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Psychosocial Issues in Palliative Care

Oxford University Press "**Psychosocial Issues in Palliative Care is for anyone working the field of palliative care, both in the community and in hospitals; this includes those in medicine, nursing, social work, chaplaincy, counseling, primary care, and mental health.**"--Jacket.

Psychosocial Palliative Care

Oxford University Press, USA **Psychosocial Palliative Care reviews the most effective practices for the psychiatric care of patients with advanced illnesses. This handbook contains basic concepts and definitions of palliative care and the experience of dying, guidelines for the assessment and management of major psychiatric complications of life-threatening illness, including psychopharmacologic and psychotherapeutic approaches.**

Psychosocial Issues in Palliative Care

A Community Based Approach for Life Limiting Illness

Oxford University Press **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.**

Psychosocial Issues in Palliative Care

A community based approach for life limiting illness

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Palliative Care

Psychosocial and Ethical Considerations

A great number of cancer patients will suffer some form of social, emotional or psychological distress and challenges as a result of the disease and its treatment. Unattended, psychosocial issues can leave patients and families ill-equipped to cope and manage their cancer diagnosis and treatment. When psychosocial care is properly integrated into clinical care, it has a direct impact on the patients quality of life. Psychosocial aspects must be integrated into routine cancer care. The patient should be screened at their initial visit for psychosocial needs, and survivors should have a treatment plan that includes attention to possible increased anxiety on completing treatment, development of posttraumatic stress symptoms, mixed anxiety and depressive symptoms. In this book, the authors cover clinical, psychosocial and end-of-life aspects.

Good Practices in Palliative Care

A Psychosocial Perspective

Routledge A team of two practitioners in psychosocial palliative care and an academic have drawn together the work of twenty-eight highly experienced practitioners. **Good Practices in Palliative Care : a psychosocial perspective** provides detailed descriptions of innovatory practices and how they were developed, together with clear practice principles. This unique contribution to palliative care literature is suitable for a wide range of health and social care professionals at student and experienced levels and is written in a user-friendly style.

Psychosocial Palliative Care

Good Practice in the Care of the Dying and Bereaved

Nelson Thornes Based firmly on practical experience to provide students with a realistic outlook. Reflects the growing body of research in the field, stressing its significance in today's practice. Emphasises the importance of the cultural and social context in which death and bereavement take place.

Psychosocial Interventions in End-of-Life Care

The Hope for a “Good Death”

Routledge The concept of a "good death" has been hotly debated in medical circles for decades. This volume delves into the possibility and desirability of a "good death" by presenting the psychosocial measures of care as a crucial component, such as religion, existentialism, hope and meaning-making. The volume also focuses on oncologic psychiatry and the influence of technology as a means to alleviate pain and suffering, and potentially provide relief to those at the end of life. Such initiatives are aimed at diminishing pain and are socially bolstering and emotionally comforting to ensure a peaceful closure with life as opposed to a battle waged. Utilizing the most recent information from medical journals and books to present the latest on healthcare and dying today, this volume crosses the boundaries of thanatology, psychology, religion, spirituality, medical ethics and public health.

Transitions in Dying and Bereavement

A Psychosocial Guide for Hospice and Palliative Care

Find the words--and the deeds--to meet the psychosocial needs of chronically ill and dying people, their families, and caregivers in this first-ever strengths-based, step-by-step guide through the labyrinthine process from diagnosis to death to bereavement. *Transitions in Dying and Bereavement* puts a human face on a difficult yet unavoidable topic. This book comprehensively and compassionately covers the key transitions that dying people and their families face and the most effective interventions to facilitate the transitions. Employing their many years of experience in hospice and palliative care, this team of counselors and other health care professionals provides: clear explanations of current theory and research related to hospice, palliative, and bereavement care ways to help alleviate anxiety, fear, fatigue, and feelings of denial and powerlessness ways to improve communication about the experience of dying help in planning for death the Palliative Performance Scale, a functional assessment tool sensitive explanations on navigating the three phases of grief perspectives on difficult issues such as body image, sexuality, and intimacy multicultural and interdenominational perspectives on death and dying ways to support staff and much more! Activities, exercises, case studies, personal essays, poetry, and illustrations are liberally and strategically located throughout the text, forming the perfect in-service, classroom, or professional development tool for nurses, physicians, counselors, social workers, allied health professionals, volunteers, and others who work with people traversing the end-of-life experience.

Psychosocial Oncology and Palliative Care in Hong Kong

The First Decade

Hong Kong University Press As the evidence-base for clinical practice in the management of life-threatening diseases and care at the end of life increases, it is apparent that psychosocial factors play a most profound role, influencing outcomes at every level from quality of life and satisfaction with clinical services through to duration of survival and mortality. This book documents some, but by no means all, of the developments that have occurred in the past decade in the area of psychosocial oncology and palliative care in Hong Kong. Contributions describing interventions by practitioners involved in service development in nursing, social work and clinical psychology, are complemented by chapters describing academic research and theoretical perspectives. The unique cultural mix of Hong Kong is given rich emphasis in the adaptations made by practitioners and academics to the interventions and theoretical issues outlined. As both a documentation of the efforts of some of those who helped psychosocial oncology and palliative care evolve in Hong Kong, and as a reflection of the need to more critically evaluate the impact of intervention efforts in health care, this volume provides a valuable resource. Nurses, social workers, psychologists and doctors involved in delivering or planning cancer treatment or palliative care will find this book useful. This book challenges many attitudes prevalent in Hong

Kong and will, we hope, begin to break some of the taboos that continue to generate unnecessary suffering among the people cared for by our health care systems. The valuable experience documented in these pages can help others build the next generation of services to those with life-threatening illness and those at the end of life.

Palliative Care Perspectives

Oxford University Press Drawing from his extensive clinical experience and many years of teaching, Dr. Hallenbeck has written a guide to palliative care for clinicians. Topics addressed range from an overview of death and dying to specific approaches to symptom management. As an introduction to both the art and science of palliative care, this book reflects the perspectives of one physician who has dedicated his career to this rapidly evolving field. The book links real stories of illness with practical advice, thereby delineating clinical practice in a way that reflects the daily concerns of clinicians.

A Field Manual for Palliative Care in Humanitarian Crises

Oxford University Press, USA As humanitarian aid organizations have evolved, there is a growing recognition that incorporating palliative care into aid efforts is an essential part of providing the best care possible. *A Field Manual for Palliative Care in Humanitarian Crises* represents the first-ever effort at educating and providing guidance for clinicians not formally trained in palliative care in how to incorporate its principles into their work in crisis situations. Written by a team of international experts, this pocket-sized manual identifies the needs of people affected by natural hazards, political or ethnic conflict, epidemics of life-threatening infections, and other humanitarian crises. Later chapters explore topics including pain management, skin conditions, non-communicable diseases, palliative care emergencies, the law and ethics of end of life care, and more. Concise and highly accessible, this manual is an ideal educational tool pre-deployment or during fieldwork for clinicians involved in planning and providing humanitarian aid, local care providers, and medical trainees.

Psychosocial Aspects of Palliative Care

Psychosocial Issues in Palliative Care

This practical text reviews the literature and evidence on the practice and provision of psychosocial support as applied to palliative care patients. It looks at the nature of services required to provide effective psychosocial care and related issues.

Palliative Psychology

Clinical Perspectives on an Emerging Specialty

Oxford University Press *Palliative Psychology* provides clinical, evidence-based training in palliative and end of life care for clinical psychologists to accomplish specific therapeutic goals. Chapters provide a clear road map for approaching assessment and treatment by reviewing the use of psychotropic medications for patients with advanced and terminal illness, basic but important aspects of pain medication, in depth psychological and psychiatric assessment for patients with advanced illness and their caregivers, and assessment tools, highlighting the specific clinical contexts for their use. The volume also includes evidence-based psychotherapy models that have been shown effective in treating various manifestations of psychological distress in patients and caregivers. In addition to clinical topics, *Palliative Psychology* addresses crucial and often sensitive professional issues, including communication and collaboration with medical providers and issues of stress and burnout. Psychologists will learn how to best communicate the results of their assessments and treatment plan goals to other care providers in order to foster collaboration and better position themselves as advocates for their patients. Insight-oriented and practical suggestions will help clinicians manage the emotional intensity of this work and channel the therapeutic potential of their emotional reactions into their work with patients and caregivers.

Feeling better

psychosocial care in specialist palliative care

Feeling Better

Psychosocial Care in Specialist Palliative Care : a Discussion Paper

Transforming Palliative Care in Nursing Homes

The Social Work Role

Columbia University Press **This volume outlines the belief that nursing homes can and should support the physical, psychological, and social needs of residents, and that residents can thrive in nursing homes when these needs are met. The book's contributors explore the role that palliative or comfort care plays in enhancing the quality of life of nursing home residents as well as the medical, familial, psychological, cultural, and financial issues that influence decision-making about end-of-life care. The book is designed to be a tool to prepare social workers to advocate for a greater incorporation of palliative care and psychosocial care into the culture of nursing home care. The book includes discussions of the psychosocial needs of nursing home residents and families, the financing of long-term care and end-of-life care, ethical issues in chronic care and end of life, trends and characteristics in nursing home care, rituals and grief at end-of-life, and considerations for the future. Each chapter includes case examples to further illustrate points made.**

Palliative Care

Psychosocial Perspectives

Routledge

Dignity Therapy

Final Words for Final Days

OUP USA **Maintaining dignity for patients approaching death is a core principle of palliative care. Dignity therapy, a psychological intervention developed by Dr. Harvey Max Chochinov and his internationally lauded research group, has been designed specifically to address many of the psychological, existential, and spiritual challenges that patients and their families face as they grapple with the reality of life drawing to a close. In the first book to lay out the blueprint for this unique and meaningful intervention, Chochinov addresses one of the most important dimensions of being human. Being alive means being vulnerable and mortal; he argues that dignity therapy offers a way to preserve meaning and hope for patients approaching death. With history and foundations of dignity in care, and step by step guidance for readers interested in implementing the program, this volume illuminates how dignity therapy can change end-of-life experience for those about to die - and for those who will grieve their passing.**

Cancer Care for the Whole Patient Meeting Psychosocial Health Needs

National Academies Press **Cancer care today often provides state-of-the-science biomedical treatment, but fails to address the psychological and social (psychosocial) problems associated with the illness. This failure can compromise the effectiveness of health care and thereby adversely affect the health of cancer patients. Psychological and social problems created or exacerbated by cancer--including depression and other emotional problems; lack of information or skills needed to manage the illness; lack of transportation or other resources; and disruptions in work, school, and family life--cause additional suffering, weaken adherence to prescribed treatments, and threaten patients' return to health. Today, it is not possible to deliver high-quality cancer care without using existing approaches, tools, and resources to address patients' psychosocial health needs. All patients with cancer and their families should expect and receive cancer care that ensures the provision of appropriate psychosocial health services. Cancer Care for the Whole Patient recommends actions that oncology providers, health policy makers, educators, health insurers, health planners, researchers and research sponsors, and consumer advocates should undertake to ensure that this standard is met.**

Perspectives on Palliative and End-Of-Life Care Disease, Social and Cultural Context

Routledge **Perspectives on Palliative and End-of-Life Care: Disease, Social and Cultural Contexts** explores how interventions can be used to address a range of issues across social and cultural contexts for those in need of end of life care.

A Clinical Data Mining Study of the Psychosocial Status of Chinese Cancer Patients in Palliative Care

Open Dissertation Press **This dissertation, "A Clinical Data Mining Study of the Psychosocial Status of Chinese Cancer Patients in Palliative Care" by Chi-ho, Chan, 陳志豪, was obtained from The University of Hong Kong (Pokfulam, Hong Kong) and is being sold pursuant to Creative Commons: Attribution 3.0 Hong Kong License. The content of this dissertation has not been altered in any way. We have altered the formatting in order to facilitate the ease of printing and reading of the dissertation. All rights not granted by the above license are retained by the author. Abstract: Abstract of thesis entitled A Clinical Data Mining Study of the Psychosocial Status of Chinese Cancer Patients in Palliative Care Submitted by Chan Chi Ho for the degree of Doctor of Philosophy at the University of Hong Kong in August 2007 Psychosocial studies of Chinese cancer patients in palliative care of Hong Kong are limited. Little has been known on the psychosocial issues of Chinese cancer patients in palliative care of Hong Kong. As a professional who is responsible for managing the psychosocial needs of cancer patients and families in palliative care, social workers are in pressing need of local research to inform indigenous social work practice. Thus, this study attempts to answer the research questions that emerge from the social work practice in palliative care, and aims at integrating social work research with practice. Clinical Data Mining (CDM), which emphasizes the use of the existing clinical data in conducting practice-based research, is adopted as the research approach in this study. Research gaps and research questions are proposed in relation to the clinical concerns in social work practice in palliative care, and the availability of clinical data. Based on the clinical data of patients who died in 2002 to 2005, a total of 935 Chinese cancer patients in palliative care are included in this study. Findings are discussed under five major themes, including 1) moving towards good death, 2) family and patient, 3) body-mind connection, 4) search for meaning, 5) psychosocial assessment. Findings may reflect the benefits of palliative care in managing physical and psychosocial symptoms of cancer patients. Patients who were at initially higher psychosocial risk level, experienced significant improvement in most of the psychosocial symptoms from the time of admission to palliative care to final days before death. This may imply palliative care, and social work interventions in palliative care may facilitate patients to move towards good death Family shows its unique importance on the patient outcomes. Chinese cancer patients may experience a cultural phenomenon of**

"support paradox," in which patients may desire for and benefit from family support, but also worry about and are influenced by the family distress in receiving support from family. Inter-relationship between body and mind is reflected from the findings. Inevitable physical deterioration, such as the "anorexia-cachexia" syndrome, may bring psychosocial distress to patients. "Negative somatization" is conceptualized to illustrate the impact of psychosocial distress on patients' under-reporting of physical symptoms. Findings also show the universal need for meaning among Chinese cancer patients in palliative care, despite many of them are atheists. Making sense of the illness, and searching for meaning in life with terminal illness may play an important role on the patient outcomes. Importance of psychosocial assessment on admission to palliative care is also highlighted. It shows to be an essential component in understanding patients' needs, comparing differences, identifying protective and risk factors, as well as monitoring changes of patients. Implications for social work practice and social work research are discussed. Cultural sensitivity in psychosocial intervention is particularly emphasized. Recommendations are proposed to enhance service improvemen

Pediatric Psychosocial Oncology: Textbook for Multidisciplinary Care

[Springer](#) This textbook walks clinicians through the psychosocial issues and challenges faced by children and adolescents with cancer and their families. Through a developmental lens, the text provides guidance and resources that will enable clinicians to understand the physical and emotional impact of the disease from diagnosis onwards, to work with families in distress, and to diagnose and treat a range of behavioral, psychological, and psychiatric issues. The book also addresses the burgeoning fields of social media, complementary therapies, palliative care, and survivorship. Among the variety of useful resources supplied are assessment tools, websites, and additional reading materials. The psychosocial issues that arise for children and their families during the course of treatment are an important yet often overlooked aspect of pediatric oncology care. The reader will find that *Pediatric Psychosocial Oncology: Textbook for Multidisciplinary Care* covers these issues at the forefront of clinical care in a direct and approachable way, integrating research literature with practical clinical guidance.

Children's Palliative Care in Africa

[Oxford University Press, USA](#) Children's palliative care has developed rapidly as a discipline, as health care professionals recognise that the principles of adult palliative care may not always be applicable to children at the end of life. The unique needs of dying children are particularly evident across Africa, where the scale of the problem is overwhelming and the figures so enormous that they are barely comprehensible. Written by a group with wide experience of caring for dying children in Africa, this book provides practical, realistic guidance on improving access to, and delivery of, palliative care in this demanding setting. It looks at the themes common to palliative care - including communication, assessment, symptom management, psychosocial issues, ethical dilemmas, end of lifecare, and tips for the professional on compassion and conservation of energy - but always retains the focus on the particular needs of the health care professional in Africa. Whilst containing some theory, the emphasis is on practical action throughout. It will provide health care professionals working in Africa, and other resource-poor settings, with the confidence, knowledge, and capacity to improve care for the terminally ill child in constrained and demanding environments.

Identifying and Addressing the Needs of Adolescents and Young Adults with Cancer

Workshop Summary

[National Academies Press](#) *Identifying and Addressing the Needs of Adolescents and Young Adults with Cancer* is the summary of a workshop convened by the Institute of Medicine's National Cancer Policy Forum in July 2013 to facilitate discussion about gaps and challenges in caring for adolescent and young adult cancer patients and potential strategies and actions to improve the quality of their care. The workshop featured invited presentations from clinicians and other advocates working to improve the care and outcomes for the adolescent and young adult population with cancer. Cancer is the leading disease-related cause of death in adolescents and young adults. Each year nearly 70,000 people between the ages of 15 and 39 are diagnosed with cancer, approximately 8 times more than children under age 15. This population faces a variety of unique short- and long-term health and psychosocial issues, such as difficulty reentering school, the workforce, or the dating scene; problems with infertility; cardiac, pulmonary, or other treatment repercussions; and secondary malignancies. Survivors are also at increased risk for psychiatric conditions such as anxiety, depression, substance abuse, and suicide and may have difficulty acquiring

health insurance and paying for needed care. **Identifying and Addressing the Needs of Adolescents and Young Adults with Cancer** discusses a variety of topics important to adolescent and young adult patients with cancer, including the ways in which cancers affecting this group differ from cancers in other age groups and what that implies about the best treatments for these cancer patients. This report identifies gaps and challenges in providing optimal care to adolescent and young adult patients with cancer and to discuss potential strategies and actions to address them.

Psychosocial aspects of palliative care in advanced cancer

Good Practices in Palliative Care

A Psychosocial Perspective

[Ashgate Publishing](#)

Oncology

[John Wiley & Sons](#) **The Mount Sinai Expert Guides** provide quick access to point-of-care specialist clinical information for physicians across a range of therapeutic areas. For each specialty area, the guides emphasize: Common pitfalls in diagnosis and management How to prevent complications Diagnostic and treatment algorithms Key scientific trials Evidence-based clinical content Major society guidelines in order to summarize the very latest in diagnosis and management of diseases and related symptoms. Edited by renowned specialists from the Icahn School of Medicine at Mount Sinai, key faculty members provide the content for all chapters. To ensure consistency within each book and the series as a whole, chapters follow a unified structure throughout. Bonus features such as online case studies with related questions for self-assessment, key patient advice, and ICD codes are provided via the series companion website. Each guide also has its own mobile app available for purchase, allowing you rapid access to the key features wherever you may be.

Oxford Textbook of Palliative Care for Children

[Oxford University Press](#) **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.**

Handbook of Supportive Oncology and Palliative Care

Whole-Person and Value-based Care

[Springer Publishing Company](#) **Handbook of Supportive Oncology and Palliative Care** is a practical guide to providing evidence-based and value-based care to adult and pediatric cancer patients experiencing severe symptoms and stressors due to cancer diagnosis, cancer treatment, and comorbid conditions. This accessible reference provides the art and science behind the whole-person and family approach to care by delivering the best practices to relieving a cancer patient's symptoms across physical, psychosocial, and spiritual dimensions. Unlike other resources, this book covers all dimensions of palliative care but with a special emphasis on primary palliative care. Part One of the handbook provides the essential background and principles of supportive oncology and palliative care, including chapters on understanding the adult and pediatric patient and family illness experience, the roles and responsibilities of the palliative care team, and the art of the palliative care assessment interview. Part Two covers symptom management and includes ten chapters considering the major physical and psychosocial symptoms a cancer patient may face—neurologic, cardiac, respiratory, gastrointestinal, genitourinary, psychiatric, sleep and

fatigue, pain, and psychosocial and spiritual distress. Part Three addresses special considerations and issues that an oncologist, physician, nurse or other healthcare provider often face in these settings, including chapters on intimacy, sexuality, and fertility issues, grief and bereavement, running a family meeting, care for the caregiver, and survivorship. Written by expert clinicians, this state-of-the-art handbook is a necessary resource for any oncologist, nurse, primary care physician, psychosocial expert, or related practitioner who endeavors to improve quality of life and provide healing to those suffering from cancer and its treatment. Key Features: Provides the binding principles of palliative care for pediatrics, adults and families from diverse cultures and spiritual beliefs Easy-to-read format makes extracting content fast and convenient for both the clinical and educational setting Guides the clinician and practitioner through the palliative care assessment process, including the appropriate questions for the palliative care interview

Geriatric Psycho-Oncology

A Quick Reference on the Psychosocial Dimensions of Cancer Symptom Management

Apos Clinical Reference Handbo Geriatric Psycho-Oncology is a comprehensive handbook that provides best practice models for the management of psychological, cognitive, and social outcomes of older adults living with cancer and their families. Chapters cover a wide range of topics including screening tools and interventions, psychiatric emergencies and disorders, physical symptom management, communication issues, and issues specific to common cancer sites. A resource section is appended to provide information on national services and programs. This book features contributions from experts designed to help clinicians review, anticipate and respond to emotional issues that often arise in the context of treating older cancer patients. Numerous cross-references and succinct tables and figures make this concise reference easy to use. Geriatric Psycho-Oncology is an ideal resource for helping oncologists and nurses recognize when it may be best to refer patients to their mental health colleagues and for those who are establishing or adding psychosocial components to existing clinics.

Psychosocial Nursing: A Guide To Nursing The Whole Person

A Guide to Nursing the Whole Person

McGraw-Hill Education (UK) Nursing involves caring for the whole person, and taking care of both physical and psychosocial needs. This book aims to help the reader to develop the knowledge, skills and confidence to care for the whole person and to ensure the patient is at the centre of the care-giving experience. The book considers different aspects of therapeutic nursing care and explains how you can make the most of opportunities to nurse the whole person. Key topics covered include: Understanding the personal experience of illness Communication and listening skills Developing nurse-patient relationships Psychosocial assessment skills Using psychotherapeutic approaches Since it is common for those who are physically ill to feel psychologically vulnerable, the book also has chapters exploring how to care for patients who are anxious, depressed, psychotic or display challenging behaviours. All chapters include case studies and personal stories of illness to help illustrate the range of practice opportunities for improving your nursing care of the whole person. An essential read for all nurses looking to hone their craft and for those preparing to qualify who want a deeper understanding of the true art of nursing.

Pathy's Principles and Practice of Geriatric Medicine

John Wiley & Sons This new edition of the comprehensive and renowned textbook Principles and Practice of Geriatric Medicine offers a fully revised and updated review of geriatric medicine. It covers the full spectrum of the subject, features 41 new chapters, and provides up-to-date, evidence-based, and practical information about the varied medical problems of ageing citizens. The three editors, from UK, USA and France, have ensured that updated chapters provide a global perspective of geriatric medicine, as well as reflect the changes in treatment options and medical conditions which have emerged since publication of the 4th edition in 2006. The book includes expanded sections on acute stroke, dementia, cardiovascular disease, and respiratory diseases, and features a new section on end-of-life care. In the tradition of previous editions, this all-encompassing text continues to be a must-have text for all clinicians who deal with older people, particularly geriatric medical specialists, gerontologists, researchers, and general practitioners. This title is also available as a mobile App from MedHand Mobile Libraries. Buy it now from Google Play or the MedHand Store. Praise for the 4th edition: "...an excellent reference for learners at all

clinical and preclinical levels and a useful contribution to the geriatric medical literature." —Journal of the American Medical Association, November 2006 5th edition selected for 2012 Edition of Doody's Core Titles™

Handbook of Psychiatry in Palliative Medicine

Oxford University Press, USA Psychiatric, or psychosocial, palliative care has transformed palliative medicine. Palliation that neglects psychosocial dimensions of patient and family experience fails to meet contemporary standards of comprehensive palliative care. While a focus on somatic issues has sometimes overshadowed attention to psychological, existential, and spiritual end-of-life challenges, the past decade has seen an all encompassing, multi-disciplinary approach to care for the dying take hold. Written by internationally known psychiatry and palliative care experts, the Handbook of Psychiatry in Palliative Medicine is an essential reference for all providers of palliative care, including psychiatrists, psychologists, mental health counselors, oncologists, hospice workers, and social workers.

Oxford Textbook of Palliative Medicine

Oxford University Press, USA The definitive Oxford Textbook of Palliative Medicine, now in its fifth edition, has again been thoroughly updated to offer a truly global perspective in this field of extraordinary talent and thoughtfulness. Updated to include new sections devoted to assessment tools, care of patients with cancer, and the management of issues in the very young and the very old, this leading textbook covers all the new and emerging topics since its original publication in 1993. In addition, the multi-disciplinary nature of palliative care is emphasized throughout the book, covering areas from ethical and communication issues, the treatment of symptoms and the management of pain. The printed book is complemented with 12 months free access to the online version, which includes expanded chapter information and links from the references to primary research journal articles, ensuring this edition continues to be at the forefront of palliative medicine. This fifth edition of the Oxford Textbook of Palliative Medicine is dedicated to the memory of Professor Geoffrey Hanks, pioneer in the field of palliative medicine, and co-editor of the previous four editions. A truly comprehensive text, no hospital, hospice, palliative care service, or medical library should be without this essential source of information.

What Do We Mean by Psychosocial?

A Discussion Paper on Use of the Concept Within Palliative Care

Symbiotic Niceness

A Study of Psychosocial Care in Palliative Care Settings

Hospice and Palliative Care

The Essential Guide

Taylor & Francis In the past 35 years, the hospice movement has undergone major changes and has grown enormously. Palliative care is of growing importance to society as our culture struggles with how to provide compassionate end-of-life care to a growing segment of the population. This book provides professionals with a comprehensive overview of the hospice practice, as well as the challenges faced by and the future direction of the hospice movement. Chapters address the fundamentals of hospice and palliative care, including key topics such as the goals and importance of community involvement, outcome measurement, and the manner in which hospices address death, grief, and bereavement.

Also provided is a detailed analysis of the business side of hospice and offers strategies for proper management for those working in hospice programs that are growing and bringing in new staff and volunteers. This book is being published in partnership with the National Hospice and Palliative Care Organization (NHPCO).

Cancer: Palliative Care

Springer Science & Business Media **Cancer: Palliative Care** examines the nature of the care and support that can be provided to those in need of palliative care and their families. This covers not only the physical treatment, such as pain management, but also the psychological well-being of patients. Health workers, clinicians, specialist nurses and medical students will find a balanced and thoughtful overview of the subject which will be of value in managing patients and helping them to come to terms with their condition.