Palliative Care Examples For Progress Notes

Collaborative Practice in Palliative Care explores how different professions work collaboratively across professional, institutional, social, and cultural boundaries to enhance palliative care. Analysing palliative care as an interaction between different professionals, clients, and carers, and the social context or community within which the interaction takes place, it is grounded in up-to-date evidence, includes global aspects of palliative care and cultural diversity as themes running throughout the book, and is replete with examples of good and innovative practice. Drawing on experiences from within traditional specialist palliative care settings like hospices and community palliative care services, as well as more generalist contexts of the general hospital and primary care, this practical text highlights the social or public health model of palliative care. Designed to support active learning, it includes features such as case studies, summaries, and pointers to other learning resources. This text is an important reference for all professionals engaged in palliative care, particularly those studying for post-qualification programmes in the area. Updated with stories from people who have been inspired by the original text, a guide to connecting with what matters most identifies four phrases for honoring relationships, letting go of unhealthy emotions, and living life fully.

This book offers a comprehensive overview of the compatibility of palliative care with the vision of human dignity in the Catholic moral and theological traditions. The unique value of this book is that it presents expert analysis of the major domains of palliative care and how they are compatible with, and enhanced by, the holistic vision of the human person in Catholic health care. This volume will serve as a critically important ethical and theological resource on palliative care, including care at the end of life, for bioethicists, theologians, palliative care specialists, other health care professionals, Catholic health care sponsors, health care administrators and executives, clergy, and students. Patients receiving palliative care and their families will also find this book to be a clarifying and reassuring resource.

The Oxford Textbook of Palliative Social Work is a comprehensive, evidence-informed text that addresses the needs of professionals who provide interdisciplinary, culturally sensitive, biopsychosocial-spiritual care for patients and families living with life-threatening illness. Social workers from diverse settings will benefit from its international scope and wealth of patient and family narratives. Unique to this scholarly text is its emphasis on the collaborative nature inherent in palliative care. This definitive resource is edited by two leading palliative social work pioneers who bring together an array of international authors who provide clinicians, researchers, policy-makers, and academics with a broad range of content to enrich the guidelines recommended by the National Consensus Project for Quality Palliative Care.

Providing an overview of the myriad ways that we are touched by death and dying, both as an individual and as a member of society, this book will help readers understand our relationship with death. Kastenbaum and Moreman show how various ways that individual and societal attitudes influence both how and when we die and how we live and deal with the knowledge of death and loss. This landmark text draws on contributions from the social and behavioral sciences as well as the humanities, such as history, religion, philosophy, literature, and the arts, to provide thorough coverage of understanding death and the dying process. Death, Society, and Human Experience was originally written by Robert Kastenbaum, a renowned scholar who developed one of the world's first death education courses. Christopher Moreman, who has worked in the field of death studies for almost two decades specializing in afterlife beliefs and experiences, has updated this edition.

Physician Assistant: A Guide to Clinical Practice E-Book

Rethinking palliative care

Living with Dying

Improving Palliative Care for Cancer

Health Care Policy in an Age of New Technologies

Textbook of Palliative Medicine and Supportive Care

Collaborative Practice in Palliative CareRoutledge

Providing an understanding of the relationship with death, both as an individual and as a member of society. This book is intended to contribute to your understanding of your relationship with death, both as an individual and as a member of society. Kastenbaum shows how individual and societal attitudes influence both how and when we die and how we live and deal with the knowledge of death and loss. Robert Kastenbaum is a renowned scholar who developed one of the world's first death education courses and introduced the first text for this market. This landmark text draws on contributions from the social and behavioral sciences as well as the humanities, such as history, religion, philosophy, literature, and the arts, to provide thorough coverage of understanding death and the dying process. Learning Goals Upon completing this book, readers should be able to: -Understand the relationship with death, both as an individual and as a member of society -See how social forces and events affect the length of our lives, how we grieve, and how we die -Learn how dying people are perceived and treated in our society and what can be done to provide the best possible care -Master an understanding of continuing developments and challenges to hospice (palliative care). -Understand what is becoming of faith and doubt about an afterlife

"It is so important to advocate for things that may not always seem possible. Getting to work with patients/families at the end of their life is the ultimate honor." - Lauren G Markham, MSW, LCSW, APHSW-C "In this work, one witnesses both depths of human suffering and heights of human transcendence that can inspire both awe and fear. At those times, I have found that surrendering my need to be "an expert" and instead, allow myself to simply be a "human" is the wisest action." - Kerry Irish, LCSW, OSW-C, FAOSW"--

Dr. Hallenbeck has written a guide for clinicians who want an introduction to palliative care that addresses "big picture" issues as well as provides specific practical advice. Topics addressed range from an overview of death and dying in the modern world to discussions of pain and symptom management, communication techniques and palliative care consults.

Each year, neonatal Intensive care units (NICUs) in the U.S. and around the world help thousands of sick or premature newborns survive. NICUs are committed to the ideals of family-centered care, which encourages shared decision-making between parents and NICU caregivers. In cases of infants with conditions marked by high mortality, morbidity, or great suffering, family-centered care affirms the right of parents to assist in making decisions regarding aggressive treatment for their infant. Often, these parents' difficult and intimate decisions are shaped profoundly by their religious beliefs. In light of this, what precisely are the teachings of the major world religious traditions about the status and care of the premature or sick newborn? Few studies have grappled with what major religious traditions teach about the care of the newborn or how these teachings may bear on parents' decisions. This volume seeks to fill this gap, providing information on religious teachings about the newborn to the multidisciplinary teams of NICU professionals (neonatologists, advance practice nurses, social workers), as well as to parents of NICU patients, and students of bioethics. In chapters dealing with Judaism, Catholicism, Denominational Protestantism, Evangelical Protestantism, African American Protestantism, Sunni and Shi'a Islam, Hinduism, Buddhism, Navajo religion, and Seventh Day Adventism, leading scholars develop the teachings of these traditions on the status, treatment, and ritual accompaniments of care of the premature or sick newborn. This is an essential book that will serve as a first resort for clinicians who need to understand the religious dynamics influencing anyone making a difficult decision about her sick newborn.

Oxford Textbook of Palliative Social Work

Exploring Issues of Care, Dying and the End of Life

Issues in Palliative Care Research

Counting What Counts: Workshop Summary

The Routledge History of Death since 1800

The Path to Continuously Learning Health Care in America

In Exploring Issues of Care, Dying and the End of Life, practitioners and academics from a range of disciplines and nationalities discuss matters pertinent to the end of life. Together they explore a variety of issues including communication, facing up to and handling death, as well as investigating what constitutes the 'good death'.

Tens of millions of people around the world live with chronic pain - many in such severe pain they are disabled by it. The Institute of Medicine estimates that chronic pain costs the U.S. alone $560 to $635 billion a year in direct medical costs and lost productivity. Morphine, an effective painkiller, costs only three cents a dose, yet because of excessive regulation in many countries, it is unavailable to millions of people who need it, even at the end of life. The World Health Organization notes that in addition to the one million end-stage AIDS/HIV patients who can't get morphine and other controlled medications, 5.5 million terminal cancer patients, nearly a million people suffering from accidents or violence, and an incalculable number of people living with chronic illnesses or recovering from surgery don't have access to it, either. Furthermore, women, children, older people, and the poor are disproportionally affected by inadequate pain relief. Physicians know almost nothing about chronic pain, much less how to treat it, for two reasons: medical schools barely teach it and government institutions allot almost nothing to the pain research budget. In The Global Pain Crisis: What Everyone Needs to Know®, renowned health journalist Judy Foreman addresses the most important questions about chronic pain: what is it, whom does it affect most, which pain relief methods in Western and alternative medicine are effective, what are the risks and benefits for opioids and marijuana, and how can the chronic pain crisis be resolved for good? Foreman's book is a wake-up call for a health problem that affects people across the globe, from all walks of life. Written in the classic, easy-to-read and quick reference style of the What Everyone Needs to Know® series, The Global Pain Crisis is a must-read for anyone whose life or work is affected by chronic pain.

Comprehensive in scope, exhaustive in detail, and definitive in authority, this third edition has been thoroughly updated to cover new practices, current epidemiological data, and the evolving models that support the delivery of palliative medicine to children. This book is an essential resource for anyone who works with children worldwide.

Health care quality and its affordability have become very pressing issues in the United States. All sectors of the country are attempting to push forward initiatives that will improve the health care system as well as the health of the American population in general. Despite the economical dedication to health care, about 1/5, the system remains uneven and fragmented, patient harm is quite common, care is often uncoordinated, and many more mishaps occur. There exists many obstacles to improve the nation's health care system; these include the capacity to reliably and consistently measure progress. In 2006 the Institute of Medicine (IOM) established the Roundtable on Value & Science-Driven Health Care which has since accelerated the development of a learning health system- one in which science, informatics, incentives, and culture are aligned to create a continuous learning loop. This learning loop would thus help make the health care system better. In response, the IOM organized a 2-day workshop to explore in depth the core measurement needs for population health, health care quality, and health care costs. The workshop hoped to gain a full understanding of how to improve the nation's measurement capacity to track progress in the health care system. Having this knowledge would help the nation get one step closer to the creation of an efficient learning loop. The workshop was divided into a series of sessions that focused on different aspects of measurement. Core Measurement Needs for Better Care, Better Health, and Lower Costs: Counting What Counts: Workshop Summary includes explanations and key details for these sessions: Vision, Current Measurement Capabilities, Specifying the Shape of a Core Metric Set, and Implementation. The report also features common themes within these areas, the workshop agenda, and information about those involved.

Looks at how the principles of human rights can be applied to older people in hospitals and care homes to ensure they are treated with greater dignity and respect. This report covers the leadership of the Department of Health; the implementation of the Human Rights Act by service providers.

Surgical Palliative Care

The human rights of older people in healthcare

Hospice Palliative Home Care and Bereavement Support

Principles and Evidence for Practice

Death, Society and Human Experience (1-download)

Transforming the Care of Serious Illness

The contributors examine the challenges faced by this multidisciplinary speciality as it seeks to combine high grade pain and symptom control with sensitive psychological, spiritual and social care. Ethical and resourcing aspects are discussed.

Palliative Care is the first book to provide a comprehensive understanding of the new field that is transforming the way Americans deal with serious illness. Diane E. Meier, M.D., one of the field's leaders and a recipient of a MacArthur Foundation "genius award" in 2009, opens the volume with a sweeping overview of the field. In her essay, Dr. Meier examines the roots of palliative care, explores the key legal and ethical issues, discusses th care, and presents ideas on policies that can improve access to palliative care. Dr. Meier's essay is followed by reprints of twenty-five of the most important articles in the field. They range from classic pieces by some of the field's pioneers, such as Eric Cassel, Balfour Mount, and Elizabeth Kübler-Ross, to influential newer articles on topics such as caregiving and cost savings of palliative care. The reprints cover a wide range of topics inclu seriously ill is so important Efforts to cope with advanced illness Legal and ethical issues Pain management Cross-cultural issues Philosophical perspective The demand for palliative care has been nothing short of stunning—largely because of palliative care's positive impact on both the quality and the cost of care provided to seriously ill individuals. By providing a wide-ranging perspective on this growing field, this book will serve as a guide approaches that will lead to better health care for all Americans.

"It has been a true pleasure to have had the opportunity to peruse the second edition ofPalliative Care Nursing.... This book, authored predominately by UK-based experts, succeeds in presenting sophisticated thoughts in readily accessible language… Each chapter begins with a summary of key points, with both classic and new relevant literature well integrated into the text. I have also been particularly impressed with the editors' final chap number of crucial issues for the future development of palliative care… this second edition makes a significant contribution to both the palliative care literature as well as to nursing literature." Carol Tishelman, Karolinska Institutet, Stockholm, Sweden "I findPalliative Care Nursinga very attractive book for nurses but also for other disciplines to learn about nursing and to learn about palliative care. The book is voluminous, informative and edu Frameworks and models in this book will give nurses the opportunity to make up their own process to offer support and be a carer for the incurably ill person and his/her family as a skilled companion…. This book gives the possibility for nurses to spread one clear voice about palliative care nursing. Congratulations to all the authors...." Martine De Vlieger, Palliatieve Hulpverlening Antwerpen v.z.w., University of Antwerp, Belgium "This book reading for nurses and other health care workers who are involved in the care of people in the final stages of life. It provides a comprehensive account of the major issues (clinical, professional, sociological and political) that confront contemporary palliative care while also offering strategies to move forward. The 'real world' of palliative care is described and critiqued and the rhetoric is dispensed with. This book is a vital resource for nursing teaching." Associate Professor, Peter Hudson (RN, PhD). Director of the Centre for Palliative Care Research and Education, St Vincent's Hospital and The University of Melbourne, Australia. "This is an excellent book for anyone completing either an academic qualification or who wants to understand the who, what and where of palliative care both in the UK and abroad. Its detail is balanced with case studies and practical illustrations that br its writing to life...For reference purposes for anyone completing academic work it has to be an absolute must." Nursing Times The second edition of this innovative textbook has been extensively revised and updated to reflect new global developments in palliative care. This textbook reviews current research and examines the evidence base for palliative care policy and practice. Over a third of the chapters are newly commissioned from lea Building on the widely acclaimed original edition, the textbook focuses on palliative care for adults in a variety of care environments. The first three sections use a novel framework – the trajectory of life-limiting illness – to cover key issues including: What happens to people as they become ill How individuals cope as they near death and are dying How families and friends deal with bereavement and loss The final section addresses contemp inter-professional working. The book is written with helpful overviews and in an informative and reader-friendly style. There are numerous examples of clinical situations and research studies which are examined in depth to illustrate debates in palliative care. The textbook spans the range of end-of-life contexts which are of relevance to practitioners, educationalists and researchers. Palliative Care Nursingis essential reading for post-qualifica nurses and health and social care professionals who provide care to people with advanced illness and those who are near the end of life. This broad ranging critical text will be invaluable to students and practitioners working with people and their families near the end of life.

"the thoroughness of the text has to be admired. It is an excellent starting point for students of palliative care which makes an important contribution to any library."-British Journal of Hospital Medicine" covers a plethora of topics ranging from the development of palliative medicine in different countries to clinical topics and bioethics an

Revolutionary advances in biomedical research and information systems technology pose new and difficult issues for American health care policy, especially in the context of managed care. Health Care Policy in a New Millennium takes on this challenging array of issues where the dignity of individual life meets the imperatives of national-level health-care systems - patients' rights, rationing of care, organ transplants, genetic research, confid the right to die, and other ethical dilemmas. The book places these critical questions about the quality of life in our society in their political, legal, social, economic, and ethical contexts.

Best Care at Lower Cost

Palliative Care in Oncology

A Guide to Palliative Care

Participatory Research in Palliative Care

Palliative Care and Catholic Health Care

Ageing populations mean that palliative and end of life care for older people must assume greater priority. Indeed, there is an urgent need to improve the experiences of older people at the end of life, given that they have been identified as the 'disadvantaged dying'. To date, models of care are underpinned by the ideals of specialist palliative care which were developed to meet the needs of predominantly middle-aged and 'young old' people, and evidence suggests these may not be adequate for the older population group. This book identifies ways forward for improving the end of life experiences of older people by taking an interdisciplinary and international approach. Providing a synergy between the currently disparate literature of gerontology and palliative care, a wide range of leading international experts contribute to discussions regarding priority areas in relation to ageing and end of life care. Some authors take a theoretical focus, others a very practical approach rooted in their clinical and research experience. The issues covered are diverse, as are the countries in which discussions are contextualised. Those working in both palliative care and gerontology will find the issues and advice discussed in this book hugely topical and of real practical value.

Prepare for every stage of your physician assistant career with Physician Assistant: A Guide to Clinical Practice, 5th Edition - the one text that takes you from your PA coursework through clinical practice! Concise, easy to read, and highly visual, this all-in-one resource by Ruth Ballweg, Edward M. Sullivan, Darwin Brown, and Daniel Vetrosky delivers the current, practical guidance you need to know to succeed in any setting. Consult this title on your favorite e-reader with intuitive search tools and adjustable font sizes. Elsevier eBooks provide instant portable access to your entire library, no matter what device you're using or where you're located. Master all the core competencies you need to know for certification or recertification. Navigate today's professional challenges with new chapters on NCCPA Specialty Recognition; Communication Issues; the Electronic Health Record; Patient Safety and Quality of Care; Population-Based Practice; and Physician Assistants and Supervision. Meet ARC-PA accreditation requirements with coverage of key topics such as Student Safety in Clinical Settings, Health Care Delivery Systems, Population-Based Practice, and Mass Casualties/Disasters. Keep up with the PA competencies that are endorsed by the AAPA, PAEA, NCCPA, and ARC-PA. Master key concepts and clinical applications thanks to a succinct, bulleted writing style; convenient tables; practical case studies; and clinical application questions throughout. Retain what you've learned and easily visualize every aspect of clinical practice with a new full-color design and illustrations throughout. Explore global options with expanded coverage of physician assistants in international medicine.

This book's striking message is that palliative care does not deliver on its aims to value people who are dying and make death and dying a natural part of life. This book draws from wider social science perspectives and critically and specifically applies these perspectives to palliative care and its dominant medical model. Applying Social Role Valorisation, the author argues for the de-institutionalisation of palliative care and the development of an alternative framework to the approaches found in hospices, palliative care units and community-based palliative care services. He offers a new conceptualisation of death and loss that refines and expands modern understandings in a way that also resonates with traditional religious views concerning death. Wide-ranging recommendations advise fundamental change in the concept of palliative care, the way support and services are organised and the day to day practice of palliative care. Rethinking palliative care will be of interest to academics, students and practitioners in palliative care as well as those in disability, social policy, sociology, social work, religion, thanatology, nursing and other health related fields.

In our society's aggressive pursuit of cures for cancer, we have neglected symptom control and comfort care. Less than one percent of the National Cancer Institute's budget is spent on any aspect of palliative care research or education, despite the half million people who die of cancer each year and the larger number living with cancer and its symptoms. Improving Palliative Care for Cancer examines the barriersâ€"scientific, policy, and socialâ€"that keep those in need from getting good palliative care. It goes on to recommend public- and private-sector actions that would lead to the development of more effective palliative interventions; better information about currently used interventions; and greater knowledge about, and access to, palliative care for all those with cancer who would benefit from it.

The Routledge History of Death Since 1800 looks at how death has been treated and dealt with in modern history – the history of the past 250 years – in a global context, through a mix of definite, often quantifiable changes and a complex, qualitative assessment of the subject. The book is divided into three parts, with the first considering major trends in death history and identifying widespread patterns of change and continuity in the material and cultural features of death since 1800. The second part turns to specifically regional experiences, and the third offers more specialized chapters on key topics in the modern history of death. Historical findings and debates feed directly into a current and prospective assessment of death, as many societies transition into patterns of ageing that will further alter the death experience and challenge modern reactions. Thus, a final chapter probes this topic, by way of introducing the links between historical experience and current trajectories, ensuring that the book gives the reader a framework for assessing the ongoing process, as well as an understanding of the past. Global in focus and linking death to a variety of major developments in modern global history, the volume is ideal for all those interested in the multifaceted history of how death is dealt with in different societies over time and who want access to the rich and growing historiography on the subject. Chapter 1 of this book is freely available as a downloadable Open Access PDF under a Creative Commons Attribution-Non Commercial-No Derivatives 4.0 license at https://tandfbis.s3-us-west-2.amazonaws.com/rt-files/docs/Open+Access+Chapters/9780429028274_oachapter1.pdf.

Palliative Care

Oxford Textbook of Palliative Medicine

Charting Spiritual Care

Palliative Care Nursing

Key Concepts in Palliative Care

The Four Things That Matter Most - 10th Anniversary Edition

Taking account of the British government's "End of Life Care Strategy", contributors set out the key issues affecting practice across a range of health and social care contexts. The book covers topics ranging from dying and death to symptom management and spiritual care, backed up with practical examples. Each entry comprises: a snapshot definition of the topic, key points, a discussion of the main debates, links to practice through thought-provoking case histories, and suggestions for further reading.

Participatory research is a relatively new method of researching practice especially within palliative care. It differs from other methodologies in that there is an expectation of action within the research process. The values that underpin participatory research are collaboration, empowerment, and reflection. In the current climate of collaboration and working with people in healthcare, participatory research methods are gaining increasing interest when there is a desire to bring about change. Organisational change is becoming an important focus as we look at ways of not only reducing costs but at the same time improving quality of care. While palliative care puts the patient and family at the centre, Participatory Research in Palliative Care discusses a new research methodology that puts practitioners at the heart of the research process as collaborators who work together with researchers to resolve problems in practice. Divided into three sections, it provides theoretical groundings of action research, a greater focus on exemplars from studies within palliative care, and discusses prominent issues when using such a methodology. All three sections are illustrated by an action research study undertaken by the author within a palliative care setting. Participatory Research in Palliative Care is written by international, multi-disciplinary authors who explore a collaborative approach to embark on research. It will appeal to health and social care professionals, academics undertaking research within palliative care, and the management of organisations where people with end of life care needs are cared for, including long-term care homes.

Volunteers have a long been involved in supporting the delivery of palliative care. Indeed in some countries, the range and quality of hospice and palliative care services depends on the involvement of volunteers. Hospice and palliative care services and volunteering are changing. As society develops, so too does volunteering. Volunteers have growing expectations of organizations, and increasingly seek roles that meet their needs and aspirations, rather than fitting in with organizational approaches. As hospice and palliative care services experience increasing and changing demands for their services due to aging populations with complex healthcare needs, we need to recognize that volunteers have a vital role to play in supporting the delivery of services of the future. The Changing Face of Volunteering in Hospice and Palliative Care explores the complex phenomenon that is volunteering in hospice and palliative care in different countries. It considers how and why volunteering is changing, through the contributions of authors from Western and Eastern Europe, North America, Australia, Africa, and India. It reflects on the influence of culture and organisational contexts, in addition to management approaches, legislative, and political influences, highlighting factors that contribute to the success of volunteering. Contributing to knowledge and understanding in the field of volunteering in hospice and palliative care internationally, this book highlights the factors that contribute to the success of volunteering models, allowing readers to see possibilities for change and find new ideas for innovative practice in their own setting.

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.

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.

The Global Pain Crisis

Collaborative Practice in Palliative Care

Research Methods in Palliative Care

Nursing Interventions and Supportive Care

Living with Ageing and Dying

Oxford Textbook of Public Health Palliative Care

This open access volume is the first academic book on the controversial issue of including spiritual care in integrated electronic medical records (EMR). Based on an international study group comprising researchers from Europe (The Netherlands, Belgium and Switzerland), the United States, Canada, and Australia, this edited collection provides an overview of different charting practices and experiences in various countries and healthcare contexts. Encompassing case studies and analyses of theological, ethical, legal, healthcare policy, and practical issues, the volume is a groundbreaking reference for future discussion, research, and strategic planning for inter- or multi-faith healthcare chaplains and other spiritual care providers involved in the new field of documenting spiritual care in EMR. Topics explored among the chapters include: Spiritual Care Charting/Documenting/Recording/Assessment Charting Spiritual Care: Psychiatric and Psychotherapeutic Aspects Palliative Chaplain Spiritual Assessment Progress Notes Charting Spiritual Care: Ethical Perspectives Charting Spiritual Care in Digital Health: Analyses and Perspectives Charting Spiritual Care: The Emerging Role of Chaplaincy Records in Global Health Care is an essential resource for researchers in interprofessional spiritual care and healthcare chaplaincy, healthcare chaplains and other spiritual caregivers (nurses, physicians, psychologists, etc.), practical theologians and health ethicists, and church and denominational representatives.

The research base for palliative care must improve as it continues to develop in increasingly evidence-based health care systems, and to provide the needs of patients and families. This is the first research methods textbook focusing on the unique needs of palliative care, aimed at improving current research and stimulating new research in the field.

This book provides an unique resource for registered nurses working in hospice palliative care at home and for the community, outside of acute care settings and also incorporates literature related to palliative care in acute health care settings, as part of the overall services and supports required. Very few resources exist which specifically address hospice palliative care in the home setting, despite the fact that most palliative care occurs outside acute care settings and is primarily supported by unpaid family caregivers. An overview of the concerns for individuals and families, as well as specific nursing interventions, from all ages would be an excellent support for nursing students and practicing registered nurses alike. The book structure begins with a description of the goals and objectives of hospice palliative care and the nursing role in providing excellent supportive care. Chapters include research findings and specifically research completed by the authors in the areas of pediatric palliative care, palliative care for those with dementia, and the needs of family caregivers in bereavement. Interventions developed by the editors are provided in this book, such as the Finding Balance Intervention for bereaved caregivers; the Reclaiming Yourself tool for bereaved spouses of partners with dementia; and The Keeping Hope Possible Toolkit for families of children with life threatening and life limiting illnesses. The development and application of these theory-based interventions are also highlighted. Videos and vignettes written by family caregivers about what was helpful for them, provide a patient-and family-centered approach./div The book will benefit nursing students, educators and practicing registered nurses by providing information, theory, and evidence from research.

The third edition of this successful book has been updated to incorporate the latest advances in the treatment of the terminally ill and includes a new section on non-malignant diseases and home care. Based on many years of clinical practice and research at St. Christopher's Hospice in London, the first teaching hospice, Living with Dying presents practical, balanced advice on the general ethical and medical principles of caring for dying patients and considers the details of the control of pain and other symptoms. The book discusses how doctors and other professionals can support the patient and family by sharing the truth with them and understanding their emotional suffering. This will continue to be an invaluable handbook for all family practitioners, hospice physicians, nurses and other health professionals. Patients and their relatives will also find this a compassionate source of factual information.

This text is the definitive resource for practicing palliative social work clinicians. It is designed to meet the needs of professionals who seek to provide culturally sensitive biopsychosocial-spiritual care for patients and families living with life-threatening illness.

What Everyone Needs to Know®

Palliative and End of Life Care for Older People

A Book About Living

eighteenth report of session 2006-07, Vol. 2: Oral and written evidence

Oxford Textbook of Palliative Care for Children

Palliative Medicine E-Book

Palliative care is rapidly evolving as a multidimensional therapeutic model devoted to improving the quality of life of all patientswith life-threatening illness. Symptom control, management of psychosocial and spiritual concerns, decision making consistent with values and goals, and care of the imminently dying that is appropriate and sensitive to the unique needs of the individual and the family--these are among the critical issues addressed through pallaitive care. As this discipline has evolved, the need for research in all these areas has become widely acknowledged. Issues in Palliative Care Research describes both the progress that already has been made in the investigation of these issues and the methodologic elements that must be addressed in future studies. The perspective is broad and the overriding goal is to inform about the state of the art in these rapidly evolving areas of research.

America's health care system has become too complex and costly to continue business as usual. Best Care at Lower Cost explains that inefficiencies, an overwhelming amount of data, and other economic and quality barriers hinder progress in improving health and threaten the nation's economic stability and global competitiveness. According to this report, the knowledge and tools exist to put the health system on the right course to achieve continuous improvement and better quality care at a lower cost. The costs of the system's current inefficiency underscore the urgent need for a systemwide transformation. About 30 percent of health spending in 2009--roughly $750 billion--was wasted on unnecessary services, excessive administrative costs, fraud, and other problems. Moreover, inefficiencies cause needless suffering. By one estimate, roughly 75,000 deaths might have been averted in 2005 if every state had delivered care at the quality level of the best performing state. This report states that the way health care providers currently train, practice, and learn new information cannot keep pace with the flood of research discoveries and technological advances. About 75 million Americans have more than one chronic condition, requiring coordination among multiple specialists and therapies, which can increase the potential for miscommunication, misdiagnosis, potentially conflicting interventions, and dangerous drug interactions. Best Care at Lower Cost emphasizes that a better use of data is a critical element of a continuously improving health system, such as mobile technologies and electronic health records that offer significant potential to capture and share health data better. In order for this to occur, the National Coordinator for Health Information Technology, IT developers, and standard-setting organizations should ensure that these systems are robust and interoperable. Clinicians and care organizations should fully adopt these technologies, and patients should be encouraged to use tools, such as personal health information portals, to actively engage in their care. This book is a call to action that will guide health care providers; administrators; caregivers; policy makers; health professionals; federal, state, and local government agencies; private and public health organizations; and educational institutions.

Palliative care has become increasingly important across the spectrum of healthcare, and with it, the need for education and training of a broad range of medical practitioners not previously associated with this field of care. Part of the Integrating Palliative Care series, this volume on surgical palliative care guides readers through the core palliative skills and knowledge needed to deliver high value care for patients with life-limiting, critical, and terminal illness under surgical care. Chapters explore the historical, philosophical, and spiritual principles of surgical palliative care, and follow the progression of the seriously ill surgical patient's journey from the pre-operative encounter, to the invasive procedure, to the post-operative setting, and on to survivorship. An overview of the future of surgical palliative care education and research rounds out the text. Surgical Palliative Care is an ideal resource for surgeons, surgical nurses, intensivists, and other practitioners who wish to learn more about integrating palliative care into the surgical field.

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.

Death, dying, loss, and care giving are not just medical issues, but societal ones. Palliative care has become increasingly professionalised, focused around symptom science. With this emphasis on minimizing the harms of physical, psychological, and spiritual stress, there has been a loss of how cultures and communities look after their dying, with the wider social experience of death often sidelined in the professionalisation and medicalisation of care. However, the people we know and love in the places we know and love make up what matters most for those undergoing the experiences of death, loss, and care giving. Over the last 25 years the theory, practice, research evidence base, and clinical applications have developed, generating widespread adoption of the principles of public health approaches to palliative care. The essential principles of prevention, harm reduction, early intervention, and health and wellbeing promotion can be applied to the universal experience of end of life, irrespective of disease or diagnosis. Compassionate communities have become a routine part of the strategy and service development in palliative care, both within the UK and internationally. The Oxford Textbook of Public Health Palliative Care provides a reframing of palliative care, bringing together the full scope of theory, practice, and evidence into one volume. Written by international leaders in the field, it provides the first truly comprehensive and authoritative textbook on the subject that will help to further inform developments in this growing specialty.

Palliative Care Perspectives

The Emerging Role of Chaplaincy Records in Global Health Care

Two Millennia of Caring for the Whole Person

Core Measurement Needs for Better Care, Better Health, and Lower Costs

The Oxford Textbook of Palliative Social Work

Death, Society, and Human Experience

Written by an established, comprehensive, multidisciplinary focused, internationally-recognized team, the sixth edition has been fully revised and brought up to date with the inclusion of recent developments in the speciality, to ensure that it retains its reputation as the definitive reference on palliative care.

A social role valorisation approach

Actions and reflections

Religion and Ethics in the Neonatal Intensive Care Unit

The Changing Face of Volunteering in Hospice and Palliative Care
New Themes In Palliative Care