a prognostic indicator for greater disability. The biopsychosocial model draws further support from evidence that patients with chronic pain often benefit from cognitive- and behavioralbased interventions. Ultimately, the biopsychosocial model helped to evolve pain treatment approaches to more holistic ones, encompassing treatments for both the body and the mind.

Overview of the Current Pain Pathway

When considered in aggregate, these early pain theories provide substantial evidence in illustrating pain as a biological process with a dedicated pathway in the nervous system. This section will bring together these theories to outline the current scientific thinking of nociception and its mechanism, which is an important consideration in understanding the various treatment modalities of today. Future chapters will also provide more detailed information on the various components of the pain pathway.

The pain pathway starts with dedicated receptors and nociceptors, which respond to local tissue damage including mechanical (pressure and pinch), heat, and chemical. If an original stimulus exceeds the activation threshold, nociceptors convert the original stimulus into an electrical signal that is transmitted to the central nervous system for processing. When activated, these nociceptors produce a cascade of inflammatory mediators, which only further activate and sensitize them (a process known as peripheral sensitization). Examples of these mediators include prostaglandins, bradykinin, and histamine.

From here, the transduced electrical signal is carried via A-delta (A δ) and C fibers, which serve as the primary afferent nerve fibers in the pain pathway. Thinly myelinated $A\delta$ fibers are responsible for sharp and stinging pain sensations, whereas C fibers typically convey duller pain. These fibers carry signal to the spinal cord where they eventually terminate in the dorsal horn and synapse with second-order neurons. The signal then travels up to the brain along the spinothalamic and spinoreticular tracts, which are located in the anterolateral white matter of the spinal cord. Once in the brain, the signal is processed by the thalamus and projected to the somatosensory cortices, insula, anterior cingulate cortex, and prefrontal cortex, all of which are important for pain perception. The lateral system is thought to be responsible for analyzing pain location, duration, intensity, and quality, whereas the medial system is thought to be responsible for perception.

Throughout the pathway lie mechanisms that can inhibit the propagation of pain signal. Information about nonnoxious stimuli are carried by alpha-beta fibers and travel adjacent to the alpha-delta and C fibers; as postulated by the gate control theory, these non-noxious fibers may inhibit the fibers of nociception if activated simultaneously. Furthermore, the signal can be modulated by a variety of spinal mechanisms, which include endogenous opioids and cannabinoid systems, as well as inhibitory aminos such as GABA and nitric oxide. Lastly, the signal can be inhibited by descending pathways from the brain. These descending pathways are mediated by noradrenaline and serotonin; the two parts of the brain most important in the descending inhibition pathway are the periaqueductal gray and the nuclei raphe magnus.

Permanent lesions or persistent damage to different parts of the pain pathway can lead to central and peripheral sensitization. The modulation of the pathway via sensitization can also lead to many different types of pain, including chronic pain states and hyperalgesia.

Types of Pain

To better understand the treatment of pain as a disease state, one should first understand the various types of pain we know to exist today. With continued and increasingly advanced research within the field of pain medicine, physicians have developed more standardized methods in classifying the many different types of pain; one way to identify pain type is to analyze its characteristics of duration, anatomic source, and etiology. In 2002, the SCI Pain Task Force of the International Association of the Study of Pain developed a taxonomy to subdivide the various pain states, which is now one of the most common methods used to classify pain. This classification system uses a physiological/etiological approach, stratifying pains by the physiological mechanism behind each type based on etiology. In this method of pain classification, three types of pain can be identified. The first two types, known as nociceptive and neuropathic, carry distinguishing characteristics, while the third type (mixed) combines characteristics of both nociceptive and neuropathic pain.

Nociceptive Pain

Nociceptive pain describes pain that typically arises from the injury of somatic structures such as the skin, muscles, tendons, ligaments, bones, and joints. Nociceptive pain is the consequence of the transduction of noxious stimuli by A-delta and C fibers, as previously described in this chapter. Nociceptive pain can be further subdivided into somatic pain and visceral pain.

- A. Somatic pain is pain associated with peripheral tissue damage sensed by cutaneous nociceptors. Specialized nociceptors sense somatic pain in response to various harmful stimuli, most notably thermal (hot vs. cold), chemical (such as those produced by cuts and the inflammatory mediators produced by damaged tissues), and mechanical (e.g., stretching and pinching). Somatic pain can be thought of as a protective mechanism; it allows the body to localize tissue damage and withdraw itself from the source.
- B. While somatic pain is typically associated with external stimuli, visceral pain draws on information from internal organs (more specifically and most typically, the mucosal lining of hollow organs). In further contrast, somatic pain withdraws the body from potential harm, where visceral pain is the inner organs' defense mechanism to signal danger given their inability to elicit motor activity and move away from the pain signal. For example, the mucosal lining of the GI tract contains receptors with the ability to detect harmful changes such as dangerously high/low pH levels or the overstretching of the organ, which in turn produce symptoms of nausea and bowel distention and lead to defecation, vomiting, etc. Simplistically, visceral pain can be viewed as an attempt to maintain an internal homeostasis of the body.

Visceral pain is the product of vagal and spinal visceral afferent neurons. In contrast to somatic pain, there are no dedicated centers in the central nervous system specifically concerned with the transmission of visceral pain signals. This broadly explains why patients with visceral pain often have difficulty localizing the exact location of their pain. Furthermore, visceral pain can be felt in locations distant from the source of the stimulation, which is termed referred pain. The exact mechanism of referred pain is not known. One hypothesis suggests that sensory afferents supply multiple locations and that the stimulation of one location activates its other branches which in turn obscures the exact localization of the original stimulus.

Neuropathic Pain

Neuropathic pain is pain caused by damage of either the peripheral or central nervous system. The International Association of the Study of Pain formally defines neuropathic pain as "pain caused by a lesion or disease of the somatosensory nervous system." Central neuropathic pain is sourced from lesions within the spinal cord or brain. Some common disease states implicated in central neuropathic pain include spinal cord lesions, multiple sclerosis, transverse myelitis, and syringomyelia. In contrast, peripheral neuropathic pain is caused by lesions within the small unmyelinated C fibers and the myelinated A fibers. The peripheral neuropathies can be further subdivided based on distribution. Most peripheral neuropathies generally fall within two categories: peripheral focal/multifocal lesions and peripheral generalized lesions. Some examples of focal and multifocal lesions include entrapment syndromes, post-traumatic neuralgia, postherpetic neuralgia, and diabetic mononeuropathy; generalized lesions include polyneuropathies associated with diabetes mellitus, alcoholism, and HIV.

Research reveals a complex pathophysiology behind the etiology of neuropathic pain. In general, however, it can be stated that the peripheral nervous system responds to the presence of noxious stimuli when, in the absence of such stimuli, it would otherwise remain dormant. With lesion/ damage/chronic disease, these nerves are sensitized (as detailed previously) and develop pathological baseline activity that causes neuropathic pain, even in the absence of noxious stimuli.

Acute Versus Chronic Pain

Pain can also be categorized by duration; these designations include acute and chronic pain. Acute pain is normally described as pain that extends for less than 3 days. It is the consequence of a specific disease or injury and is normally self-limited. For example, patients often experience a degree of pain in the days immediately following surgery, but this pain tends to subside thereafter with the passage of a moderate amount of time. Conversely, chronic pain is pain that persists beyond the expected healing period for a specific injury or disease; typically pain greater than 3 months is defined as chronic pain. Chronic pain is hypothesized to be the result of permanent changes within the nervous system (central and

peripheral sensitization, as discussed previously), and the associated syndromes of chronic pain are typically the focus of pain management and treatment. Some examples include chronic lower back pain, headache, myofascial pain, and fibromyalgia. These chronic pain syndromes comprise components of both neuropathic and nociceptive pain and are marked by distinguishing biological etiologies.

Treatment: A Multimodal Approach

Today's approach to pain treatment is the result of many decades of research into the effectiveness and efficiencies of treatment methods and reflects the depth of the patient pool currently seeking treatment. It has been estimated that roughly 100 million adults suffer from chronic pain and that chronic pain is responsible for 90 million physician visits, 14% of all prescriptions, and 50 million lost workdays annually. The total healthcare cost associated with pain treatment in the United States is estimated to be \$650 billion dollars per year. With such a prevalent burden on both society and medicine, the American Board for Hospital Accreditation has adopted pain as the "fifth vital sign." Similarly, Canada added pain assessment and management to its Achieving Improved Measurement Standards in 2005. Continued increases in the number of patients seeking treatment for pain have led to the formal protocolization of pain assessment and treatment in the inpatient setting.

The task of developing efficacious and cost-effective treatment methods has challenged physicians and scientists for decades. A well-received and widely utilized solution was developed by anesthesiologist John J. Bonica, who pioneered the multidisciplinary approach to pain treatment. As a physician who treated chronic pain in soldiers after World War II, Bonica was one of the first physicians to consult experts from different disciplines (neurology, psychiatry, surgery) in an attempt to find more effective and allencompassing treatments for his patients. Utilizing this multidisciplinary approach, Dr. Bonica witnessed vast improvements in his patients' pain ratings and led him to open the first multidisciplinary pain clinic in 1950. This multidisciplinary approach to treatment produced greater patient satisfaction and decreased healthcare costs.

Since the landmark events of Dr. Bonica's multidisciplinary approach to treatment, physicians have accepted the multifaceted mechanism involved in the generation, adaptation, and chronicity of pain, as illustrated throughout this chapter. With this acceptance, pain management has evolved into a multimodal approach, and pain treatment now often encompasses a combination of pharmaceutical, surgical, behavioral, and alternative treatments. Utilizing a multidisciplinary approach, the goal of pain management today is not only to reduce a patient's pain symptoms but to improve the individual's overall physical function, better manage chronic diseases, and improve psychological well-being in relation to a patient's pain in the hopes of creating an improved overall general state.

This section of the chapter serves as an introduction to the current modalities of pain treatment and the diseases they treat. Pain treatment can be simplified into two broad categories: pharmacological and nonpharmacological strategies. Future chapters will be dedicated to more detailed descriptions and analyses of each of the modalities.

Pharmacological Treatments

Pharmacological treatment is one of the oldest methods of pain treatment that is still widely used today. Specific pharmacological algorithms, which will be discussed in a future chapter, have been proposed to aid practitioners in treating pain. When implementing pharmacological strategies, it is important to consider the type of pain that is being treated, whether it be nociceptive, neuropathic, or mixed; specific classes of medication treat different types of pain with varying efficacies. Furthermore, a multimodal approach to pharmacological treatment is considered most beneficial for patients. The aim of a multimodal approach is to use medications that act on different receptors and/or parts of the pain pathway to create synergistic effects and consequently stronger pain relief. Some of the most commonly utilized pharmacological treatments include nonsteroidal anti-inflammatory drugs (NSAIDS), antidepressants, anticonvulsant medications, and opioids.

Nonsteroidal Anti-inflammatory Drugs (NSAIDS)

Nonsteroidal anti-inflammatory drugs (NSAIDS) are one of the most popular agents in the treatment of multiple pain states, most notably inflammatory pain. NSAIDS provide pain relief via anti-inflammatory mechanisms by inhibiting prostaglandins-producing enzymes, which are present in both the central and peripheral parts of the pain pathway. Prostaglandins are produced by the COX class of enzymes, with COX-1 and COX-2 being the most pertinent isoforms to the pain pathway [62]. Although NSAIDS inhibit both COX-1 and Cox-2, it has been suggested that most of its analgesic properties arise from its COX-2 inhibition. Nonselective COX agents inhibit both COX-1 and Cox-2 isoforms; some examples of nonselective inhibitors include aspirin, paracetamol, ibuprofen, and naproxen. Selective COX inhibitors, or those that only inhibit COX-2, were later designed to prevent the common side effects associated with nonselective agents (most notably GI intolerance and renal injury). Examples of selective COX inhibitors include celecoxib, meloxicam, and rofecoxib. It is important to note that NSAIDS, although frequently used in the treatment of neuropathic pain, have been shown to have little utility in the

treatment of nociceptive pain. NSAIDS have been found most effective in treating diseases with an inflammatory mechanism such as osteoarthritis and rheumatoid arthritis.

Antidepressants

Antidepressants, although originally intended for the treatment of psychiatric illnesses, have also been found effective in pain treatment. Antidepressant medications work by inhibiting the reuptake of certain neurotransmitters, most notably serotonin and norepinephrine, causing a buildup that potentiates inhibitory pain pathways in the spinal cord.

The three classes of antidepressants most commonly utilized in pain management are tricyclic antidepressants (TCAs), selective serotonin reuptake inhibitors (SSRIs), and serotonin-norepinephrine reuptake inhibitors (SNRIs). Tricyclic antidepressants were the first class to be used in pain medicine and work by blocking the reuptake of serotonin and noradrenaline. TCAs have been found beneficial in the treatment of neuropathic pain, fibromyalgia, and headaches. However, notable side effects of TCAs include cardiac arrhythmia and hypertension.

Selective serotonin reuptake inhibitors, unlike tricyclic antidepressants, only inhibit the reuptake of serotonin and were developed in an attempt to avoid the side effects of TCAs. There has also been increased research on newer SNRI drugs, which inhibit the reuptake of serotonin and noradrenaline, showing their helpfulness in treatment of neuropathic pain and fibromyalgia.

Anticonvulsant Medications

Anticonvulsant medications have been effective in the field of chronic pain management since their inception in the due 1960s to their neuromodulating properties. Anticonvulsants assert their activity by enhancing the inhibitory pathway of the gamma-aminobutyric acid (GABA) pathway; gabapentin and pregabalin are the most researched and most commonly utilized anticonvulsants in the field of pain management. These drugs have become clinically favored as adjunct agents in the treatment of chronic neuropathic pain, particularly in postherpetic neuralgia and diabetic neuropathy. Carbamazepine, also in this class of drugs, is considered a mainstay in the treatment of chronic neuropathic pain associated with trigeminal neuralgia. The popularity of anticonvulsants has risen over time because of their wide safety margin. The most common side effect of anticonvulsants is dose-dependent sedation.

Opioids

Opioids are one of the oldest, most potent, and most popular drug classes used in pain management and treatment. Opiates is the term used to encompass naturally occurring derivatives of opium such as morphine and codeine; opioids is a broader term used to describe any drug that produces morphine-like actions, whether they be naturally occurring (opiates) or artificially synthesized (as with fentanyl, heroin, and methadone). The effectiveness of these drugs in treating pain, particularly acute pain within the first 6 weeks of an acute pain trigger, is undisputed, but there is currently no proven data to show efficacy beyond the acute pain period. In addition, adverse effects have been a significant source of scrutiny in the medical field; due to the current rise in opioid misuse, addiction, and overprescribing, there has been a sway in the pendulum of pain treatment to minimize the use of opioids at this time.

Opioids exert their desired effects via G-coupled protein receptors, which are subclassified as mu, kappa, and delta receptors. Simplistically, opioid receptors inhibit adenylyl cyclase, which subsequently inhibit cyclic AMP (cAMP) and the release of certain neurotransmitters implicit in pain production and perception. Opioids for chronic pain treatment are usually prepared in either oral or transdermal forms.

Although many would accept their role in the management of acute pain, opioids have been highly scrutinized in the context of chronic pain management, predominantly due to their adverse side effects and potentially abusive qualities. The primary side effects of opioids are reported to be constipation, nausea, and sedation. More serious side effects, such as hypogonadism, amenorrhea in women, and impairment in neuropsychological function, have also been associated with chronic opioid use. Furthermore, opioids have been shown to have addictive properties, and the concepts of drug addiction, tolerance, and dependence must be addressed with patients when administered for prolonged use in the treatment of chronic pain.

Interventional Techniques

Although pharmacological and noninvasive techniques are typically the first and primary line of defense in the treatment of chronic pain, it has been estimated that approximately 10% of patients cannot be adequately treated with these methods. In these select patients, the role of interventional techniques has been explored to find potential solutions in the treatment and management of their pain. Interventional techniques are usually utilized when a patient's pain arises from a spinal etiology, such as discogenic or sacroiliac joint pain. Interventional techniques have been used to both help alleviate pain, as well as to locate and diagnose the source of a patient's pain. Common interventional techniques include injection therapy, intrathecal drug delivery, and discography.

Injection Therapy

Injection therapy encompasses a wide array of injection types used in the treatment of chronic pain and were first reported to be utilized in 1901. With the development of modern techniques, injection therapy has become an accepted tool in treating pain patients.

Peripheral injection therapies include trigger point and intra-articular joint injections. Trigger point injections involve dry needling into tender muscle points, which attempts to elicit a twitch in the targeted muscle and subsequently alleviate any painful preexisting muscle contractures. The dry needling can also be supplemented with concomitant injections of local anesthetics, though this has not been scientifically proven effective. Anecdotally, physicians have witnessed the benefits of trigger point injections in the treatment of myofascial pain, while intra-articular joint injections have been utilized in the treatment of chronic joint pain. These injections are performed with corticosteroids, which are aimed at reducing inflammation in the joint.

Central injection therapies include intrathecal therapy, epidural injections, and radiofrequency ablation. In the United States, epidural steroid injections are the most common intervention form of this type utilized in pain treatment. Epidural steroid injections achieve their pain-alleviating properties by reducing inflammation of nerve roots and their surrounding environment and have been shown to be useful in the treatment of radiculopathies caused by disc pathology such as disc herniation. Traditionally, epidural steroid injections had been administered via the interlaminar approach, where the injection is placed in the space between two laminae. Other approaches, such as caudal and transforaminal, have also become more common with the advancement of technique and the advent of fluoroscopic-guided practices.

Radiofrequency ablation (RFA) is the process by which a generated radio frequency wave is injected via an insulated needle to produce heat energy from the induction of ion oscillation. The heat energy produced subsequently causes the thermal ablation of the surrounding environment. In relation to pain treatment, RFA has been targeted at dorsal root ganglions and nerve plexus in various diseases in an attempt to cause nerve denervation and block pain signal propagation. RFA has been shown helpful in the treatment of trigeminal neuralgia (gasserian ganglion), cluster headaches (sphenopalatine ganglion), and postherpetic neuralgia (percutaneous dorsal root ganglions). The most common use of radiofrequency ablation has been witnessed in the treatment of facet-mediated back pain, given RFAs particular effectiveness in its ability to denervate medial branch nerves.

Neuromodulation

Neuromodulation is one of the newest emerging treatment concepts in the field of pain management. Although largely still considered an evolving treatment, the foundation of neuromodulation originated in the mid-1960s with Melzack and Wall's gate control theory of pain, which states that the transmission of pain depends upon the opening and closing of a "gate" by large and small fibers. Neuromodulation functions under the assumption that it may be possible to alter the activation of these large and small fibers in different parts of the pain pathway, resulting in the inhibition pain. The alteration of these fibers can be achieved via two forms: electrical and chemical neuromodulation.

Electrical neuromodulation is accomplished by placing implantable electrodes, attached to a generator, in the peripheral and central nervous system to generate a pulse onto the targeted nerves. Different types of electrical neuromodulation include spinal cord stimulation (SCS), peripheral nerve stimulation (PNS), occipital nerve field stimulation (ONFS), deep brain stimulation (DBS), and motor cortex stimulation (MCS).

Spinal cord stimulation has been most frequently implemented in the treatment of persistent back pain following failed back surgery. Spinal cord stimulation has also been used in the treatment of ischemic leg pain from peripheral vascular disease, unstable angina, and complex regional syndrome. Deep brain stimulation, similar to spinal cord stimulation, asserts its pain-alleviating effects through the electrical stimulation of subcortical structures. Although not completely understood, the stimulation of the periaqueductal and periventricular gray matter of the thalamus may cause the release of endogenous opiates that provide pain relief. Motor cortex stimulation has been indicated in the treatment of central post-stroke pain.

Neuromodulation can also be accomplished through chemical means by delivering drugs directly into the central nervous system. Chemical neuromodulation is most commonly accomplished via intrathecal drug delivery systems.

Psychological Treatment of Pain

As defined by the biopsychosocial model, pain is not solely a biological mechanism but can also be influenced by an individual's environment and mental well-being. Pain can often produce anxiety in patients and has been shown to ultimately worsen the overall perception of pain. Thus, it is not surprising that today's treatment protocols often include components aimed at assessing and treating the psychological impacts of pain.

The most common technique used to treat the psychological effects of pain is cognitive behavioral therapy (CBT). Cognitive behavioral therapy identifies the negative perceptions, emotions, and beliefs an individual associates with his or her physical pain, which often cause increased stress and anxiety. Furthermore, CBT seeks to identify any negative coping mechanisms individuals have learned or developed to cope with pain. Once identified, CBT attempts to equip patients with new techniques and positive tools to cope, adapt, and manage their pain independently, counteracting the previously held negative beliefs and ineffective coping

Conclusion: Putting It All Together

The concept of pain remains a medical phenomenon. Today's definition of pain is complex and continues to evolve with new scientific discoveries and the overall advancement of medicine. Given the dynamic nature of pain, its definition is likely to continue to change over time; it remains well established, however, that pain is an integrative process comprising an interplay between the human anatomy/physiology, psychological factors, and external environment. Under the current definition, it is also widely accepted that many types of pain and a multitude of pain states exist. With such diversity among the mechanics and types of pain, the field of pain management has evolved to encompass multimodal and multidisciplinary approaches to treatment. As physicians continue to seek a full understanding of pain in an effort to better serve patients, it is expected that treatments will continue to evolve and new modalities will be pioneered. Nevertheless, the historical context provided in this chapter serves as the foundation of today's treatment methods and provides a strong base for future developments in the field of pain management.

Recommended Reading

- 1. Anwar K. Pathophysiology of pain. Dis Mon. 2016;16(9):324-9.
- Ashburn M, Staats P. Management of chronic pain. Lancet. 1999;353(9167):1865–9.
- Baron R. Mechanisms of disease: neuropathic pain—a clinical perspective. Nat Clin Pract Neurol. 2006;2(2):95–106.
- Bashir U, Colvin LA. The place of pharmacological treatment in chronic pain. Anaesth Intensive Care Med. 2013;14(12):528–32.
- Beissner F, Brandau A, Henke C, et al. Quick discrimination of A delta and C fiber mediated pain based on three verbal descriptors. Ikeda K, ed. PLoS One. 2010;5(9):e12944.
- Binshtok AM. Mechanisms of nociceptive transduction and transmission: a machinery for pain sensation and tools for selective analgesia. Int Rev Neurobiol. 2011;97:143–77.
- Bittar RG, Teddy PJ. Peripheral neuromodulation for pain. J Clin Neurosci. 2009;16(10):1259–61.
- Braz J, Solorzano C, Wang X, Basbaum AI. Transmitting pain and itch messages: a contemporary view of the spinal cord circuits that generate gate control. Neuron. 2014;82(3):522–36.
- 9. Buyten JP. Radiofrequency or neuromodulation treatment of chronic pain, when is it useful? Eur J Pain Suppl. 2008;2(S1):57–66.
- Campbell LC, Clauw DJ, Keefe FJ. Persistent pain and depression: a biopsychosocial perspective. Biol Psychiatry. 2003;54(3):399–409.
- Chapman CR. New directions in the understanding and management of pain. Soc Sci Med. 1984;19(12):1261–77.
- Cheatle MD. Biopsychosocial approach to assessing and managing patients with chronic pain. Med Clin N Am. 2016;100(1):43–53.
- 13. Chen J. History of pain theories. Neurosci Bull. 2011;27:343.

- Chen M, Hoshino H, Saito S, Yang Y, Obata H. Spinal dopaminergic involvement in the antihyperalgesic effect of antidepressants in a rat model of neuropathic pain. Neurosci Lett. 2017;649:116–23.
- Cohen S, Raja S. Goldman's Cecil medicine. 24th ed. Philadelphia: Elsevier; 2012. Chapter 29, Pain. p. 133–40.
- Colloca L, Ludman T, Bouhassira D, Baron R, Dickenson A, et al. Neuropathic pain. Nat Rev Dis Primers. 2017;3:17002.
- Craig AD. Pain mechanisms: labeled lines versus convergence in central processing. Annu Rev Neurosci. 2003;26(1):1–30.
- Dale R, Stacey B. Multimodal treatment of chronic pain. Med Clin N Am. 2016;100(1):55–64.
- Dedeli O, Kaptan G. Spirituality and religion in pain and pain management. Health Psychol Res. 2013;1(3):e29.
- Deer TR. An overview of interventional spinal techniques. Semin Pain Med. 2004;2(3):154–66.
- Dietrich J. Psychology and chronic pain. Anaesth Intensive Care Med. 2011 Feb;12(2):42–3.
- D'Mello R, Dickenson AH. Spinal cord mechanisms of pain. Br J Anaesth. 2008;101(1):8–16.
- 23. Doleys DM. Chronic pain as a hypothetical construct: a practical and philosophical consideration. Front Psychol. 2017;8:664.
- Dowell D, Haegerich TM, Chou R. CDC guideline for prescribing opioids for chronic pain – United States, 2016. MMWR Recomm Rep. 2016;65(RR-1):1–49.
- Duan B, Cheng L, Ma Q. Spinal circuits transmitting mechanical pain and itch. Neurosci Bull. 2018;34(1):186–93.
- Dubin AE, Patapoutian A. Nociceptors: the sensors of the pain pathway. J Clin Invest. 2010;120(11):3760–72.
- Dworkin RH, O'Connor AB, Audette J, et al. Recommendations for the pharmacological management of neuropathic pain: an overview and literature update. Mayo Clin Proc. 2010;85(3 Suppl):S3–S14.
- Finnerup NB, Haroutounian S, Kamerman P, Baron R, Bennett DL, Bouhassira D, et al. Neuropathic pain. Pain. 2016;157(8):1599–606.
- Grichnik KP, Ferrante FM. The difference between acute and chronic pain. Mt Sinai J Med. 1991;58(3):217–20.
- Hadidi S, Baranidharan G. Implantable technology for pain management. Anaesth Intensive Care Med. 2016;17(11):536–42.
- Hamann MCJ, Tsai EC, Tator CH, Shoichet MS. Novel intrathecal delivery system for treatment of spinal cord injury. Exp Neurol. 2003;182(2):300–9.
- 32. Handwerker H. Itch hypotheses. Itch Front Neurosci. 2014:1-8.
- High KNW. Pain pathways: peripheral, spinal, ascending, and descending pathways. In: Rajs practical management of pain. Elsevier Inc. 2008. p. 119–34. https://doi.org/10.1016/B978-0-323-04184-3.X5001-8.
- Hillinger MG, Wolever RQ, Mckernan LC, Elam R. Integrative medicine for the treatment of persistent pain. Prim Care. 2017;44(2):247–64.
- 35. Hoffert M. The gate control theory re-revisited. J Pain Symptom Manag. 1986;1(1):39–41.
- Honey CM, Tronnier VM, Honey CR. Deep brain stimulation versus motor cortex stimulation for neuropathic pain: a minireview of the literature and proposal for future research. Comput Struct Biotechnol J. 2016;14:234–7.
- Kansal A. Visceral pain. Anaesth Intensive Care Med. 2016;17(11):543–7.
- Keele KD. Anatomies of pain. Oxford: Pl. XIII. Blackwell Scientific Publications; 1957.
- Kremer MCA, Salvat E, Muller A, Yalcin I, Barrot M. Antidepressants and gabapentinoids in neuropathic pain: mechanistic insights. Neuroscience. 2016;338:183–206.
- Kroenke K, Krebs EE, Bair MJ. Pharmacotherapy of chronic pain: a synthesis of recommendations from systematic reviews. Gen Hosp Psychiatry. 2009;31(3):206–19.
- 41. Lima D. Endogenous pain modulatory system in the light of the gate control theory. Pain Forum. 1996;5(1):31–9.

- Lynch ME, Watson CPN. The pharmacotherapy of chronic pain: a review. Pain Res Manag. 2006;11(1):11–38.
- Maixner W, Fillingim RB, Williams DA, Smith SB, Slade GD. Overlapping chronic pain conditions: implications for diagnosis and classification. J Pain. 2016;17(9):T93–107.
- Manworren RC. Multimodal pain management and the future of a personalized medicine approach to pain. AORN J. 2015;101(3):307–18.
- 45. Marchand S. The phenomenon of pain. Seattle: IASP Press; 2012.
- Mcdonald J, Lambert DG. Opioid mechanisms and opioid drugs. Anaesth Intensive Care Med. 2016;17(9):464–8.
- 47. Melzack R. From the gate to the neuromatrix. Pain. 1999;82:S121-6.
- Melzack R, Wall PD. Pain mechanisms: a new theory. Pain Forum. 1996;5(1):3–11.
- Mendell LM. Constructing and deconstructing the gate theory of pain. Pain. 2014;155(2):210–6.
- Mendis V, Mottaleb R, Fung M. Radiofrequency techniques in pain management. Anaesth Intensive Care Med. 2016;17(11):564–7.
- Moayedi M, Davis KD. Theories of pain: from specificity to gate control. J Neurophysiol. 2012;109(1):5–12.
- Nasmerow L, Kutner E, Wakefiled E, Rzepski B, Sahl R. Pain amplification syndrome: a biopsychosocial approach. Semin Pediatr Neurol. 2016;23(3):224–30.
- 53. Nijs J, Wilgen CPV, Oosterwijck JV, Ittersum MV, Meeus M. How to explain central sensitization to patients with 'unexplained' chronic musculoskeletal pain: practice guidelines. Man Ther. 2011;16(5):413–8.
- Ochs S-A. History of nerve functions: from animal spirits to molecular mechanisms. Cambridge: Cambridge University Press; 2004.
- Ossipov MH, Dussor GO, Porreca F. Central modulation of pain. J Clin Invest. 2010;120(11):3779–87.
- Pearce JMS. Von Frey's pain spots. J Neurol Neurosurg Psychiatry. 2006;77(12):1317.
- Penney JN. The biopsychosocial model of pain and contemporary osteopathic practice. Int J Osteopathic Med. 2010;13(2):42–7.
- Pergolizzi J, Ahlbeck K, Aldington D, Alon E, Coluzzi F, Dahan A, et al. The development of chronic pain: physiological CHANGE necessitates a multidisciplinary approach to treatment. Curr Med Res Opin. 2013;29(9):1127–35.
- Perl ER. Pain mechanisms: a commentary on concepts and issues. Prog Neurobiol. 2011;94(1):20–38.
- Pleuvry B. Non-opioid analgesics. Anaesth Intensive Care Med. 2005;6(1):25–9.
- 61. Rey R. The history of pain. Cambridge, MA: Harvard Univ. Press; 1998.
- Roe M, Sehgal A. Pharmacology in the management of chronic pain. Anaesth Intensive Care Med. 2016;17(11):548–51.
- Sanchis MN, Lluch E, Nijs J, Struyf F, Kangasperko M. The role of central sensitization in shoulder pain: a systematic literature review. Semin Arthritis Rheum. 2015;44(6):710–6.
- 64. Serpell MG, Makin A, Harvey A. Acute pain physiology and pharmacological targets: the present and future. Acute Pain. 1998;1(3):31–47.
- Seth B, Gray LD. Genesis of chronic pain. Anaesth Intensive Care Med. 2016;17(9):431–5.
- 66. Sousa MVPD, Ferraresi C, Magalhães ACD, Yoshimura EM, Hamblin MR. Building, testing and validating a set of home-made von Frey filaments: a precise, accurate and cost effective alternative for nociception assessment. J Neurosci Methods. 2014;232:1–5.
- Stanos S, Tyburski M, Harden NR. Braddom's physical medicine and rehabilitation. 5th ed. Philadelphia: Elsevier; 2015. Chapter 37, Chronic Pain. p. 809–17.
- Steeds C. The anatomy and physiology of pain. Surgery Oxford International Edition. 2009;27(12):507–11.
- Stein C. Opioids, sensory systems and chronic pain. Eur J Pharmacol. 2013;716(1–3):179–87.

- Tashani O, Johnson M. Transcutaneous Electrical Nerve Stimulation (TENS) a possible aid for pain relief in developing countries? Libyan J Med. 2009;4(2):62–5.
- Taylor LEV, Stotts NA, Humphreys J, Treadwell MJ, Miaskowski C. A biopsychosocial-spiritual model of chronic pain in adults with sickle cell disease. Pain Manag Nurs. 2013;14(4):287–301.
- 72. Tompkins DA, Hobelmann JG, Compton P. Providing chronic pain management in the "Fifth Vital Sign" Era: historical and treatment perspectives on a modern-day medical dilemma. Drug Alcohol Depend. 2017;173:S11–21.
- Turk DC, Wilson HD, Cahana A. Treatment of chronic non-cancer pain. Lancet. 2011;377(9784):2226–35.
- 74. Wolff BB. Gate control theory and the brain. Pain Forum. 1996;5(2):147–9.
- 75. Woolf CJ. Central sensitization: implications for the diagnosis and treatment of pain. Pain. 2011;152(3 Suppl):S2–15.
- Wrigley P, Siddall P. Wall & Melzack's textbook of pain. 6th ed. Philadelphia: Elsevier/Saunders; c2013. Chapter 68, Pain following Spinal Cord Injury. p. 978–89.

Algorithms of Pain Treatment

Andrew So and Karina Gritsenko

Introduction

Chronic pain is one of the most common symptoms for which patients seek medical attention. It is estimated that up to 11.2% of adults in the United States suffer from chronic pain [1] and that patients in the United States alone spend between \$560 billion and \$635 billion on chronic pain treatment annually [2]. Chronic pain also remains widely prevalent due to its ubiquitous nature; it is associated with a plethora of conditions, such as cancer, lower back pain, and neuropathic pain conditions [3].

At present, there is not one universally accepted formula to treat chronic pain, and many times treatment is based on anecdotal outcomes and non-evidence-based personal experiences [4]. Dr. John Bonica, who was a trained anesthesiologist during World War II, was one of the first physicians to formalize the treatment of chronic pain, leading to the eventual transformation of chronic pain into its own subspecialty. Dr. Bonica was also one of the first physicians to treat pain in a multidisciplinary way in an attempt to formalize a more effective approach to pain treatment [5]. Since then, numerous societies have been established that are dedicated to the fields of pain treatment and management.

Even with the advent of a multidisciplinary approach by Dr. Bonica, it remains impossible to create a single set of criteria to treat the wide array of pain states and disease mechanisms. Various tools and guidelines have been set forward to aid physicians in their treatment approaches and to streamline a more protocolized approach to the treatment of pain. As such, pain physicians should understand the current

A. So (🖂)

K. Gritsenko

approach to treating chronic pain patients and the tools that are available to aid in this journey.

To provide a better understanding of the common algorithms used in chronic pain treatment, this chapter will discuss the importance of the initial patient evaluation in the treatment process and highlight several effective tools physicians can implement. It will also highlight the currently accepted treatment guidelines for some pain states most commonly encountered by physicians.

Assessment Algorithm

Although treatment algorithms typically vary depending upon the type of pain being treated (cancer, neuropathic, lower back pain, etc.), a macro-level analysis of pain algorithms reveals strong underlying points of commonality, including the importance of the initial evaluation of a chronic pain patient. Therefore, we begin our analysis of pain algorithms by exploring the initial process of patient assessment and its role in framing overall treatment plans.

A comprehensive initial assessment provides vital information on an individual's pain and aids in treatment decisionmaking that will provide the best suited outcome for individual patients [6]. In formulating their Pain Taxonomy framework, the American Pain Society defines the importance of a reliable and valid pain assessment in the treatment of chronic pain patients; this initial assessment functions as a vital tool in helping determine the severity of pain, surveying the course of the pain, and establishing the pathophysiology of the pain. With this information, physicians are equipped with a better understanding of the patient and thus are more ably prepared to craft effective treatment plans [7]. There is a myriad of literature focused in detail on the specific domains that should be investigated in the initial pain assessment, but given the purpose and scope of this chapter, this section will focus on the most highlighted domains. Each of these domains is discussed in brief below.



Albert Einstein College of Medicine/Montefiore Medical Center, New York City, NY, USA

Albert Einstein College of Medicine/Montefiore Medical Center, Montefiore Multidisciplinary Pain Program, The Bronx, NY, USA

T. R. Deer et al. (eds.), Deer's Treatment of Pain, https://doi.org/10.1007/978-3-030-12281-2_2

History

Nearly all pain treatment algorithms emphasize the importance of an initial history review. Gathering and understanding a detailed, holistic background on a patient and his/her medical history often supply practitioners with significant information that eventually aids in treatment planning. The result of this initial interview can guide a practitioner in making medical diagnoses, determining a need for further workup, and/or assessing potential treatment options [6].

In their most recently published Practice Guidelines for Chronic Pain Management, the American Society of Anesthesiologists (ASA) recommends that the initial evaluation of a chronic pain patient includes a detailed medical history with specific focus on the chronology of the pain. Attention should be paid to obtaining details such as the intensity, quality, location, duration, and distribution of a patient's reported pain. In addition to symptom history, social history (with a focus on abuse), surgical history, and family history should also be investigated and recorded. Furthermore, the ASA recommends a review of previous diagnostic tests, medication lists (both past and present), and any other previously implemented treatments to investigate their historical effectiveness [8].

Physical Exam

A thorough physical exam is also a fundamental component of the initial evaluation. The aim of the physical exam is to elicit information to help make a proper diagnosis and may also help eliminate underlying and treatable pathology [9]. In general, a physical exam should follow that of a normal generalized comprehensive human systems exam. Particular attention should be given to the system under which the initial chief complaint falls. Furthermore, special attention should be paid to the musculoskeletal and neuromuscular system, as the majority of chronic pain falls under this system [10]. Eliciting symptoms, such as focal weaknesses, site tenderness, and neurological deficits may help providers make accurate diagnosis.

Pain Intensity: Pain Intensity Rating Scales

As covered, one purpose of the initial interview is to determine the intensity of a patient's pain, which some argue to be the most important dimension investigated in the initial pain assessment [11]. With an understanding of pain intensity, clinicians are better prepared to gauge the severity of a patient's pain and its realistic impact/burden on the patient's functional capacity. Furthermore, clinicians can make more informed treatment decisions based on a patient's pain intensity and, even more importantly, assess the efficacy of a treatment by evaluating positive/neutral/negative changes in reported intensity [12].

An objective and accurate system of measuring pain intensity remains difficult to define, predominantly due to the multitude of factors that can influence a patient's perception of pain, from pathophysiological to psychosocial aspects [13]. Given its subjective nature and with only a limited amount of time for clinicians to ascertain a patient's pain intensity, the role of pain measurement scales has proven to be essential. The verbal rating scale (VRS), the visual analog scale (VAS), and the numerical rating scale (NRS) are among the most commonly used pain measurement scales in clinical practice. While all of these scales have proven to be helpful, distinct advantages and disadvantages can be identified for each [14].

The verbal rating scale has been implemented in practice since the 1940s and consists of a list of qualitative descriptors used to define various levels of pain intensity. An example of a verbal rating scale would be no pain, mild pain, moderate pain, and severe pain. Each adjective is also assigned a numeric value in an ascending order of pain intensities; in the previous example, no pain = 0, mild pain = 1, and so forth [15]. Patients are asked to select the term from the rating scale which best correlates to their perceived pain intensity, and the numeric value associated with that adjective is then assigned. A major advantage of the verbal rating scale is its relative simplicity and easy implementation [13]. Some criticize the verbal rating scale since patients may have to be familiar with the terms to properly use it.

The visual analog scale is a linear model anchored at one end by the non-existence of pain symptoms and opposed at the other by the worst imaginable pain. A numerical value is also plotted on the line to coincide with the various pain intensity descriptors, with 0 equating to no pain and 10 with the worst imaginable pain [16]. Patients are asked to mark where on the line their pain intensity lies. The biggest advantage of the VAS noted is its high degree of "sensitivity" when compared to the other scale rating systems [17]. However, the VAS has been characterized as more complex and may be difficult to implement for patients with cognitive impairments.

The numeric rating scale consists of a numerical scale, usually 0 to 10 (0 = no pain and 10 = most severe pain ever). Patients are instructed to pick a number that best reflects their pain intensity. The numeric rating scale has been seen to be easier to use than the visual analog scale [7].

Pain Quality: The McGill Pain Questionnaire

Along with the varying pain intensities discussed previously, pain can also be characterized by its perceived qualities. Even though two varying pain symptoms may be ranked as