

The Psychosocial Aspects Death Dying

Dr. Irmo Marini, Dr. Mark A. Stebnicki

The Psychosocial Aspects of Death and Dying, 2nd Edition John D. Canine, 2019-05

The Psychosocial Aspects of Death and Dying John Canine, 1996-10-21 This book provides an insightful and concise approach to the psychosocial process of death and bereavement.

Top Five Regrets of the Dying Bronnie Ware, 2019-08-13 Revised edition of the best-selling memoir that has been read by over a million people worldwide with translations in 29 languages. After too many years of unfulfilling work, Bronnie Ware began searching for a job with heart. Despite having no formal qualifications or previous experience in the field, she found herself working in palliative care. During the time she spent tending to those who were dying, Bronnie's life was transformed. Later, she wrote an Internet blog post, outlining the most common regrets that the people she had cared for had expressed. The post gained so much momentum that it was viewed by more than three million readers worldwide in its first year. At the request of many, Bronnie subsequently wrote a book, *The Top Five Regrets of the Dying*, to share her story. Bronnie has had a colourful and diverse life. By applying the lessons of those nearing their death to her own life, she developed an understanding that it is possible for everyone, if we make the right choices, to die with peace of mind. In this revised edition of the best-selling memoir that has been read by over a million people worldwide, with translations in 29 languages, Bronnie expresses how significant these regrets are and how we can positively address these issues while we still have the time. *The Top Five Regrets of the Dying* gives hope for a better world. It is a courageous, life-changing book that will leave you feeling more compassionate and inspired to live the life you are truly here to live.

Death Attitudes and the Older Adult Adrian Tomer, 2018-10-24 This innovative and informative new text bridges the fields of gerontology and thanatology.

Psychosocial Issues in Palliative Care Mari Lloyd-Williams, 2008-05-08 *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.

The Psychological and Social Impact of Illness and Disability Irmo Marini, PhD, DSc, CRC, CLCP, Mark A. Stebnicki, PhD, LPC, DCMHS, CRC, CCM, CCMC, 2017-12-28 Praise for the Sixth Edition: Adds an important international perspective on illness and disability. The personal narratives help bring the real world of people who are [survivors] to the forefront of the

scientific discourse. —Doody's Medical Reviews Now in its seventh edition, this bestselling classic continues to be the most comprehensive and diverse text available on the psychosocial aspects of illness and disability. It is substantially revised to reflect the growing disparity between the haves and the have-nots and incorporates social justice issues throughout the text. In addition to new and updated information integrated throughout the book, the seventh edition features two new chapters addressing social justice in regards to depression and disability, and the psychosocial aspects of grief, death, and dying. Additionally, the text now includes an Instructor's Manual and PowerPoint slides. Combining a mix of seminal work from rehabilitation counseling legends with current theoretical and treatment approaches, the book provides a practical, real-life perspective and offers broad and inclusive coverage of the day-to-day challenges of working with a diverse and marginalized population. Additionally, the text analyzes barriers to enabling patients with disabilities and improving their quality of life. Chapter objectives, review questions, and personal narratives in each chapter facilitate in-depth learning. New to the Seventh Edition: Completely updated to incorporate social justice issues, from the medical and psychosocial aspects of combat trauma to the impact of mental and physical disabilities on immigrants, refugees and asylum-seekers, throughout Includes two new chapters addressing Social Justice/Depression and Disability and the Psychosocial Aspects of Grief, Death, and Dying Includes an Instructor's Manual and PowerPoint slides Enhanced coverage of topics concerning diverse and marginalized populations, including Women with Disabilities, Sexuality and Disabilities, LBGTQ Issues, Aging with Disabilities, Trauma, and more Key Features: Presents the most comprehensive and diverse coverage of psychosocial aspects of disability of any text Emphasizes the negative impact of societal attitudes and treatment of disabled individuals on their psychological adjustment to disability Examines both seminal and current thinking and treatment approaches Provides a bridge between theory and practice with abundant narratives Includes objectives and reviews questions in each chapter

The Psychological and Social Impact of Illness and Disability, 6th Edition Dr. Irmo Marini, Dr. Mark A. Stebnicki, 2012-02-24 Print+CourseSmart

Psychology of Death and Dying John C. Morgan, Richard L. Morgan, 2006-04-01 Sooner or later each one of us faces death, our own or others we care about. And yet, few take time beforehand to think about these endings, and in the process may lose the wisdom of the ages that comes after facing death. Perhaps this explains why when Plato was asked to summarize his philosophy he reportedly said: Practice dying. He understood that dying is what each one of us does throughout our lives, whether it is leaving home the first time to go to school or departing from this planet when our lives end. If we learn how to die--to let go and get our egos out of the way--we will have gained wisdom about how best to live. This brief yet comprehensive book deals not only with the philosophical and psychological meaning of death but its practical implications for our lives. Written by two brothers who have taught philosophy, ethics, psychology, and religion at community colleges and four-year private colleges, this book can be used in many learning situations, whether part of courses in

philosophy, ethics, psychology, or counseling; or for short-term workshops or continuing education courses for students in human services, health care, social work, or any of the helping professions.

The Psychological and Social Impact of Chronic Illness and Disability Irmo Marini, PhD, DSc, CRC, CLCP, Allison R. Fleming, PhD, CRC, Malachy Bishop, PhD, CRC, 2023-08-09 Reflects the most significant and fundamental shifts in the experience of disability in human history With the release of its eighth edition, this bestselling text remains the most comprehensive and current text addressing the psychological and social issues dealt with by persons with disabilities. The new edition is almost completely rewritten and expanded by expert voices in disability and rehabilitation policy, research, and lived experience. It presents many new chapters covering topics such as disability identity, the impact of US laws and policies, the impact of micro-aggressions and discrimination, applications of well-being and positive psychology, and mental health implications of social media usage for people with disabilities. The eighth edition also includes new Personal Perspectives from individuals with various disabilities. The text provides an informed, critical, and engaging exploration of the impact of chronic illness and disability (CID) for a wide range of students, educators, and professionals who work with this population. It delivers a comprehensive understanding of CID topics ranging from the impact of law and policies, social justice issues, personal and professional rehabilitation, and the psychosocial experiences of CID. The book continues to investigate a diverse range of topics, from the historical and cultural perspectives on illness and disability to the personal, familial, and social impacts of disability. Chapters include Learning Objectives, Pre-reading Questions, Class Activities, and Case Studies with accompanying Discussion Questions to promote engagement. Instructors will also have access to the Instructor Manual, Test Bank, and chapter PowerPoints. Extensively rewritten and updated with expert voices in disability and rehabilitation policy and research Brand new chapters on disability identity, the impact of US laws and policies, the impact of micro-aggressions and discrimination, and more New Personal Perspectives from persons who have lived with various disabilities New Class Activities to help reinforce content Key Features: Presents the most comprehensive and diverse coverage of psychosocial aspects of disability of any text Addresses how people with CID have been viewed and treated throughout history and examines the changes and developments over the past decade Provides a bridge between theory and practice with abundant narratives Includes Learning Objectives, Pre-reading Questions, and Case Studies with Discussion Questions to enhance learning Delivers a comprehensive instructor package including Instructor Manual, Test Bank, and chapter PowerPoints

Psychosocial Aspects of Pediatric Oncology Shulamith Kreitler, Myriam Weyl Ben Arush, 2004-08-13 This book is a comprehensive text on the all-important psychosocial aspects of cancer in children. Edited by an experienced psycho-oncologist and an equally experienced pediatric oncologist, the book brings together an international group of contributors composed of pediatric oncologists and psychologists/psycho-oncologists. This unique balance of contributors gives the book a

focus on the real-life practical aspects of children undergoing treatment for cancer. The book helps health care professionals, who look after children and adolescents with cancer, in dealing with the difficult and complex problems that face the child, his siblings and his parents. It deals with critical issues such as the neuropsychological impact of treatment, (including bone marrow transplantation), pain, quality of life in survivors, palliative care, talking to children and their families about death, and providing support to the dying. An important part of the book provides tools for evaluation and methods of psychological intervention. It is thus a very practical work for psycho-oncologists, pediatric oncologists, pediatricians, psychologists, nurses, social workers and art therapists dealing with the problems that children with cancer have to confront. A comprehensive book on the clinical approach to psychosocial problems of children with cancer Includes evaluation tools and psychological intervention techniques Contributions from both pediatric oncologists and psycho-oncologists highlight the team approach to the treatment of childhood cancers

In the Face of Death Danai Papadatou, PhD, 2009-04-06 Occasionally I read a book and say to myself that this is a book that I wished I had written. This is such a book! It is a delightful, practical, interesting, and inspiring book! Danai has written a soon-to-be classic in the field! Her writing is excellent! Her use of theories, concepts, history, and models are wonderful!--
Illness, Crisis and Loss Danai Papadatou [presents] an approach of Relational Care, care based on an understanding of relationships, that should be essential reading. [S]he writes in an engaging and non-technical language, and manages to convey complex ideas in a manner that is accessible to all. --Colin Murray Parkes, OBE, MD, FRCPsych President, *Cruse: Bereavement Care* (From the Foreword) [A] wonderful contribution to the literature on caregiving and her well thought out ideas about relational caregiving are on the cutting edge. Congratulations. -- Lynne Ann DeSpelder CoAuthor, *The Last Dance: Encountering Death and Dying*, 8th edition This is no ordinary academic textbook. It is nothing short of a masterpiece in which Papadatou provides the field of death, dying and bereavement with invaluable and constructive insights into the world of caring for the dying and bereaved.--*Grief Matters* This is a fascinating book, applying important theoretical models in order to describe and speculate about how professionals manage to work in an environment where suffering and grief are constantly present. This is an important and substantial addition to the mostly self-help literature about self-care for caregivers.--*Doody's Medical Reviews* *In the Face of Death* explores the experiences of health care professionals who care for the seriously ill, the dying, and the bereaved. In this book, Danai Papadatou offers a practical approach to caregiving, as well as a breadth and depth of insight into both the patient's and the caregiver's responses to death. The author discusses the issues and challenges health care professionals face when treating dying and bereaved patients. Topics include: compassion fatigue, the inevitability of suffering and the potential for growth, suffering in the workplace, team functioning in death situations, and team resilience. The main themes are: The Caring Relationship focuses on the relationship between the care provider and the person who is dying or grieving, and proposes a new, relationship-based model of care The Care Provider in

Death Situations addresses the health professional's personal responses to death, using a model that illustrates the grieving process of the health professional. *The Team in the Face of Death* provides recommendations for effective, interdisciplinary care services that support dying or bereaved patients as well as the health care provider.

Towards Death with Dignity Sylvia Poss, 2021-11-07 The explosion of literature on the once taboo topic of death and dying in the late 1970s had tended to pass the professional social worker by. Originally published in 1981, it was to fill this important gap that *Towards Death with Dignity* was written. Not since Kubler-Ross's now classic *On Death and Dying* has a book in the field of terminal care been informed by so much first-hand experience, and so much case material, allowing the caregiver to learn from the dying person himself how best to help him towards a dignified death. Sylvia Poss's sensitive elucidation of what the dying person must do for himself in order to master his terminal crisis was welcomed as a major contribution to psychosocial knowledge at the time. Having outlined the dying person's side of the crisis, she turns to the perspective of those who hope to help him towards death - other patients, nurses, doctors, paramedical staff and social workers, chaplains, volunteers, employers, relatives and friends. *Towards Death with Dignity* focuses on three of social work's major methods: social casework, community work and teaching. Not only does Sylvia Poss outline what may need to be done by the caregiver, but she also illustrates how; she further outlines how to prepare for social work in the terminal care field and suggests an effective method for teaching terminal care skills. Her book also provided, for the first time, a synthesis of other recent work in the field, to help social workers through what had become a plethora of specialist psychosocial and medical literature. *Towards Death with Dignity* was thus a useful, practical guide, both for laymen and for the many professionals involved in this aspect of the health care field. It will also be valuable for those who are involved personally in moving towards their own death, or are being called upon to be involved in some way in the death of a relative, neighbour or friend.

Psychosocial Aspects of Oncology Jimmie C. Holland, Robert Zittoun, 2012-12-06 The European School of Oncology came into existence to respond to a need for information, education and training in the field of the diagnosis and treatment of cancer. There are two main reasons why such an initiative was considered necessary. Firstly, the teaching of oncology requires a rigorously multidisciplinary approach which is difficult for the Universities to put into practice since their system is mainly disciplinary orientated. Secondly, the rate of technological development that impinges on the diagnosis and treatment of cancer has been so rapid that it is not an easy task for medical faculties to adapt their curricula flexibly. With its residential courses for organ pathologies and the seminars on new techniques (laser, monoclonal antibodies, imaging techniques etc.) or on the principles of chemotherapy (conservative or mutilating surgery, primary or adjuvant chemotherapy, radiotherapy alone or integrated), it is the ambition of the European School of Oncology to fill a cultural and scientific gap and, thereby, create a bridge between the University and Industry and between these two and daily medical

practice. One of the more recent initiatives of ESO has been the institution of permanent study groups, also called task forces, where a limited number of leading experts are invited to meet once a year with the aim of defining the state of the art and possibly reaching a consensus on future developments in specific fields of oncology.

Death and Dying Glennys Howarth, 2007-01-16 Glennys Howarth provides a cutting-edge, comprehensive discussion of the key topics in death and dying and in so doing demonstrates that the study of mortality is germane to all areas of sociology. The book is organized thematically, utilizing empirical material from cross-national and cross-cultural perspectives. It carefully addresses questions about social attitudes to mortality, the social nature of death and dying, and explanations for change and diversity, and explores traditional and contemporary experiences of death.--Jacket.

Gerotranscendence Lars Tornstam, PhD, 2005-06-20 Given the 2006 GREAT GERONTOLOGY AWARD for outstanding contribution to gerontological research by the Swedish Gerontological Society Received a VALUE GROUND AWARD from the journal *Aldreomsorg* (Old Age Care) Expanding upon his earlier writings, Dr. Tornstam's latest book explores the need for new theories in gerontology and sets the stage for the development of his theory of gerotranscendence. This theory was developed to address what the author sees as a perpetual mismatch between present theories in social gerontology and existing empirical data. The development towards gerotranscendence can involve some overlooked developmental changes that are related to increased life satisfaction, as self-described by individuals. The gerotranscendent individual typically experiences a redefinition of the Self and of relationships to others and a new understanding of fundamental existential questions: The individual becomes less self-occupied and at the same time more selective in the choice of social and other activities. There is an increased feeling of affinity with past generations and a decreased interest in superfluous social interaction. The individual might also experience a decrease in interest in material things and a greater need for solitary meditation. Positive solitude becomes more important. There is also often a feeling of cosmic communion with the spirit of the universe, and a redefinition of time, space, life and death. Gerotranscendence does NOT imply any state of withdrawal or disengagement, as sometimes erroneously believed. It is not the old disengagement theory in new disguise. Rather, it is a theory that describes a developmental pattern beyond the old dualism of activity and disengagement. The author supports his theory with insightful qualitative in-depth interviews with older persons and quantitative studies. In addition, Tornstam illustrates the practical implications of the theory of gerotranscendence for professionals working with older adults in care settings. A useful Appendix contains suggestions of how to facilitate personal development toward gerotranscendence. For Further Information, Please Click Here!

Dying to be Men Will Courtenay, 2011-05-09 Masculinity has a powerful effect on the health of men and boys. Indeed, many of the behaviors they use to be men actually increase their risk of disease, injury, and death. In this book, Dr. Will Courtenay, an internationally recognized expert on men's health, provides a foundation for understanding this troubling

reality. With a comprehensive review of data and literature, he identifies specific gender differences in the health-related attitudes, beliefs, and behaviors of men and boys and the health consequences of these differences. He then describes the powerful social, environmental, institutional, and cultural influences that encourage their unhealthy behaviors and constrain their adoption of healthier ones. In the book's third section, he more closely examines the health needs of specific populations of men, such as ethnic-minority men, rural men, men in college, and men in prisons. Courtenay also provides four empirical studies conducted with multidisciplinary colleagues that examine the associations between masculinity and men and boys' health beliefs and practices. Finally, he provides specific strategies and an evidence-based practice guideline for working with men in a variety of settings, as well as a look to the future of men's health. Medical professionals, social workers, public health professionals, school psychologists, college health professionals, mental health practitioners, academics, and researchers from a broad array of disciplines, and anyone interested in this topic will find it to be an extensively researched and accessible volume.

Dying in America Institute of Medicine, Committee on Approaching Death: Addressing Key End-of-Life Issues, 2015-03-19 For patients and their loved ones, no care decisions are more profound than those made near the end of life. Unfortunately, the experience of dying in the United States is often characterized by fragmented care, inadequate treatment of distressing symptoms, frequent transitions among care settings, and enormous care responsibilities for families. According to this report, the current health care system of rendering more intensive services than are necessary and desired by patients, and the lack of coordination among programs increases risks to patients and creates avoidable burdens on them and their families. *Dying in America* is a study of the current state of health care for persons of all ages who are nearing the end of life. Death is not a strictly medical event. Ideally, health care for those nearing the end of life harmonizes with social, psychological, and spiritual support. All people with advanced illnesses who may be approaching the end of life are entitled to access to high-quality, compassionate, evidence-based care, consistent with their wishes. *Dying in America* evaluates strategies to integrate care into a person- and family-centered, team-based framework, and makes recommendations to create a system that coordinates care and supports and respects the choices of patients and their families. The findings and recommendations of this report will address the needs of patients and their families and assist policy makers, clinicians and their educational and credentialing bodies, leaders of health care delivery and financing organizations, researchers, public and private funders, religious and community leaders, advocates of better care, journalists, and the public to provide the best care possible for people nearing the end of life.

Encyclopedia of Behavioral Medicine Marc D. Gellman, J. Rick Turner,

Encyclopedia of Death and Dying Glennys Howarth, Oliver Leaman, 2003-12-16 In recent years there has been a massive upsurge in academic, professional and lay interest in mortality. This is reflected in academic and professional literature, in

the popular media and in the proliferation of professional roles and training courses associated with aspects of death and dying. Until now the majority of reference material on death and dying has been designed for particular disciplinary audiences and has addressed only specific academic or professional concerns. There has been an urgent need for an authoritative but accessible reference work reflecting the multidisciplinary nature of the field. This Encyclopedia answers that need. The Encyclopedia of Death and Dying consolidates and contextualizes the disparate research that has been carried out to date. The phenomena of death and dying and its related concepts are explored and explained in depth, from the approaches of varied disciplines and related professions in the arts, social sciences, humanities, medicine and the sciences. In addition to scholars and students in the field—from anthropologists and sociologists to art and social historians - the Encyclopedia will be of interest to other professionals and practitioners whose work brings them into contact with dying, dead and bereaved people. It will be welcomed as the definitive death and dying reference source, and an essential tool for teaching, research and independent study.

Dealing with Dying, Death, and Grief during Adolescence David E. Balk, 2014-02-18 For some, life's introduction to death and grief comes early, and when it does it can take many forms. Not only does *Dealing with Dying, Death, and Grief during Adolescence* tackle them all, it does so with David Balk's remarkable sensitivity to and deep knowledge of the pressures and opportunities adolescents face in their transition from childhood to adulthood. In seamless, jargon-free language, Balk brings readers up to date with what we know about adolescent development, because over time such changes form the backstory we need to comprehend the impact of death and bereavement in an adolescent's life. The book's later chapters break down the recent findings in the study of life-threatening illness and bereavement during adolescence. And, crucially, these chapters also examine interventions that assist adolescents coping with these difficulties. Clinicians will come away from this book with both a grounded understanding of adolescent development and the adolescent experience of death, and they'll also gain specific tools for helping adolescents cope with death and grief on their own terms. For any clinician committed to supporting adolescents facing some of life's most difficult experiences, this integrated, up-to-date, and deeply insightful text is simply the book to have. David E. Balk is professor in the department of health and nutrition sciences at Brooklyn College (CUNY), where he directs the graduate program in thanatology. He is the author of *Adolescent Development: Early Through Late Adolescence*, *Helping the Bereaved College Student*, and several other books on death and bereavement. He is also co-editor of the 2nd edition of the *Handbook of Thanatology* (Routledge, 2013).

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