

National Strategy For Palliative Care

Catherine Walshe, Nancy Preston, Bridget Johnston

Oxford Textbook of Palliative Medicine Nathan I. Cherny, Marie Fallon, Stein Kaasa, Russell K. Portenoy, David Currow, 2015 Emphasising the multi-disciplinary nature of palliative care the fourth edition of this text also looks at the individual professional roles that contribute to the best-quality palliative care.

Textbook of Palliative Care Roderick Duncan MacLeod, Lieve Van den Block, 2025-02-25 This second edition provides the most up-to-date information on all aspects of palliative care including recent developments (including COVID-19), global policies, service provision, symptom management, professional aspects, organization of services, palliative care for specific populations, palliative care emergencies, ethical issues in palliative care, research in palliative care, public health approaches and financial aspects of care. This new Textbook of Palliative Care remains a unique, comprehensive, clinically relevant and state-of-the art book, aimed at advancing palliative care as a science, a clinical practice and as an art. Palliative care has been part of healthcare for over fifty years but it still needs to be explained to many. Healthcare education and training has been slow to recognize the vital importance of ensuring that all practitioners have a good understanding of what is involved in the care of people with serious or advanced illnesses and their families. However, the science of palliative care is advancing and this new edition will contribute to a better understanding. This new edition offers 86 updated or new chapters out of 108, written by experts in their given fields, providing up-to-date information on a wide range of topics of relevance to those providing care towards the end of life no matter what the disease may be. We present a global perspective on contemporary and classic issues in palliative care with authors from a wide range of disciplines involved in this essential aspect of care. The Textbook includes sections addressing aspects such as symptom management and care provision, organization of care in different settings, care in specific disease groups, palliative care emergencies, ethics, public health approaches and research in palliative care. This new Textbook will be of value to practitioners in all disciplines and professions where the care of people approaching death is important, specialists as well as non-specialists, in any setting where people with serious advanced illnesses are residing. It is also an important resource for researchers, policy- and decision-makers at national or regional levels. Neither the science nor the art of palliative care will stand still so the Editors and contributors from all over the world aim to keep this Textbook updated so that the reader can find new evidence and approaches to care.

Cor orans. Instrukcja wykonawcza o ?e?skim ?yciu kontemplacyjnym Congregazione per gli Istituti di Vita Consacrata e le Società di Vita Apostolica, 2019

EAPC Carlos Centeno, European Association for Palliative Care, 2013

Cancer Control World Health Organization, 2007 In 2005, 7.6 million people died of cancer. More than 70% of those deaths occurred in low and middle income countries. WHO has developed a series of six modules that provides practical advice for programme managers and policy-makers on how to advocate, plan and implement effective cancer control programmes, particularly in low and middle income countries. The WHO guide is a response to the World Health Assembly resolution on cancer prevention and control (WHA58.22), adopted in May 2005, which calls on Member States to intensify action against cancer by developing and reinforcing cancer control programmes.

Fragility Fracture Nursing Karen Hertz, Julie Santy-Tomlinson, 2018-06-15 This open access book aims to provide a comprehensive but practical overview of the knowledge required for the assessment and management of the older adult with or at risk of fragility fracture. It considers this from the perspectives of all of the settings in which this group of patients receive nursing care. Globally, a fragility fracture is estimated to occur every 3 seconds. This amounts to 25 000 fractures per day or 9 million per year. The financial costs are reported to be: 32 billion EUR per year in Europe and 20 billion USD in the United States. As the population of China ages, the cost of hip fracture care there is likely to reach 1.25 billion USD by 2020 and 265 billion by 2050 (International Osteoporosis Foundation 2016). Consequently, the need for nursing for patients with fragility fracture across the world is immense. Fragility fracture is one of the foremost challenges for health care providers, and the impact of each one of those expected 9 million hip fractures is significant pain, disability, reduced quality of life, loss of independence and decreased life expectancy. There is a need for coordinated, multi-disciplinary models of care for secondary fracture prevention based on the increasing evidence that such models make a difference. There is also a need to promote and facilitate high quality, evidence-based effective care to those who suffer a fragility fracture with a focus on the best outcomes for recovery, rehabilitation and secondary prevention of further fracture. The care community has to understand better the experience of fragility fracture from the perspective of the patient so that direct improvements in care can be based on the perspectives of the users. This book supports these needs by providing a comprehensive approach to nursing practice in fragility fracture care.

Approaching Death Institute of Medicine, Committee on Care at the End of Life, 1997-10-16 When the end of life makes its inevitable appearance, people should be able to expect reliable, humane, and effective caregiving. Yet too many dying people suffer unnecessarily. While an overtreated dying is feared, untreated pain or emotional abandonment are equally frightening. Approaching Death reflects a wide-ranging effort to understand what we know about care at the end of life, what we have yet to learn, and what we know but do not adequately apply. It seeks to build understanding of what constitutes

good care for the dying and offers recommendations to decisionmakers that address specific barriers to achieving good care. This volume offers a profile of when, where, and how Americans die. It examines the dimensions of caring at the end of life: Determining diagnosis and prognosis and communicating these to patient and family. Establishing clinical and personal goals. Matching physical, psychological, spiritual, and practical care strategies to the patient's values and circumstances. Approaching Death considers the dying experience in hospitals, nursing homes, and other settings and the role of interdisciplinary teams and managed care. It offers perspectives on quality measurement and improvement, the role of practice guidelines, cost concerns, and legal issues such as assisted suicide. The book proposes how health professionals can become better prepared to care well for those who are dying and to understand that these are not patients for whom nothing can be done.

Report on the National Strategy for Palliative Care Singapore. Ministry of Health,Duke-NUS Graduate Medical School Singapore,Lien Centre for Palliative Care,2011 Lien Centre for Palliative Care was commissioned by Ministry of Health to formulate a National Strategy for Palliative Care, in consultation with key stakeholders in the healthcare system. A National Strategy for Palliative Care Workgroup was appointed.

Dying in America Institute of Medicine,Committee on Approaching Death: Addressing Key End-of-Life Issues,2015-03-19 For patients and their loved ones, no care decisions are more profound than those made near the end of life. Unfortunately, the experience of dying in the United States is often characterized by fragmented care, inadequate treatment of distressing symptoms, frequent transitions among care settings, and enormous care responsibilities for families. According to this report, the current health care system of rendering more intensive services than are necessary and desired by patients, and the lack of coordination among programs increases risks to patients and creates avoidable burdens on them and their families. Dying in America is a study of the current state of health care for persons of all ages who are nearing the end of life. Death is not a strictly medical event. Ideally, health care for those nearing the end of life harmonizes with social, psychological, and spiritual support. All people with advanced illnesses who may be approaching the end of life are entitled to access to high-quality, compassionate, evidence-based care, consistent with their wishes. Dying in America evaluates strategies to integrate care into a person- and family-centered, team-based framework, and makes recommendations to create a system that coordinates care and supports and respects the choices of patients and their families. The findings and recommendations of this report will address the needs of patients and their families and assist policy makers, clinicians and their educational and credentialing bodies, leaders of health care delivery and financing organizations, researchers, public and private funders, religious and community leaders, advocates of better care, journalists, and the public to provide the best care possible for people nearing the end of life.

Health Literacy and Palliative Care National Academies of Sciences, Engineering, and Medicine,Health and Medicine

Division, Board on Population Health and Public Health Practice, Roundtable on Health Literacy, 2016-06-18 The Institute of Medicine Roundtable on Health Literacy convened a 1-day public workshop to explore the relationship between palliative care and health literacy, and the importance of health literate communication in providing high-quality delivery of palliative care. Health Literacy and Palliative Care summarizes the discussions that occurred throughout the workshop and highlights the key lessons presented, practical strategies, and the needs and opportunities for improving health literacy in the United States.

Textbook of Palliative Medicine Eduardo Bruera, Irene Higginson, Charles F von Gunten, 2009-01-30 Textbook of Palliative Medicine provides an alternative, truly international approach to this rapidly growing specialty. This textbook fills a niche with its evidence-based, multi-professional approach and global perspective ensured by the international team of editors and contributing authors. In the absence of an international curriculum for the study of palliative medicine, this textbook provides essential guidance for those both embarking upon a career in palliative medicine or already established in the field, and the structure and content have been constructed very much with this in mind. With an emphasis on providing a service anywhere in the world, including the important issue of palliative care in the developing nations, Textbook of Palliative Medicine offers a genuine alternative to the narrative approach of its competitors, and is an ideal complement to them. It is essential reading for all palliative care physicians in training and in practice, as well as palliative care nurses and other health professionals in the palliative care team

Care of the Imminently Dying Judith Paice, 2015-11-20 Palliative care is an essential element of our health care system and is becoming increasingly significant amidst an aging society and organizations struggling to provide both compassionate and cost-effective care. Palliative care is also characterized by a string interdisciplinary approach. Nurses are at the center of the palliative care team across settings and populations. The seventh volume in the HPNA Palliative Nursing Manuals series, *Care of the Imminently Dying* provides an overview of symptom management when a patient is reaching the end of their life. This volume covers delirium and the advantages of early diagnosis, determining the presence of dyspnea, death rattle, or cough, urgent syndromes that may appear the end of life, palliative sedation, and the withdrawal of life-sustaining therapies. The content of the concise, clinically focused volumes in the HPNA Palliative Nursing Manuals series is one resource for nurses preparing for specialty certification exams and provides a quick-reference in daily practice.

Compassionate Communities Klaus Wegleitner, Katharina Heimerl, Allan Kellehear, 2015-06-26 Compassionate communities are communities that provide assistance for those in need of end of life care, separate from any official health service provision that may already be available within the community. This idea was developed in 2005 in Allan Kellehear's seminal volume- *Compassionate Cities: Public Health and End of Life Care*. In the ensuing ten years the theoretical aspects of the idea have been continually explored, primarily rehearsing academic concerns rather than practical ones. Compassionate

Communities: Case Studies from Britain and Europe provides the first major volume describing and examining compassionate community experiments in end of life care from a highly practical perspective. Focusing on community development initiatives and practice challenges, the book offers practitioners and policy makers from the health and social care sectors practical discussions on the strengths and limitations of such initiatives. Furthermore, not limited to providing practice choices the book also offers an important and timely impetus for other practitioners and policy makers to begin thinking about developing their own possible compassionate communities. An essential read for academic, practitioner, and policy audiences in the fields of public health, community development, health social sciences, aged care, bereavement care, and hospice & palliative care, Compassionate Communities is one of only a handful of available books on end of life care that takes a strong health promotion and community development approach.

Oxford Handbook of Palliative Care Max Watson, Caroline Lucas, Andrew Hoy, Jo Wells, 2009-03-26 The Oxford Handbook of Palliative Care covers all aspects of palliative care in a concise and succinct format suited to busy professionals who need to access key information in their daily care of patients. This new edition is revised throughout, with an additional emphasis on the nursing aspects of Palliative Care. The authors have included new sections on international palliative care, self care and liaison palliative care in acute hospitals. There is also extended material on the use of antibiotics, palliative care research and quality of life issues. The second edition of the Oxford Handbook of Palliative Care continues to be an invaluable resource for all health professionals working with adults, children and families with palliative care needs.

Palliative Care in Oncology Bernd Alt-Epping, Friedemann Nauck, 2015-03-26 Palliative care provides comprehensive support for severely affected patients with any life-limiting or life-threatening diagnosis. To do this effectively, it requires a disease-specific approach as the patients' needs and clinical context will vary depending on the underlying diagnosis. Experts in the field of palliative care and oncology describe in detail the needs of patients with advanced cancer in comparison to those with non-cancer disease and also identify the requirements of patients with different cancer entities. Basic principles of symptom control are explained, with careful attention to therapy for pain associated with either the cancer or its treatment and to symptom-guided antineoplastic therapy. Complex therapeutic strategies for palliative cancer patients are highlighted that involve both cancer- and symptom-directed options and address a range of therapeutic aims. Issues relating to drug use in palliative cancer care are fully explored, and a separate section is devoted to care in the final phase. A range of organizational and policy issues are also discussed, and the book concludes by considering likely future developments in palliative care for cancer patients. Palliative Care in Oncology will be of particular interest to palliative care physicians who are interested in broadening the scope of their disease-specific knowledge, as well as to oncologists who wish to learn more about modern palliative care concepts relevant to their day-to-day work with cancer patients.

LGBTQ-Inclusive Hospice and Palliative Care Kimberly D. Acquaviva, 2017-05-23 This is the only handbook for

hospice and palliative care professionals looking to enhance their care delivery or their programs with LGBTQ-inclusive care. Anchored in the evidence, extensively referenced, and written in clear, easy-to-understand language, LGBTQ-Inclusive Hospice and Palliative Care provides clear, actionable strategies for hospice and palliative physicians, nurses, social workers, counselors, and chaplains.

Ensuring Balance in National Policies on Controlled Substances World Health Organization, 2011 This book provides guidance on policies and legislation with regards to availability, accessibility, affordability and control of medicines made from substances regulated under the international drug control conventions, herein referred to as controlled medicines. Their scope encompasses all controlled medicines, but with a specific focus on essential medicines. Controlled medicines play an important role in several areas of medicine, including pain treatment, treatment of opioid dependence, emergency obstetrics, psychiatry and neurology. The availability, accessibility and affordability of controlled medicines are important issues for all countries, but problematic for most of them. The World Health Organization (WHO) promotes governments, civil society and other interested individuals to strive for the maximum public health outcome of policies related to these medicines. WHO considers the public health outcome to be at its maximum (or balanced) when the optimum is reached between maximizing access for rational medical use and minimizing substance abuse. Policy-makers, academia, civil society and other individuals whose area of work or interest is drug control or public health may potentially work with these guidelines in order to ensure that better use is made of controlled medicines and that more patients benefit from the advantages that their rational use can offer.

Palliative Care Nursing: Principles and Evidence for Practice Catherine Walshe, Nancy Preston, Bridget Johnston, 2018-05-18 What can nurses do to support those receiving palliative care? How do you ensure clear communication and maintain patients' and families' preferences? Palliative Care Nursing is essential reading for nursing students, professional nurses and other health and social care professionals providing supportive and palliative care to those with advanced illness or who are towards the end of life. This third edition of the acclaimed textbook has been extensively revised and examines important research studies, key debates around care and strategies to advance palliative care nursing. In four sections, the book covers key elements of nursing practice towards the end of life: • Defining the palliative care patient • Providing palliative nursing care • Caring around the time of death • Challenging issues in palliative care nursing Leading authors address contemporary issues and explore how to provide high quality person-centred palliative care, encouraging application to practice through exercises and case studies. Chapters completely reworked or new for this edition include those on communication, living with uncertainty, bereavement care, the costs of caring, nurses' decision-making and capacity, and palliative care worldwide. The clarity of evidence presented and coverage of a diverse range of topics make this the foundational textbook for all studying palliative care at pre-registration level, postgraduate level or as part of CPD study.

With a foreword by last edition editor, Professor Sheila Payne, Lancaster University, UK. 'I welcome this third edition of Palliative Care Nursing and congratulations to the new team who have provided us with a dynamic and innovative development of a core text for palliative nursing practice. As the largest workforce in palliative care, and given the changing face of clinical practice for nurses, including increased educational opportunity and expanding roles and responsibilities, this book is timely in its focus on critical issues which frame and scope the reality of palliative care and the nursing contribution to that discipline. The learning exercises, in particular, offer tools for educators and clinicians to reflect on practice and understand new ways of knowing in palliative care. It will be an excellent resource for nursing, both in the UK and Ireland and to the wider international audience, having drawn on the breadth of global nursing expertise to bring this book together'. Philip Larkin, Professor of Clinical Nursing (Palliative Care), University College Dublin and Our Lady's Hospice and Care Services, Dublin, Ireland; President, European Association for Palliative Care 'This is a book of substance that captures the current status of palliative nursing, including the values and research evidence that underpin it. The changing nature of palliative nursing as an evidence-based specialism is balanced with practical skills and insights from experts, and also considers the needs of those working with, or concerned about, the dying person's well-being. It covers a range of challenging issues as well as drawing on the wisdom of those who actually undertake this work on a daily basis. I hope that students and practitioners from all disciplines will find this a useful resource to understand the art and craft of good palliative nursing'. Professor Daniel Kelly, Fellow of the Royal College of Nursing and Royal College of Nursing Chair of Nursing Research, Cardiff University, UK

Handbook of Public Policy Implementation Fritz Sager, Céline Mavrot, Lael R. Keiser, 2024-07-05 In this comprehensive Handbook, international experts examine theoretical and empirical research to analyse a core element of the public policy process: implementation. Traversing numerous sub-disciplines and traditions including top-down and bottom-up approaches to public policy implementation research, the chapters present a synthesis of the state of scholarship and stimulate future thinking in the field.

Palliative Medicine E-Book T. Declan Walsh, Augusto T. Caraceni, Robin Fainsinger, Kathleen M. Foley, Paul Glare, Cynthia Goh, Mari Lloyd-Williams, Juan Nunez Olarte, Lukas Radbruch, 2008-10-07 As a palliative medicine physician, you struggle every day to make your patients as comfortable as possible in the face of physically and psychologically devastating circumstances. This new reference equips you with all of today's best international approaches for meeting these complex and multifaceted challenges. In print and online, it brings you the world's most comprehensive, state-of-the-art coverage of your field. You'll find the answers to the most difficult questions you face every day...so you can provide every patient with the relief they need. Equips you to provide today's most effective palliation for terminal malignant diseases • end-stage renal, cardiovascular, respiratory, and liver disorders • progressive neurological conditions • and HIV/AIDS.

Covers your complete range of clinical challenges with in-depth discussions of patient evaluation and outcome assessment • ethical issues • communication • cultural and psychosocial issues • research in palliative medicine • principles of drug use • symptom control • nutrition • disease-modifying palliation • rehabilitation • and special interventions. Helps you implement unparalleled expertise and global best practices with advice from a matchless international author team. Provides in-depth guidance on meeting the specific needs of pediatric and geriatric patients. Assists you in skillfully navigating professional issues in palliative medicine such as education and training • administration • and the role of allied health professionals. Includes just enough pathophysiology so you can understand the whys of effective decision making, as well as the how tos. Offers a user-friendly, full-color layout for ease of reference, including color-coded topic areas, mini chapter outlines, decision trees, and treatment algorithms. Comes with access to the complete contents of the book online, for convenient, rapid consultation from any computer.

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