palliative care research methods, end-of-life care studies, qualitative palliative research, quantitative palliative research, hospice care research design

Research Methods In Palliative Care

#palliative care research methods #end-of-life care studies #qualitative palliative research #quantitative palliative research #hospice care research design

Research methods in palliative care are crucial for advancing our understanding of patient comfort, quality of life, and treatment efficacy for individuals facing serious illnesses. This field employs a diverse range of qualitative and quantitative approaches to explore clinical outcomes, patient experiences, and caregiver support strategies, ensuring evidence-based practices enhance compassionate care.

Research methods in palliative care

Palliative and end of life care are concerned with the physical, social, psychological and spiritual care of people with advanced disease. It currently has a poorly developed research base, but the need to improve this is increasingly recognised. One of the reasons for the lack of research - and the variable quality of the research that is undertaken - is the difficulty of conducting research with very ill and bereaved people. Standard and well-established research methods may need to be adapted to work in this context. This means that existing research methods textbooks may be of limited use to palliative care practitioners seeking to do research for the first time, or to more experienced researchers wanting to apply their knowledge in palliative care settings. This research methods textbook is the first to be written specifically for palliative care. It has been edited by four experienced palliative care academics with acknowledged expertise and international reputations in this field. It encompasses methods used in both clinical and health services research in palliative care, with sections on clinical, epidemiological, survey and qualitative research, as well as a section covering skills needed in any research project. Each chapter provides readers with an up to date overview of the research method in question, an understanding of its applicability to palliative care and of the particular challenges of using it in this setting. It is essential reading for all palliative care researchers.

Participatory Research in Palliative Care

Participatory Research in Palliative Care discusses participatory research methods within the discipline of palliative care. Providing an overview of the action research methods, it uses exemplars from studies within palliative care, as well as discusses the prominent issues currently faced in this methodology from a global perspective.

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Participatory Research in Palliative Care: Actions and reflections

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.

Researching Palliative Care

There are many texts available on research methods but few that are related directly to palliative treatment. This book fills the gap in the literature and provides a useful resource for students engaged in such activity.

Research Methods For Nurses And The Caring Professions

Praise for the first edition of Research into Practice and Research Methods for Nurses and the Caring Professions: These books provide a good introduction for the uninitiated to reading and doing research. Abbott and Sapsford provide a clearly written and accessible introduction to social research ... One of their aims is to 'de-mystify' research, and in this they succeed admirably... After reading the text and the articles in the reader, and working through the various research exercises, readers should have a clear appreciation of how to evaluate other people's research and how to begin their own. -David Field, Journal of Palliative Medicine This book, now substantially revised in its second edition, is about the appreciation, evaluation and conduct of social research. Aimed at nurses, social workers, community workers and others in the caring professions, the book is particularly focused on research which evaluates and contributes to professional practice. The authors have provided many short, practical exercises in the text, and the examples are drawn mostly from projects carried out by one or two people rather than large research teams. The clear, accessible style will make this the ideal introductory text for those undertaking or studying research for the first time. The book may be used in conjunction with Research into Practice (Open University Press), a reader of useful examples selected by the same authors.

Evidence-Based Palliative Care

This book describes the current state of the art in the field of palliative care in children and adults. Special emphasis is placed on addressing the efficacy and effectiveness of palliative care models, pain and symptom management, and on measuring quality of life. In addition the book evaluates current research methods in palliative care and suggests suitable alternatives. Finally the book bridges the gap between science and practice by providing the reader with the current evidence and how it can be applied in the practice setting.

Palliative Care Nursing: Principles And Evidence For Practice

This textbook in palliative care nursing draws together the principles and evidence that underpins practice to support nurses working in specialist palliative care settings and those whose work involves end-of-life care.

Issues in Palliative Care Research

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.

Palliative Day Care in Practice

This edited book provides a comprehensive overview of the current philosophy, patterns and policies of palliative day care. It places emphasis on the need to evaluate performance in palliative day care. Audit, health economics and research in palliative day care, and their associated problems and pitfalls, are therefore described.

Palliative Care

The incidence of cancer is alarmingly increasing worldwide. The major problem that the medical profession is currently facing refers to "late presentation" patients who, for the most part, have reached the terminal stage of the illness. For these people, the only treatment option left is palliative care. Various patterns of palliation have been in practice in every culture and in every ethnic group for generations. Unfortunately, we still lack significant and sustained investment in research related to the practice of palliative care. Authors from around the globe seek more investment of public and private funds to investigate ways to improve the bedside practice of palliative care. Modern palliative care concepts were established by Dame Cicely Saunders, from London, UK, right after World War II. It is only in the past 15 to 20 years that this new discipline started to develop in the developing world. However, we still lack the essential basic biological processes involved in relieving the suffering of cancer patients while receiving palliative measures throughout the trajectory of the disease. This book owes its origins in large measure to physicians and nurses in 30 countries globally, who decided to devote their time, energy, compassion and goodwill, to the promotion of palliative care in their countries and communities, yet they lack solid evidence-based data to rely upon while extending their treatment to both patients and family members. The goal, in part, is to bridge the gap between scientists and clinicians from developed countries and those in developing countries. We have been aware of the variances between cultures, traditions, beliefs and practices. I am continually struck by the seemingly diametrical views of "knowing" and cultures and the strong overlaps that might give rise to new ideas. We hope that these new volumes will serve to inspire health professionals' and administrators' interests and appreciation for the investment in basic and clinical research that will serve to advance our understanding of the underlying physical and emotional factors involved while extending palliative care to patients suffering from cancer and other non-communicable illnesses.

A Postgraduate's Guide to Doing a Literature Review in Health and Social Care, 2e

This text is a comprehensive, highly readable guide to how to undertake a literature review in health and social care, tailored specifically for postgraduate study. Essential reading for all those undertaking any study at post-graduate level, the book provides clarity and a step by step approach to doing a literature review from start to finish which will enable you to: • Identify which type of review is appropriate for your study • Select the literature that you need to include in your review • Search for, appraise and analyse relevant literature • Write up your review Crucially the book explores the common features of a broad range of types of literature review, which serve different functions – including the literature review that is a pre-requisite prior to a larger empirical study, and the literature review that is a study in its own right. With real-life examples of written research and succinct summaries at the end of each chapter, A Post-Graduate’s Guide to Doing a Literature Review in Health and Social Care is the ideal text for students wanting to get the very most from their study.

Oxford Handbook of Palliative Care

This practical guide briefly covers the historical and epidemiological background of palliative care and the growth of palliative medicine as a specialty, before dealing with major physical, psychological, spiritual, and symptom management issues from diagnosis to bereavement care.

A Public Health Perspective on End of Life Care

Focusing on population health and discussing studies using different methodologies, this title presents a synthesis and overview of relevant research and empirical data on the end of life that can bear a basis for a more systematic 'public health of the end of life'.

How to Read and Critique Research

Do you find research challenging to read? Do you struggle to get to grips with a research paper? Understanding, critiquing and using research is a key requirement of students studying nursing and healthcare. This bookwill equip you with the skills you need to understand research and use it in your practice and academic assignments. The approach used in this book is unique: each chapter focuses on a published research paper – one you might be asked to read for a seminar or include in your academic work. In clear, straightforward language, the authors take you through each paper step by step, using it as a basis for exploring the underpinning research method or design, and how it has been reported. Key features: · Each chapter focuses on a different research method by working through a relevant research paper · Identifies the main skills you need for your course: understanding research methods and critiquing articles · Written specifically for nursing and healthcare students by experienced nursing and health care lecturers · Develops your confidence in understanding research by helping you to apply your knowledge to real research papers.

Involving Service Users in Health and Social Care Research

Service user involvement in research can range from the extremes of being the subject, to being the initiator or investigator, of a research study. The activity of the professional researcher can also range from being the person undertaking the research, to being a partner with, or mentor to, service users. This broad scope of levels of involvement is reflected in the contributions in this book, both in the research experiences reported and in the writing of the chapters themselves. With contributions coming from a range of service areas including learning disabilities, cancer care, older people and mental illness, chapters look at important research issues such as: strategies for working in true partnership avoiding ‘tokenism’ involving service users at all stages of the research process communication and terminology involving service users of different ages and experience training needs of professionals and service users problems surrounding ‘payment’ for service users other ethical and practical issues. This book is invaluable reading for researchers in health and social care from academic, professional and service user backgrounds.

Approaching Death

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."

The Patient's Wish to Die

Wish to die statements are becoming a frequent phenomenon in terminally ill patients. Those confronted by these statments need to understand the complexity of such wishes, so they can respond competently and compassionately to the requests. If misunderstood, the statements can be taken at face-value and the practitioner may not recognise that a patient is in fact experiencing ambivalent feelings at the end of life, or they may misinterpret the expressed wish to die as a sign of clinical depression. Public debate about the morality and ethics of various end-of-life care options has exploded in recent years. However, it has never been sensitive to the finer aspects of clinical reality or the experiences of patients. The Patient's Wish to Die: Research, Ethics, and Palliative Care brings together that reality and the patient's voice, combining them with different research approaches. It presents the best available knowledge and research methodologies about patients' wishes at the end-of-life, together with a series of ethical views and a discussion about the clinical implications for palliative care. The book presents material in an open and unbiased manner whilst remaining sensitive to the spiritual and existential dimensions of dying, and to the different cultural views that provide meaning to the individual. Written by the best specialists and ethics scholars from around the world, including palliative care practitioners and end-of-life scholars from countries where assisted dying practices are legalized and from those where it isn't, The The Patient's Wish to Die: Research, Ethics, and Palliative Care will prove essential reading for all those working or studying in the field of palliative care.

Research in Palliative Care

Patient participation and user involvement are central to current thinking about the effective delivery of desired healthcare outcomes. Working with the person who lies behind every patient is core to palliative care. A voice can only become significant when it is listened to and acted upon. With palliative care increasingly addressing the needs of people with a variety of conditions in a range of settings, as well as with advances in research, technology, and information, the challenge to be 'a voice for the voiceless' is greater than ever. This book addresses key aspects in the provision of patient-centred palliative care and tracks significant developments in user involvement. It sets the philosophy within the cultural, social and political context of modern healthcare, particularly addressing issues of quality, standards, education and bereavement. A key component in the delivery of high quality palliative care is the multi-professional team. Following a discussion of teamwork, five core professions present a critical analysis of their working practices. The book concludes with a commentary from a palliative care user and a bereaved carer. It is often somewhat glibly asserted that the patient is, or should be, at the centre of care. There have been few attempts to examine how to keep him or her there without professional needs and protocols crowding him or her out. This book asks how we listen and why we listen. The book focuses on the challenges of how professionals can keep the needs of the patient central in clinical care and how the patient can influence the direction of that care.

Patient Participation in Palliative Care

"Kidney Palliative (Supportive) Care is a field that has been in existence since at least 2000 but has yet to gain solid traction in terms of integration into nephrology practice or consistent visibility in publications and professional meetings. This is beginning to change. A search of PubMed for "Kidney AND Palliative Care" reveals over 1,300 articles with the majority having been written in the last 10 years with a particular upsurge in the last five years"--

Palliative Care in Nephrology

With the number of people requiring palliative and end of life care steadily increasing, it is the responsibility of every nurse, regardless of specialism, to know how to provide high quality care to this group of people. Yet caring for those nearing the end of life can throw up complex issues, including handling bereavement, cultural and ethical issues, delivering care in a wide variety of settings, symptom management and also ensuring your own emotional resilience. This book is specifically designed to equip nursing students and non-specialists with the essential knowledge in relation to the care and management of people nearing the end of life.

Palliative and End of Life Care in Nursing

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.

Oxford Textbook of Palliative Medicine

This book provides a new outlook on the practice of palliative care worldwide. All five continents are represented in this book by global leaders in this relatively new subspecialty. The chapters in the book re-emphasize the fact that in the 21st century, most patients in the world still lack this elementary tool to alleviate suffering physical, and even more so, emotional and spiritual which are so critical to people, especially when patients conditions become fatal. An issue that comes up again and again from almost all parts of the world, regardless of religion and traditional backgrounds, ethnicities, beliefs or faith, refers to the critical lack of basic and advanced training for physicians, nurses, volunteers and the public at large. Healthcare professionals are currently not equipped with the principles of communication with both the patient and his/her relatives. These kinds of drawbacks have to be corrected immediately. Moreover, training courses, symposia and conferences do not require large amounts of funds and can be carried out in local countries and/or regions which share a common language, culture and faith. Each country needs to create a nucleus of local champions who would then take it upon themselves to educate as many people in their own countries with the support, guidance and encouragement of international organizations that are dedicated to this mission. Almost all of the larger international institutions, e.g., the UN and WHO, preach for improvement of the current situation. Unfortunately, responses are extremely slow and not efficient. This book calls for the global health community to urgently respond and bring about a rapid change in a totally unjustified situation that still prevails in over three-quarters of the world.

Palliative Care -- Perspectives, Practices and Impact on Quality of Life

Compassionate communities are communities that provide assistance for those in need of end of life care, separate from any official heath 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.

Compassionate Communities

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 Medicine E-Book

Health geographers are increasingly turning to a diverse range of interpretative methodologies to explore the complexities of health, illness, space and place to gain more comprehensive understandings of well-being and broader social models of health and health care. Drawing upon postmodernism, many health geographers are concerned with issues of representation, the body and health care policy. Also related to an emphasis on the body is the growing literature in feminist health geography that investigates the metaphorical, physical and emotional challenges of the body and disease. Reflecting these interests, the chapters in this book set out the host of creative qualitative methods being used to explore the psychosocial experiences of individuals more directly, using such traditional methods as in-depth interviews and group discussions, participant observation, diaries and discourse analysis, but also more novel techniques such as 'go-along interviews', reflexive writing, illustrations, and photographic techniques. There are several areas of qualitative research unique to geographers which figure prominently in this volume including: health and place, comparative case study analysis, and qualitative approaches to the use of geographic information systems (GIS). This collection brings together a wide range of empirical concerns related to questions of health and shines a light on the diversity of qualitative methods in practice. Illustrating how qualitative methodologies are used in diverse health contexts this book fills an important niche for health geographers but will have wide appeal to health and geographic researchers.

Practicing Qualitative Methods in Health Geographies

50 Studies Every Palliative Doctor Should Know presents key studies that have shaped the practice of palliative medicine. Selected using a rigorous methodology, the studies cover topics including: palliative care, symptom assessment and management, psychosocial aspects of care and communication, and end-of-life care. For each study, a concise summary is presented with an emphasis on the results and limitations of the study, and its implications for practice. An illustrative clinical case concludes each review, followed by brief information on other relevant studies. This book is a must-read for health care professionals and anyone who wants to learn more about the data behind clinical practice.

50 Studies Every Palliative Care Doctor Should Know

“Anne’s contribution to our understanding of the needs of young people with cancer has been unparalleled and without her extraordinary insights our services would be that much poorer.” From the foreword by Simon Davies , CEO Teenage Cancer Trust This topical and timely text provides valuable insights into the choices and experiences of palliative and end of life care for young people with cancer and other life limiting illnesses. With a focus on palliative care provision across a range of different clinical settings, this comprehensive new resource explores care in the home, the hospice and hospital. It looks at how and where families and young people can access palliative care, and what support is offered to attain their preferred place of death. Bereavement support for families is discussed, as well as a discussion of multidisciplinary work, interagency co-operation and resource issues. This title is essential reading for community children’s nurses, specialist palliative care teams, children’s hospices, school nurses, social workers and student nurses as well as families. A comprehensive resource on end of palliative are provision for children and young adults with cancer and other life limiting illnesses Timely and topical, tying in with the Department of Health palliative care strategy ‘Better Care: Better Lives’ Written in an accessible style that does not assume either detailed medical or theoretical knowledge Explores palliative care provision in a range of different clinical settings including the home, hospice, and hospital Provides valuable insights into the experiences of parents, children and young people

Palliative and End of Life Care for Children and Young People

Behavioral Intervention Research in Hospice and Palliative Care: Building an Evidence Base sets forth research considerations and guidelines to build evidence-based interventions to improve end-of-life care. It is an in-depth introduction to implementation research and showcases how a clinical need is identified to inform an intervention. The book extensively examines the various phases of intervention research, including design, implementation, evaluation, dissemination and translation. The book focuses on methodological, ethical and practical issues. The science behind the quality of hospice and palliative care lags behind that of traditional medical practice, despite the continuous growth of palliative care interdisciplinary teams. Researching, developing and testing strategies is essential to advancing the effectiveness and value of this care. Informs readers how to conduct intervention research toward identifying best care Advises readers on design, implementation and evaluation of research Provides step-by-step templates to develop an intervention study Includes mock protocols from successful intervention trials Synthesizes lessons learned by established intervention researchers in hospice and palliative care

Behavioral Intervention Research in Hospice and Palliative Care

This innovative, practical guide introduces researchers to the use of the video reflexive ethnography in health and health services research. This methodology has enjoyed increasing popularity among researchers internationally and has been inspired by developments across a range of disciplines: ethnography, visual and applied anthropology, medical sociology, health services research, medical and nursing education, adult education, community development, and qualitative research ethics.

Video-Reflexive Ethnography in Health Research and Healthcare Improvement

This volume present a series of case narratives, following individual patients and families throughout the course of illness and death in the context of hospice and palliative care. Using a variety of qualitative research methods, including participant-observation, interviews, and journal-keeping, the experiences, perceptions, and feelings of the patient, the family, and a range of caregivers are recorded, providing the reader with rich, multi-textured narratives. Going beyond conventional case reports in Medicine, typically concentrating on symptoms and treatment, these narratives depict how individuals find personal meaning in illness, and how this influences the experience and outcome of care.

Crossing Over

This work complements the Oxford Textbook of Palliative Medicine and The Handbook of Psychooncology. Topics include the role of psychiatry in terminal care, diagnosis and management of depression, suicide in the terminally ill, pain management, the nature of suffering in terminal illness, and psychotherapeutic interventions. The book also takes into consideration new directions for psychosocial palliative care research.

Handbook of Psychiatry in Palliative Medicine

The rapidly evolving field of Palliative Care focuses on the management of phenomena that produce discomfort and that undermine the quality of life of patients with incurable medical disorders. The interdisciplinary clinical purview includes those factors - physical, psychological, social, and spiritual - that contribute to suffering, undermine the quality of life, and prevent a death with comfort and dignity. Palliative Care is a fundamental part of clinical practice, the "parallel universe" to therapies directed at cure or prolongation of life. All clinicians who treat patients with chronic life threatening diseases are engaged in palliative care, continually attempting to manage complex symptomatology and functional disturbances. The scientific foundation of palliative care is advancing, and similarly, methods are needed to highlight, for practitioners at the bedside, the findings of empirical research. Topics in Palliative Care Series is divided into sections that address a range of issues. Addressing aspects of sumptom control, psyshocsoical functioning, spiritual or existential concerns, ethics, and other topics, the chapters in each section review the given area and focus on a small number of salient issues for analysis. The authors present and evaluate existing data, provide a context drawn from clinical and research settings, and integrate knowledge in a manner that is both practical and readable. The specific topics covered in Volume 5 are Cultural issues in Palliative Care, Palliative Care in Geriatrics, Communication Issues in Palliative Care, Outcomes Research in Palliative Care, Opiod Tolerance; Reality of Myth?, and Pain and other symptoms: Treatment Challenges.

Topics in Palliative Care

Textbook of Palliative Care is a 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 we still find ourselves having to explain its nature and practice to colleagues and to the public in general. 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 our understanding concerning many aspects of palliative care is developing rapidly. The book is divided into separate sections for ease of use. Over 100 chapters written by experts in their given fields provide 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 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 we aim to keep this Textbook updated as the authors find new evidence and approaches to care.

Textbook of Palliative Care

It is innately human to comfort and provide care to those suffering from cancer, particularly those close to death. Yet what seems self-evident at an individual, personal level has, by and large, not guided policy at the level of institutions in this country. There is no argument that palliative care should be integrated into cancer care from diagnosis to death. But significant barriers-attitudinal, behavioral, economic, educational, and legal-still limit access to care for a large proportion of those dying from cancer, and in spite of tremendous scientific opportunities for medical progress against all the major symptoms associated with cancer and cancer death, public research institutions have not responded. In accepting a single-minded focus on research toward cure, we have inadvertently devalued the critical need to care for and support patients with advanced disease, and their families. This report builds on and takes forward an agenda set out by the 1997 IOM report Approaching Death: Improving Care at the End of Life, which came at a time when leaders in palliative care and related fields had already begun to air issues surrounding care of the dying. That report identified significant gaps in knowledge about care at the end of life and the need for serious attention from biomedical, social science, and health services researchers. Most importantly, it recognized that the impediments to good care could be identified and potentially remedied. The report itself catalyzed further public involvement in specific initiatives-mostly pilot and demonstration projects and programs funded by the nonprofit foundation community, which are now coming to fruition.

Improving Palliative Care for Cancer

The Oxford Textbook of Palliative Nursing remains the most comprehensive treatise on the art and science of palliative care nursing available. Dr. Betty Rolling Ferrell and Dr. Judith A. Paice have invited 162 nursing experts to contribute 76 chapters addressing the physical, psychological, social, and spiritual needs pertinent to the successful palliative care team. Organized within 7 Sections, this new edition covers the gamut of principles of care: from the time of initial diagnosis of a serious illness to the end of a patient's life and beyond. This fifth edition features several new chapters, including chapters on advance care planning, organ donation, self-care, global palliative care, and the ethos of palliative nursing. Each chapter is rich with tables and figures, case examples for improved learning, and a strong evidence-based practice to support the highest quality of care. The book offers a valuable and practical resource for students and clinicians across all settings of care. The content is relevant for specialty hospice agencies and palliative care programs, as well as generalist knowledge for schools of nursing, oncology, critical care, and pediatric. Developed with the intention of emphasizing the need to extend palliative care beyond the specialty to be integrated in all settings and by all clinicians caring for the seriously ill, this new edition will continue to serve as the cornerstone of palliative care education.

Oxford Textbook of Palliative Nursing

Mixed methods research combines quantitative and qualitative research methods in a single study. The use of mixed methods research is increasingly popular in nursing and health sciences research.

This growth in popularity has been driven by the increasing complexity of research problems relating to human health and wellbeing. Mixed Method Research for Nursing and the Health Sciences is an accessible, practical guide to the design, conduct and reporting of mixed method research in nursing or the health sciences. Each chapter stands alone, describing the various steps of the research process, but contains links to other chapters. Within the text, 'real-life' examples from the published literature, doctoral theses and the unpublished work of the authors, illustrate the concepts being discussed. Places mixed methods research within its contemporary context Includes international contributions from UK, Australia, NZ and USA Provides an accessible introduction to theoretical and philosophical underpinnings Demystifies strategies for analysing mixed methods data Examines strategies for publishing mixed methods research Includes learning objectives and exemplars in each chapter Final chapters provide 'real-life' examples of applied research About the Authors: Sharon Andrew is Head of Program (Postgraduate) and Elizabeth J. Halcomb is Senior Lecturer, School of Nursing & Midwifery, University of Western Sydney. Also of Interest: The Research Process in Nursing (Fifth Edition) Edited by Kate Gerrish and Anne Lacey 978-14051-3013-4 Research Handbook for Healthcare Professionals Mary Hickson 978-14051-7737-5 Real World Research: A Resource for Social Scientists and Practitioner-Researchers Second edition Colin Robson 978-0631-21305-5 Reviewing Research Evidence for Nursing Practice: Systematic Reviews Edited by Christine Webb and Brenda Roe 978-14051-4423-0

Mixed Methods Research for Nursing and the Health Sciences

This important book fills a gap in the literature by focusing specifically on the role of interventional radiology in patients receiving palliative medicine and supportive care, a group in which the need for minimally invasive therapy is especially high. Detailed information and guidance is provided on use of the tools of interventional radiology for the purpose of problem solving in relation to a wide variety of diseases and complications. Readers will find clear explanation of the ways in which interventional radiology techniques can assist with regard to intravenous access, feeding, musculoskeletal and neurological pain relief, tumor debulking, management of bleeding and obstructions, drainages, and treatment of fistulas. Throughout, helpful tips and tricks of value in daily practice are highlighted. The book is an ideal reference on the interventional management of palliative/supportive care and the effective use of interventional radiology techniques in a multidisciplinary environment. Beyond specialists and trainees in interventional radiology, it will have broad appeal to all who deal with patients on palliative and supportive care on a day-to-day basis.

Interventional Radiology in Palliative Care

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.

Oxford Textbook of Palliative Medicine