Life And Death Decisions Psychological And Ethical Considerations In End Of Life Care

End-of-Life-Care: A Practical Guide, Second EditionPalliative and End of Life Nursing CarePathways through Care at the End of LifeValues at the End of LifeTo Comfort AlwaysChanging the Way We DieEnd-of-life CareCase Studies in Palliative and End-of-Life CareThe Helping Professional's Guide to End-of-Life CareInnovations in End-of-life CareEnd-of-Life Care and OutcomesLiving at the End of LifeLiving with DyingA Public Health Perspective on End of Life CareEnd-of-Life Nursing CareApproaching DeathPerspectives on Palliative and End-of-Life CareImproving End of Life CareEnd-of-lifeThe Conversation Barry M. Kinzbrunner Beth Hardy Claire Henry Roi Livne Linda Norlander Fran Smith Kim K. Kuebler Margaret L. Campbell E. Alessandra Strada Mildred Z. Solomon Karl Lorenz Karen Whitley Bell Joan Berzoff Joachim Cohen Annie Pettifer Committee on Care at the End of Life Rebecca S Allen Bruce Jennings Lippincott Williams & Wilkins Angelo E. Volandes

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the most thorough text available on providing patients and families with quality end of life care the study learning questions at the end of each chapter make this book an excellent resource for both faculty who wish to test knowledge and individual learners who wish to assess their own learning the book is well written and easy to read 3 stars doody s review service end of life care a practical guide offers solution oriented coverage of the real world issues and challenges that arise daily for clinicians caring for those with life limiting illnesses and conditions end of life care a practical guide includes specific clinical guidance for pain management and other common end of life symptoms the second edition has been made even more essential with the addition of chapter ending q a for self assessment and board review new coverage of multicultural medicine an increased number of algorithms to assist decision making on complicated clinical legal and ethical issues six sections walk you through the complexities of caring for patients who are nearing the end of life preparing patients for end of life management of symptoms diagnostic and invasive interventions ethical dilemmas special populations diversity no other text better assists physicians and other clinicians in providing patients near the end of life with support guidance and hope in the face of hopelessness than end of life care a practical guide

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

this is a comprehensive guide to providing excellent end of life care it covers the pathway of care at the end of life from initial conversations about end of life care and assessment and planning to care in different settings multi agency care and providing a good death case studies and reflective questions feature throughout

this insightful study examines the deeply personal and heart wrenching tensions among financial considerations emotional attachments and moral arguments that motivate end of life decisions america s health care system was built on the principle that life should be prolonged whenever possible regardless of the costs this commitment has often meant that patients spend their last days suffering from heroic interventions that extend their life by only weeks or months increasingly this approach to end of life care is coming under scrutiny from a moral as well as a financial perspective sociologist roi livne documents the rise and effectiveness of hospice and palliative care and growing acceptance of the idea that a life consumed by suffering may not be worth living values at the end of life combines an in depth historical analysis with an extensive study conducted in three hospitals where livne observed terminally ill patients their families and caregivers negotiating treatment livne describes the ambivalent conflicted moments when people articulate and act on their moral intuitions about dying interviews with medical staff allowed him to isolate the strategies clinicians use to help families understand their options as livne discovered clinicians are advancing the idea that invasive expensive hospital procedures often compound a patient s suffering affluent educated families were more readily persuaded by this moral calculus than those of less means once defiant of death or even in denial many american families and professionals in the health care system are beginning to embrace the notion that less treatment in the end may be better treatment

focusing on the nurse s role as clinician advocate and guide this brief handbook provides a holistic and team based approach to end of life care it covers advance care planning pain management physical symptom management suffering active dying the period following the death pediatric issues cultural sensitivity and hospice norlander is a nurse and the director of the minnesota partnership to improve end of life care annotation copyrighted by book news inc portland or

there s a quiet revolution happening in the way we die more than 1 5 million americans a year die in hospice care nearly 44 percent of all deaths and a vast industry has sprung up to meet the growing demand once viewed as a new age indulgence hospice is now a 14 billion business and one of the most successful segments in health care changing the way we die by award winning journalists fran smith and sheila himmel is the first book to take a broad penetrating look at the hospice landscape through gripping stories of real patients families and doctors as well as the corporate giants that increasingly own the market changing the way we die is a vital resource for anyone who wants to be prepared to face life s most challenging and universal event you will learn hospice use is soaring yet most people come too late to get the full benefits with the age tsunami it becomes even more critical for families and patients to choose end of life care wisely hospice at its best is much more than a way to relieve the suffering of dying it is a way to live

nurses often develop long term relationships with the patients and families for whom they care providing quality care until the end of life is absolutely fundamental to nursing this important book provides the guidelines and tools necessary to provide this care publisher description case studies in palliative and end of life care uses a case based approach to provide students and practitioners with an important learning tool to improve critical thinking skills and encourage discussion toward improving experiences for patients and their families the book is organized into three sections covering subjects related to communication symptom management and family care each case is presented in a consistent logical format for ease of use highlighting key evidence based concepts including the case history care setting diagnosis and prognosis assessment treatment considerations and family support a key reference case studies in palliative and end of life care is an invaluable resource for clinicians who provide palliative care to patients with life limiting illnesses and those at the end of life along with their families

clinical psychologist e alessandra strada presents the helping professional s guide to end of life care a complete manual designed to help chaplains nurses physicians hospice workers psychotherapists palliative care specialists and psychologists address the psychological needs of terminally ill and dying patients and their families

the ahrq sponsors the development of technical assess to assist org in their efforts to improve the quality of health care it addresses the end of life which refers to a prolonged difficult period for patients caregivers nine tenths of medicare insured elderly live with a serious chronic condition before death ahrq conducted a systematic review to evaluate the scope of the end of life population outcome variables that are valid indicators of the quality of the end of life experience for the dying person surviving loved ones patient family healthcare system assoc with better or worse outcomes at end of life process interventions assoc with improved or worsened outcomes future research directions for improving end of life care illus

an updated edition of the most respected book on hospice care for both patients and caregivers this warm and informative resource on hospice and other end of life care options now gets an update it receives a new preface and revised guidance on elders who need more long term care and support recommendations on pain medications and advice for those living extended lives with treatable but not curable diseases written by a hospice nurse living at the end of life reassures us that this difficult time also offers an opportunity to explore and rediscover a richer meaning in life drawing on her years of experience bell has created a comprehensive insightful guide to every aspect of hospice care and the final stages of life for people in hospice as well as their friends and families this is an indispensable and trustworthy source of comfort and spiritual healing

the first resource on end of life care for healthcare practitioners who work with the terminally ill and their families living with dying begins with the narratives of five healthcare professionals who when faced with overwhelming personal losses altered their clinical practices and philosophies the book provides ways to ensure a respectful death for individuals families groups and communities and is organized around theoretical issues in loss grief and bereavement and around clinical practice with individuals families and groups living with dying addresses practice with people who have specific illnesses such as aids bone marrow disease and cancer and pays special attention to patients who have been stigmatized by culture ability sexual orientation age race or homelessness the book includes content on trauma and developmental issues for children adults and the aging who are dying and it addresses legal ethical spiritual cultural and social class issues as core factors in the assessment of and work with the dying it explores interdisciplinary teamwork supervision and the organizational and financing contexts in which dying occurs current research in end of life care ways to provide leadership in the field and a call for compassion insight and respect for the dying makes this an indispensable resource for social workers healthcare educators administrators consultants advocates and practitioners who work with the dying and their families

focusing on population health and discussing studies using different methodologies a public health perspective on end of life care 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

students and newly qualified staff make up much of the workforce delivering end of life care but because end of life care can be both technically challenging and emotionally demanding it is an aspect of nursing that can cause considerable anxiety 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 the book is tailored to the needs of student nurses working with adult patients it explores the importance of their role in end of life care and how this interfaces with the roles of other multidisciplinary professionals involved in the care of their patients it will also be helpful to students of other health care professions and support newly qualified health care professionals working in adult health

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

individuals and families face challenges at the end of life that can vary significantly depending on social and cultural contexts yet more than ever is now known about the needs that cut across the great diversity of experiences in the face of dying and death a number of behavioural interventions and clinical approaches to addressing these needs have been developed and are available to help providers care for clients and assist them in achieving their goals perspectives on palliative and end of life care disease social and cultural contexts explores how these interventions can be used to address a range of issues across social and cultural contexts for those in need of end of life care with perspectives from experienced clinicians providers and caregivers from around the world the book offers a strong foundation in contemporary evidence based practice alongside seasoned practice insights from the field and explores interventions for people as diverse as hiv caregivers in africa and individuals dying with dementia in addition readers will learn about the process of caring for individuals with chronic illnesses including severe mental illness weigh the impact of policy regulations on the availability of and access to palliative care and interventions and be able to compare the different issues experienced by family caregivers and formal caregivers as the companion volume to perspectives on behavioural interventions in palliative and end of life care this book will be of interest to a wide variety of individuals such as academics researchers and postgraduates in the fields of mental health medicine psychology and social work it will also be essential reading for healthcare providers and trainees from psychosocial and palliative medicine social work and nursing

this sensitively written book offers a wealth of insight and practical advice for nurses in every specialty and setting providing end of life care nurses will learn how to address patients spiritual concerns ensure that physical needs are met help patients maintain their dignity and provide emotional support to grieving families nurses will also learn how to cope with their own feelings about dying and end of life care coverage includes stages of dying nursing interventions for palliative care pain control alternative therapies physical and psychological signs of grieving and more vignette insights from the well known end of life specialist joy ufema offer advice on giving compassionate care

in this enlightening jane brody new york times book harvard medical school physician angelo e volandes offers a solution to traumatic end of life care talking medicine s oldest tool there is an unspoken dark side of american medicine keeping patients alive at any price two thirds of americans die in healthcare institutions tethered to machines and tubes at bankrupting costs even though research shows that most prefer to die at home in comfort surrounded by loved ones dr angelo e volandes believes that a life well lived deserves a good ending through the stories of seven patients and seven very different end of life experiences he demonstrates that what people with a serious illness who are approaching the end of their lives need most is not new technologies but one simple thing the conversation he argues for a radical re envisioning of the patient doctor relationship and offers ways for patients and their families to talk about this difficult issue to ensure that patients will be at the center and in charge of their medical care it might be the most important conversation you ever have

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