

KEY CONCEPTS IN PALLIATIVE CARE KEY CONCEPTS SAGE (DOWNLOAD ONLY)

Alan Mclaughlin

Key Concepts In Palliative Care Key Concepts Sage Introduction

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End-of-Life Nursing Care

End of life care and palliative care is core to undergraduate nursing training. Packed with case studies and action points for student reflection, this is only book on the market targeted specifically at nurses and is more succinct than the competing titles which are suitable for those on medical degrees.

Key Concepts in Nursing

Nursing can be complex and challenging to new students and trainee practitioners. Key Concepts in Nursing provides a much needed guide to the central topics and debates which shape nursing theory, policy and contemporary practice. From assessment to ethics, and leadership to risk management, the book offers a comprehensive yet concise guide to the professional field. Each entry features: " a snapshot definition of the concept; " a broader discussion addressing the main issues and links to practice; " key points relevant to the entry; " case studies to illustrate the application to practice; " examples of further reading. Highly readable, with clear indexing and cross referencing, this is an ideal book for trainees to turn to for learning more about key issues in nursing practice and education. It meets the validation requirements of all training programmes and will also be invaluable for nurses continuing their professional education, those returning to practice and for mentoring.

Key Concepts in Palliative Care

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Key Concepts in Social Gerontology

"Far from a dictionary, the concepts are portrayed as complex, and conflicting definitions and usages are

both noted and evaluated... Each article includes a (necessarily selective) set of references, and cross-references to other concepts included in the book... Moreover, the coverage and evaluation of the concepts is right at the current leading edge in a rapidly moving field.\" - Victor Marshall, Department of Sociology and Institute on Aging, University of North Carolina at Chapel Hill \"The concepts are very clear and very relevant, and fantastic for my group of undergraduate students... I found each concept to be succinctly and simply captured, holding enough information to satisfy initial cravings and complex enough to tempt further reading... I had originally thought that this book would provide supplementary reading as part of my 'Healthy Ageing' course, but have changed my mind to essential reading now I have had the opportunity to read the entire book.\" - Dr Meredith Tavener, Groningen University Social gerontology is a new and dynamic field reflecting the increasing interest in ageing across the world. This book provides a readily accessible guide to well established and contested issues, as well as new concepts emerging through cutting edge research in the discipline. The entries give concise, lucid knowledge on what constitutes the 'building blocks' of social gerontology and sets out a clear review of the core concepts, both classic and emerging, in this subject area. Each concept is explored in terms of its history, application, usefulness to theory and research, and significance in practice. They go beyond simple definitions of the concepts to look at how each issue has shaped the discipline of social gerontology today. This book is authored by social gerontologists from the UK and the USA. Together they present an interdisciplinary perspective and reflect a global approach to the presentation of key concepts in social gerontology.

Counselling for Grief and Bereavement

\"The authors have done their homework in reading and consulting with the prominent literature, especially regarding children. All this effort gives the book a solid background foundation and makes it readable, and well-usable, for both lay counsellors and professional providers, and for all of us who are engaged in the delicate and rewarding endeavor of Grief Therapy\" - Naji Abi-Hashem, Clinical & Cultural Psychologist, Berkeley, California Praise for the First Edition: \"The book provides an absorbing and challenging journey through the possible process involved in bereavement work, and encourages one to think broadly about how one can approach a bereaved person... this was a book I enjoyed reading very much, and which I found both theoretically sound and practically helpful\" - Bereavement Care (Cruse) Counselling for Grief and Bereavement, Second Edition is a bestselling, introductory guide for professionals who work with people experiencing bereavement through death and other forms of loss. Focusing on practical assessment and intervention strategies, Geraldine Humphrey and David Zimpfer guide readers through the essential theory and skills needed to work with clients in a way which sensitively facilitates the process of grief, initiates healing and promotes a sense of growth. Setting out the broad principles for practice, the authors go on to show how these can be applied in working with individuals, families and groups and in relation to specific issues including chronic and life-threatening illnesses, palliative care and complicated grief. Carefully chosen case examples illustrate the counselling process, while specific attention is paid throughout to ethical considerations and the possible need for referral. This fully revised and updated Second Edition features a new chapter on working with children and adolescents: both from the perspective of young people who are grieving losses and those who are receiving palliative care as patients. While focusing on the practical, the book provides a firm theoretical base by explaining key concepts such as attachment, grief and resilience. Geraldine M. Humphrey is Counsellor in the Department of Psychology at the North Canton Medical Foundation, specializing in death, illnesses, and non-death and grief. David G. Zimpfer is former Director of the Cancer Center of Ohio.

Palliative and End-of-Life Care

Palliative and End-of-Life Care, 2nd Edition provides clinicians with the guidelines and tools necessary to provide quality, evidenced-based care to patients with life-limiting illness. This text describes the care and management of patients with advanced disease throughout the disease trajectory, extending from diagnosis of advanced disease until death. Four units provide the general principles of palliative and end-of-life care, important concepts, advanced disease management, and clinical practice guidelines. Clinical practice

guidelines offer in-depth discussions of the pathophysiology of 19 different symptoms, interventions for specific symptom management (including in-depth rationales), and suggestions for patient and family teaching. Defines dying as a normal, healthy process aided by the support of an interdisciplinary team. Provides in-depth pathophysiology, assessment, and intervention information based upon the disease trajectory. Highlights opportunities for patient and family teaching. Describes psychosocial issues experienced by patients and their families. Reviews uncomplicated and complicated grief and mourning, providing suggestions to help the family after a patient's death. Includes case studies at the end of chapters to reinforce key concepts of compassionate care. New chapters including Advance Care Planning, Ethical Issues, Spiritual Care Across Cultures, Pharmacology, Sleep, and Nutrition. Includes a new appendix on Assessment Tools and Resources for more comprehensive coverage of palliative and end-of-life care.

Palliative Care in Oncology

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.

HIV and Aging

Despite decades of attention on building a global HIV research and programming agenda, HIV in older populations has generally been neglected until recently. This new book focuses on HIV and aging in the context of ageism with regard to prevention, treatment guidelines, funding, and the engagement of communities and health and social service organizations. The lack of perceived HIV risk in late adulthood among older people themselves, as well on the part of providers and society in general, has led to a lack of investment in education, testing, and programmatic responses. Ageism perpetuates the invisibility of older adults and, in turn, renders current medical and social service systems unprepared to respond to patients' needs. While ageism may lead to some advantages – discounts for services, for example – it is the negative aspects that must be addressed when determining the appropriate community-level response to the epidemic.

CBT for Chronic Illness and Palliative Care

There is a growing awareness of the need to address the psychological distress associated with physical ill health; however, current resources are limited and difficult to access. The best way to tackle the issue is by enhancing the skills of those professionals who have routine contact with them. CBT provides the evidence-based skills that most readily meet these requirements in a time and cost efficient manner. Based on materials prepared for a Cancer Network sponsored training programme and modified to address the needs of a larger client population of people experiencing psychological distress due to physical ill-health, this innovative workbook offers a basic introduction and guide to enable healthcare professionals to build an understanding of the relevance and application of CBT methods in everyday clinical practice.

Pharmacology Case Studies for Nurse Prescribers

Nurse prescribers are now practising within every discipline of nursing and midwifery, both in the community and the hospital setting. This much-needed book is aimed particularly at students on a non-medical prescribing (NMP) course as well as those nurses who are studying for their community practitioner non-medical prescribing qualification as part of the specialist practice programme. The book is also a useful resource for nursing students and registered nurses who administer and/or prescribe medicines. The authors offer a basic introduction to pharmacological concepts, embedded in specific conditions, through case studies and self-assessment questions. By utilising a case study approach, they enable the reader to link pharmacological concepts with clinical practice. Many of the conditions presented will be seen across all healthcare settings. In addition, the comprehensive glossary and list of abbreviations will enable students to understand some of the more technical terms used in the British National Formulary (BNF) and other prescribing sources. Reading this book, and carrying out the numerous self-assessment activities, will give the reader an appreciation of the value of having a sound pharmacological knowledge base in order to deliver safe practice, effective prescribing and improved patient care. Contents include: Introduction How the body affects drugs How drugs affect the body Types of adverse drug reactions and interactions Understanding and using the British National Formulary Adherence Children Stable angina Hypertension Heart failure Chronic obstructive pulmonary disease Neurological problems Gastrointestinal disorders Incontinence in adults Diabetes Mental health problems Eye problems Complex health needs and polypharmacy Palliative care The challenges of non-medical prescribing

Dementia

This book explores how our conception of dementia has changed since its initial discovery, taking in advancements in knowledge that translate into better ways to manage the condition. Providing detailed reports of the latest research, the book explores the myriad forms of dementia. Written in accessible language, it looks at current methods of assessing and diagnosing the condition before turning to contemporary approaches to treatment. Chapters dedicated to often overlooked issues include raising awareness about how dementia affects the lives of those with an intellectual developmental disorder, the fundamental need to consider cultural differences, and the need to fully acknowledge and support informal carers. The final section of the text examines how COVID-19 has spotlighted serious gaps in healthcare for those living with dementia. Fortified with straightforward explanations and references to clinical material throughout, the book is essential reading not only for clinical psychologists in training and those in practice seeking an overview of the field and latest developments, but for a broader audience as well.

Values-Based Health & Social Care

Is evidence-based practice really best practice? This is a hotly debated question in health and social care circles and the starting point for this book. Engaging firmly in the debate, Values-Based Health & Social Care calls into question the dominance of evidence-based practice and sets out an alternative vision of care which places holism, professional judgement, intuition and client choice at its centre. Bringing together writers from a range of health and social care backgrounds, the book describes the rise of evidence-based practice and explores major criticisms of the approach. It argues that evidence should be seen as part of a broader vision of practice which places equal value on: - a holistic vision of the needs of patients and clients. - professional knowledge and intuition, and - seeing patients and clients as partners in their care. Examples are used throughout the book to help readers link the concepts to practice. The book concludes with suggestions on how to develop a values-based approach in practice and through professional education. Values-Based Health & Social Care sets out key debates surrounding the nature of practice which will be of interest to students and practitioners alike.

Dementia Care

Dementia is both a personal and a societal challenge. The goal of *Dementia Care: A Practical Approach* is to focus on how practitioners can meet this challenge with hope and compassion, thereby enabling those with dementia to live well. The book takes a 'strengths approach' with an emphasis on exploring sustainable strategies. Its content is underpinned by relevant policies and strategies and explicitly links to research evidence while always valuing the voices of those living with dementia. Covering various dementia strategies, the book provides a clear vision of dementia care delivery and is mapped to the Curriculum for UK Dementia Education. For health care students, the content is also mapped to the requirements of the Health and Care Professions Council and the Nursing and Midwifery Council. The book includes experiences of people living with dementia, practical examples, self-assessment questions, and key point summaries. It is a valuable resource to practitioners, informal caregivers, families, individuals with dementia or those wanting to know more about the subject.

Psychosocial Issues in Palliative Care

Caring for terminally ill patients and their families is challenging. Patients with life limiting illness require the skills of many professionals but also the support of their community. While most clinicians are comfortable in assessing a broad range of physical problems, it is often the psychosocial issues that prove the most complex. These issues range from psychosocial assessment to the treatment and care of patients with life limiting illnesses. Evaluating emotional, social and spiritual needs, in particular, requires excellent teamwork. This fully-updated and expanded new edition takes a comprehensive look at current practice and provision of psychosocial support as applied to a range of palliative care patients. A number of important areas are covered including community approaches of psychosocial care, neonatal palliative care, the provision of psychosocial care to families, the role of volunteers in supporting palliative care professionals, and the needs of the frail elderly, marginalised patients, and those with dementia. Including multiple case study examples, this highly practical text examines current literature and evidence to demonstrate the best research-based practice in psychosocial care. It is an essential resource for professionals working within hospitals and communities in the fields of medicine, nursing, social work, chaplaincy, counselling, primary care, and mental health.

Compassionate Communities

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.

Choices in Palliative Care

Choices in Palliative Care brings together leading experts to spotlight core issues in the field and identify ways PC can fill gaps in current care systems. This far-sighted volume redefines palliative care as interdisciplinary and integrative, bridging acute and long-term care to respond to clients' evolving needs.

Those teaching health service delivery courses will find this material especially useful.

End-of-life Nursing Care

This very accessible, straightforward book helps to allay those concerns and enables pre-registration students to prepare confidently for the challenges they will face when they are caring for dying patients and supporting their families. Each chapter is based on a different and realistic scenario - reflecting a range of circumstances - to demonstrate the essential generic knowledge and skills they need to develop, and draws out the important practical and theoretical issues students should consider and address if patients and their families are to receive the best possible care. Written by two experienced palliative care lecturer/practitioners, and mapping closely to the NMC's 2010 domains, *End-of-Life Nursing Care* is tailored to the needs of student nurses working with adult patients. Annie Pettifer and Joanna de Souza ...

Rethinking palliative care

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.

Palliative Care E-Book

Find out all you need to know about providing high-quality care to patients with serious illnesses from the 2nd edition of *Palliative Care: Core Skills and Clinical Competencies*. Drs. Linda L. Emanuel and S. Lawrence Librach, leaders in the field, address the clinical, physical, psychological, cultural, and spiritual dimensions that are integral to the care of the whole patient. They give you a broad understanding of the core clinical skills and competencies needed to effectively approach patient assessment, care of special populations, symptom control, ethical issues, and more. Clearly written in a user-friendly, high-yield format, this resource is your ultimate guidebook to the burgeoning practice of palliative medicine. Improve your pain management and symptom management skills with a better understanding of best practices in palliative care. Quickly review specific treatment protocols for both malignant and non-malignant illnesses, including HIV/AIDS, heart failure, renal failure, pulmonary disease, and neurodegenerative disease. Better understand and manage the common and unique challenges associated with delivering palliative care in various social settings, such as the ICU, hospice, and the home; and to diverse populations, such as children, elders, and vulnerable members of society. Expand your knowledge of palliative care issues with new chapters on Veterans, Special Populations, Prognostication, Delirium, Working with Families, Wound Care, Home Care, and Dealing with Economic Hardship. Find the information you need quickly and easily with a templated, high-yield format.

Psychosocial Palliative Care

One of the most challenging roles of the psycho-oncologist is to help guide terminally-ill patients through the physical, psychological, and spiritual aspects of the dying process. Patients with cancer, AIDS, and other life-threatening illnesses are at increased risk for the development of major psychiatric complications, and have an enormous burden of both physical and psychological symptoms. Concepts of adequate palliative care must

be expanded beyond the current focus on pain and physical symptom control to include the psychiatric, psychosocial, existential, and spiritual aspects of care. The psycho-oncologist, as a consultant to or member of a palliative care team, has a unique role and opportunity to fulfill this promise of competent and compassionate palliative care for those with life-threatening illnesses. Psychosocial Palliative Care guides the psycho-oncologist through the most salient aspects of effective psychiatric care of patients with advanced illnesses. This handbook reviews basic concepts and definitions of palliative care and the experience of dying, the assessment and management of major psychiatric complications of life-threatening illness, including psychopharmacologic and psychotherapeutic approaches, and covers issues such as bereavement, spirituality, cultural sensitivity, communication and psychiatric contributions to common physical symptom control. A global perspective on death and palliative care is taken throughout the text, and an Appendix provides a comprehensive list of international palliative care resources and training programs.

Resilience in Palliative Care

The first book of its kind, *Resilience and Palliative Care - Achievement in adversity* takes the increasing international literature on resilience and applies it to palliative and end-of-life care. The book offers an overview of all key aspects of palliative care, presented through a resilience perspective. Why do some patients and families break down while others surmounts the challenges facing them? What interventions strengthen individual, family and community coping? This book aims to facilitate change with people facing the crisis of death, dying and bereavement. Much of the existing literature has focused on risk, problems and vulnerability; the emerging concept of resilience focuses on strengths and possibilities. The 'total pain'/'total care' approach pioneered by Dame Cicely Saunders and St Christopher's Hospice now needs reinterpreting in the light of changing contexts and challenges. The realities of demographic change and resource-constrained health and social care environments have generated an increasingly risk focused approach to service delivery. A narrowly medicalised approach has inevitable limitations; professional care alone will be unable to meet need and demand in the face of ageing populations, changing patterns of illness and the need for equity. The resilience approach offers a counterbalance that harnesses the strengths of individuals and the communities in which they live and in which most of their dying will take place. Resilience thinking emphasises the importance of public health and creates a partnership between patients, professionals and community structures, seeking to build community capacity and to deliver a preventive health care that will leave future generations less afraid of the dying and bereavement that will confront all of us. This book offers insights into how, at all levels of planning and delivering palliative care, there is the opportunity to maximise coping, build an infrastructure for self-help, and increase the capacity of strengthened teams and organisations.

Participatory Research in Palliative Care

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.

Palliative Care

A clinical case-based handbook has a role in general clinicians' practice of caring for patients with serious or life-limiting illness. The explosion of the field of Hospice and Palliative Medicine impacts all physicians and healthcare providers. Fellowship trained specialists graduate in greater numbers annually. These and more seasoned specialists are now certified by the American Board of Medical Specialties as subspecialists. Research in this field is expanding in scope and quantity, and peer reviewed journals devoted to this work are multiplying. Moreover, peer reviewed journals in primary care and other subspecialties of medicine regularly include papers that focus on end-of-life care, quality of life issues, and symptom management. Overall then, access to clinical information necessary to care for patients with life-limiting illness is not only essential, but also increasingly available. This case-based clinical book aims to help with the actual practice of caring for patients with life-limiting illness. Numerous texts and journals exist to provide the data to inform care, yet there remains a need to find practical points and information about the practical application of the principles of palliative care. Thus, we hope that the cases, key points, and practical tips will help health care providers who are not experts already in palliative care in the care of patients with serious illness and challenging problems. Some chapters follow one patient through the course of an illness to highlight the applicability of palliative care throughout the disease process.

Palliative Care in Nursing and Healthcare

This book helps nursing and healthcare students to prepare for the challenges of working with the increasing number of patients requiring palliative care, so that they can work in partnership with patients and their carers, providing care that is compassionate, practical and backed up by the latest evidence. Delivering palliative care can be emotionally challenging and the book focuses on supporting healthcare staff, allowing them to provide the care that is needed. Key features include: * case studies in every chapter, helping students to practically work through difficult scenarios * reflective activities that assist readers in thinking critically about their care and how to improve it * a holistic approach to palliative care that includes family, carers and interprofessional work * up to date theory and policy. Palliative Care in Nursing and Healthcare is suitable for undergraduate nursing students and allied health students and practitioners. Michelle Brown is Senior Lecturer at the University of Derby.

Key Concepts in Public Health

"This is a very good text for undergraduate students as it gives a broad overview of the concept of public health, utilising case studies to illustrate practical application. This book would be also be an excellent way for practitioners to increase their own knowledge of public health and could inform their own continuing professional development." Julie Lemprière, University of Gloucestershire

Key Concepts in Public Health identifies fifty key concepts used across the discipline of public health in order to give the reader a broad perspective of the core topics relevant to training and practice. From epidemiology to health promotion, and ethics to leadership, the book offers an exciting guide to the multiprofessional field. Each entry features: - a snapshot definition of the concept - a broader discussion addressing the main issues and links to practice - key points relevant to the entry - case studies to illustrate the application to practice - examples of further reading. Highly readable, with clear indexing and cross referencing, this is an ideal book to turn to for learning more about key issues in public health practice and education. The clever structure means the book can be read in its entirety to support a programme of study or readers can use it to dip into and update their knowledge of a particular concept. It meets the validation requirements of all allied health and nursing training programmes and will also be invaluable for policy-makers and healthcare practitioners continuing their professional education.

Oxford Textbook of Public Health 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 Nursing, Fourth Edition

Print+CourseSmart

Fragility Fracture Nursing

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.

Palliative Care

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 the development of palliative 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 including: Why the care of the 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 for developing meaningful approaches that will lead to better health care for all Americans.

Death, Dying and Bereavement

The fully revised and updated edition of this bestselling collection combines academic research with professional and personal reflections. *Death, Dying and Bereavement* addresses both the practical and the more metaphysical aspects of death. Topics such as new methods of pain relief, guidelines for breaking bad news, and current attitudes to euthanasia are considered, while the mystery of death and its wider implications are also explored. A highly distinctive interdisciplinary approach is adopted, including perspectives from literature, theology, sociology and psychology. There are wide-ranging contributions from those who come into professional contact with death and bereavement - doctors, nurses, social wo

Palliative and End of Life Nursing Care

Mapped to the 2018 NMC Standards and written with the support of a student advisory board, this is the ultimate resource for developing students' confidence and competence in end of life care.

Palliative and End of Life Care in Nursing

This book is specifically designed to equip nursing students and non-specialists with essential knowledge in relation to the care and management of people nearing the end of life, covering complex issues such as handling bereavement and cultural differences.

Key Themes in Health and Social Care

Key Themes in Health and Social Care is a learning resource for students in health and social care. It provides an overview of foundational issues and core themes in the field and introduces key areas of debate, moving from an introductory level to in-depth discussion as the book progresses. Divided into three parts: the first part sets the scene, addressing introductory psychology and sociology, social policy, equality and diversity, skills for practice, and working with people the second part considers key themes such as the contribution of philosophy and politics; criminal justice; management of services; the relationship between place and wellbeing; research in health and social care; theories of counselling; housing and the built environment the third part looks at discrete areas of practice such as mental health; substance abuse, protection work; health promotion; disability studies; working with men; child welfare and public responsibility. Each chapter begins with an outline of the content and learning outcomes and includes reflective exercises to allow students to reflect on what they have read, review their learning and consolidate their understanding. Time-pressed readers wanting to 'dip into' the book for relevant areas can do so but, read from cover to cover, the book provides a comprehensive introduction to the key areas of contemporary health and social care practice. It will be particularly helpful for students undertaking health and social care undergraduate and foundation degrees.

Enhancing Compassion in End-of-Life Care Through Drama

'Healthcare professionals spend much of their time listening to stories of sickness related by patients and their families. It thus seems appropriate that drama, which is primarily concerned with exploring narratives, change and crises and relies, like the clinical situation, on communication, is an ideal medium for healthcare professionals to gain new insights into care.' From the Introduction Good communication forms the heart of

patient-centred care and is the cornerstone of a trusting relationship. *Enhancing Compassion in End-of-Life Care Through Drama* explores a broad range of plays from Greek tragedy to the present day and investigates how particular theatrical dynamics help to understand complexities in the setting of end-of-life care. It examines fresh ways to interpret the action and subtext represented on the stage and finds symmetries in a clinical context. It is ideal for use in a range of educational contexts, with practical ideas for workshops and summaries of key concepts in each chapter. This book will motivate all members of the multidisciplinary palliative care team including palliative care professionals, doctors, nurses, psychologists, spiritual advisers and social workers. Although based in the setting of palliative care, the learning points are relevant to all areas of clinical practice.

Key Concepts in Learning Disabilities

This is a handy guide to the topics you need to know about whether studying or working in the field of learning disabilities. The A-Z format allows you to “dip in” and quickly find relevant information on topics ranging from Autism to Rights and from Advocacy to Challenging Behaviors.

Continuity in Palliative Care

The challenge of providing continuity of care in an increasingly complex and technological health service is likely to continue to be problematic. *Continuity in Palliative Care* provides the practical information needed by primary care teams as they care for what is likely to be an increasing number of patients and their carers, faced with life-threatening disease and difficult treatment choices. Understanding the issues in palliative care that face primary care teams is an important concept in preparing for future challenges in the NHS and this book clearly, concisely and very readably summarises the story.

Palliative Care Nursing

“This 5th edition is an important achievement; it is a symbol of commitment to the field of palliative nursing, where we have been and where we are going.” - Betty Rolling Ferrell, PhD, MA, FAAN, FPCN, CHPN
From the Foreword The aging population has only grown since the first edition of this comprehensive and seminal publication nearly 20 years ago. Based on the need to humanize rather than medicalize the illness experience for patients, this text delves into palliative care beyond the specific diseases affecting the patient. Instead, content focuses on the whole person and family. Palliative patients struggle with chronic, debilitating, and painful conditions, and grapple with the fact that life as they knew it has already passed away. Families and friends reciprocally suffer, not knowing how to help and therefore become the secondary victims of the disease. This is not the challenge of a lone nurse, or a single physician, therapist, or social worker. Rather, palliative and hospice care requires the expertise and unique roles of an interprofessional team to help the patient and family strengthen their resilience, continue to find meaning and purpose in life, and cure what can be cured. *Palliative Care Nursing, Fifth Edition*, delivers advanced empirical, aesthetic, ethical and personal knowledge. This new edition brings an increased focus on outcomes, benchmarking progress, and goals of care. It expounds upon the importance of the cross-disciplinary collaboration introduced in the previous edition. Every chapter in Sections I, II, and III includes content written by a non-nursing member of the interprofessional team. Based on best-evidence and clinical practice guidelines, this text presents comprehensive, targeted interventions responsive to the needs of palliative and hospice patients and family. Each chapter contains compassionate, timely, appropriate, and cost-effective care for diverse populations across the illness trajectory. **Key Features** The expanded new edition offers current, comprehensive, one-stop source of highly-relevant clinical information on palliative care Life-span approach: age-appropriate nursing considerations (e.g. geriatric, pediatric and family) Includes disease-specific and symptom-specific nursing management chapters Promotes a holistic and interdisciplinary approach to palliative care Offers important legal, ethical and cultural considerations related to death and dying Case Studies with Case Study Conclusion in each clinical chapter New to The Fifth Edition: An expanded chapter on Palliative Care incorporates most up to date scope and standards, information on Basic and Advanced

HPNA certification, self-reflection and self-care for nurses. A chapter on Interprofessional Collaboration
Instructor Resources: Power points and Test bank

The Social Worker'S Guide To The Social Sciences: Key Concepts

This book provides social work students and professionals with core knowledge of the most important concepts in the social sciences. Offering a straightforward guide to diverse and complex disciplines, the book will equip and encourage you to delve further into the way societies function and individuals behave.

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