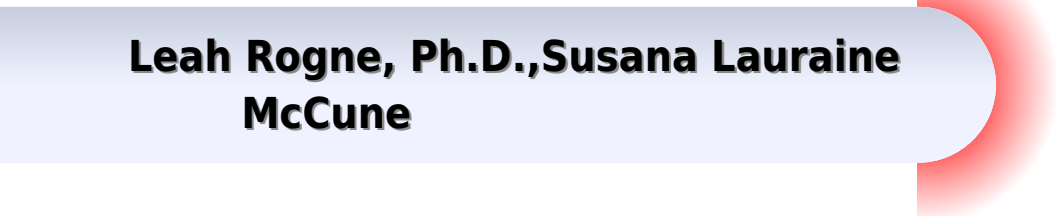




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**Leah Rogne, Ph.D., Susana Lauraine
McCune**



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The Coda Alliance Michael Brady, 1981-01-01 **Handbook of Palliative Care** Richard Kitchen, Christina Faull, Sarah Russell, Jo Wilson, 2024-05-08 Handbook of palliative care Comprehensive resource utilising up to date evidence and guidelines to support non specialists in palliative care in both hospital and community settings Building on the success of previous editions this new edition of the award winning handbook has a practical focus and provides the user with an approach to clinical challenges while also providing enough information to explain why this approach is suggested The 4th edition of Handbook of Palliative Care supports non specialists in palliative care in both hospital and community settings and focuses on holistic care and therapeutic interventions With several new chapters and content significantly updated to reflect new evidence and practice the 4th edition also presents up to date evidence guidance in a succinct format and utilises flow charts and figures to enhance the accessibility of information Written by four highly accomplished nursing and medical authors with over 100 years experience between them in hospital hospice care home and community settings Handbook of Palliative Care provides Guidance from clinicians who are experts in their field An acknowledgment of the requirements of healthcare professionals attending to patients with palliative care needs along with a dedicated chapter addressing this topic Contemporary guidance on medicine management symptom control and managing complications of cancer Palliative care in heart failure renal disease and advanced liver neurological and respiratory diseases An in depth look at patient and public involvement in palliative care and inequity Skill development including communication ethical considerations and spiritual care New chapters including frailty dementia and multi morbidity and palliative care for people living with mental illness and people with intellectual disabilities This 4th edition of Handbook of Palliative Care is an ideal supporting resource for doctors nurses and other healthcare professionals caring for patients with palliative care needs in the UK and beyond The 1st edition was the winner of the 1999 BMA Medical Book of the Year Prize *Advance Care Planning* Leah Rogne, Ph.D., Susana Lauraine McCune, 2013-07-29 Print CourseSmart Hemispheric Alliances Andrew J. Kirkendall, 2022-04-27 Hemispheric foreign policy has waxed and waned since the Mexican War and the Cold War presented both extraordinary promises and dangerous threats to U S Latin American cooperation In Hemispheric Alliances Andrew J Kirkendall examines the strengths and weaknesses of new models for U S Latin American relations created by liberal Democrats who came to the fore during the Kennedy administration and retained significant influence until the Reagan era Rather than exerting ironfisted power in Latin America liberal Democrats urged Washington to be a moral rather than a militaristic leader in hemispheric affairs Decolonization President Eisenhower s missteps in Latin America and the Cuban Revolution all played key roles in the Kennedy administration s Alliance for Progress which liberal Democrats hailed as a new cornerstone for U S Latin American foreign policy During the Vietnam War era liberal Democrats began to incorporate human rights more centrally into their agendas using Latin America as the primary arena for these policies During the long period of military dictatorship in much

of Latin America and the Caribbean liberal Democrats would see their policies dissolved by the Nixon Reagan and Bush administrations who favored militant containment of both communism and absolutism Journey'S End Victoria Brewster,Julie Saeger Nierenberg,2017-07-20 In Journey s End many and varied collaborators write about death dying and the end of life We attempt to describe real life issues and circumstances and we discuss ways to proactively deal with them Useful training resource and reference material is also included Death dying and end of life are topics many prefer to avoid This book suggests that we benefit from having frank discussions living life to the fullest and planning for our own journey s end whenever that may be Everyone who is born eventually will die whether or not we want to embrace that fact Though few of us know when we will die we and our family or friends can be well prepared We can have discussions and create written directives for what we want if we are unable to verbally state them ourselves Do we want life support Do we want interventions that may or may not have any benefit to our quality of life if we are in the hospital or in an accident Do we want to be involved in planning our funeral memorial or celebration of life The submissions within are from professionals in the field of death and bereavement support and from laypeople all of whom share stories of dying family members friends clients and patients Julie and Victoria the coauthors of this book also share stories from their personal and professional experiences Journey s End is a broadly comprehensive book about death dying and the end of life *Psychedelic Medicine at the End of Life* Richard Louis Miller,2024-11-05 Outlines 10 steps for dying gracefully with the help of psychedelics including how to navigate the complex legal landscape and find the right guide and therapy Looks at clinical studies of psychedelics from UCLA Johns Hopkins and NYU School of Medicine that show dramatic lessening of end of life anxiety in terminally ill patients Shares wisdom from experts on psychedelic research and palliative care including Roland Griffiths Katherine MacLean Ira Byock and Anthony Bossis Examining the evolving landscape that is found around end of life psychedelic care Dr Richard Louis Miller a clinical psychologist for more than half a century looks at how LSD MDMA psilocybin and ayahuasca can be vital tools in allowing individuals in all stages of life to confront fears of dying and in so doing lead richer lives Miller shares wisdom from experts on the frontiers of psychedelic research and palliative care including Roland Griffiths Katherine MacLean Ira Byock and Anthony Bossis and examines cutting edge studies from Johns Hopkins UCLA and NYU School of Medicine that show dramatically decreased anxiety in terminally ill patients through the use of psychedelics He explores how different substances can help the dying overcome their end of life distress He also provides testimony from researchers and patients participating in psychedelic assisted psychotherapy that helps convey the experience of ego death at the heart of the psychedelic experience Miller outlines 10 steps for dying gracefully without fear with the help of psychedelics He examines how to navigate the complex legal landscape and find the right guide dose and therapy He also includes reflections from key figures in the psychedelic community as well as some of his own psychedelically informed mystical and near death experiences Revealing psychedelics as a portal of transformation Miller shows how they are singularly valuable in helping

individuals face the end of life with courage and serenity *The Last Things We Talk About* Elizabeth T. Boatwright, 2021-04-06 *The Last Things We Talk About* gives readers and their loved ones the opportunity as death approaches to affirm celebrate and remember the people and experiences they cherish in life The author guides readers step by step through the process of making aging and death related decisions This includes defining personal values and wishes as well as planning for practical medical financial and legal considerations This book will help readers Identify the people experiences and things that are important to them and help define and celebrate what gives life meaning and purpose Discover and define their goals and wishes regarding transitions support and the legacy they wish to leave behind Understand important topics such as legal financial and medical documents the continuum of care and end of life decisions Find professionals to help them put together inventories for financial legal and practical matters Explore options and plan for culturally and spiritually sensitive end of life rituals and celebrations Learn what needs to be done after death and how survivors can begin to piece their lives back together

Improving Access to and Equity of Care for People with Serious Illness National Academies of Sciences, Engineering, and Medicine, Health and Medicine Division, Board on Health Sciences Policy, Board on Health Care Services, Roundtable on Quality Care for People with Serious Illness, 2019-12-06 The Centers for Disease Control and Prevention estimates that approximately 40 million people in the United States suffer from a serious illness that limits their daily activities These illnesses include heart and lung disease cancer diabetes and Alzheimer's disease and other forms of dementia However significant disparities exist across different communities in the quality and access to care for these illnesses Factors such as race ethnicity gender geography socioeconomic status or insurance status exacerbate these complex disparities It is critical to reevaluate the current models of care delivery across diverse communities and vulnerable populations On April 4 2019 The National Academies of Sciences Engineering and Medicine convened a workshop to investigate barriers policy initiatives and opportunities for improving access to and equity of care for people living with a serious illness Discussions explored the current climate of health care and opportunities to improve access to care using organizational community patient and family and clinician perspectives This publication summarizes the discussions and presentations from the workshop

Advance Care Planning in End of Life Care Keri Thomas, Ben Lobo, Karen Detering, 2017-11-28 *Advance Care Planning* ACP is an essential part of end of life care in the UK and most developed countries It enables more people to live well and die as they would choose and has significant implications for the individual person their family and carers and our wider society In the context of an ageing population and increasing possibilities for medical interventions ACP is a particularly important aspect of quality care Expanded and fully updated throughout this new edition gives a comprehensive overview of ACP and explores a wide range of issues and practicalities in providing end of life care Written by experts from around the world the book takes a comprehensive look at the subject by exploring the wide range of issues and practicalities in providing ACP framing the purpose process and outcomes of these

plans and providing an important update on national and international research policy and practice. Chapters also discuss values, goals, and priorities and include detailed case examples to aid best practice. This book is an invaluable resource for all clinicians involved in the caring for people in their final stages of life. It is of particular value to GPs, palliative care specialists, geriatricians, social care teams, researchers, and policy leads interested in improving end of life care.

Values at the End of Life

Roi Livne, 2019-06-10

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.

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