

# Crossing Over

Narratives of  
Palliative Care



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# Crossing Over Narratives Of Palliative Care

**David Barnard, Anna M.  
Towers, Patricia Boston, Yanna  
Lambrinidou**



## **Crossing Over Narratives Of Palliative Care:**

*Crossing Over* David Barnard, Anna M. Towers, Patricia Boston, Yanna Lambrinidou, 2000-03-02 *Crossing Over* provides a unique view of patients families and their caregivers striving together to maintain comfort and hope in the face of incurable illness The narratives weave together emotions physical symptoms spiritual concerns and the stresses of family life as well as the professional and personal challenges of providing hospice and palliative care Based on a vast amount of participant observation and in depth interviews *Crossing Over* moves far beyond dry technical manuals for symptom control and tired clichés about death with dignity to depict the sights sounds tastes and smells of the daily in patients homes and the palliative care unit It captures the breathtaking diversity of people's aspirations and ideals as they face death and the views of the professionals who care for them Anger and fear tenderness and reconciliation jealousy and love social support and falling through the cracks unexpected courage and unshakable faith all of these are part of facing death in late twentieth century North America and this book brings them to life in an extraordinary portrait of the processes of giving and receiving palliative care

**Crossing Over** David Barnard, Anna Towers, Patricia Boston, Yanna Lambrinidou, 2023 The revised edition of *Crossing Over* is a book of stories narratives of giving and receiving palliative care in the context of end of life It is not a textbook that portrays ideal palliative care or that prescribes specific management techniques Instead it presents stories of actual patients and families who have experienced terminal illness with the support of hospice or palliative care teams The narratives are derived from a three year qualitative ethnographic study of the experiences of patients families and caregivers

**Crossing Over** David Barnard, 2000 *Crossing Over* David Barnard, Anna Towers, Patricia Boston, Yanna Lambrinidou, 2023 The revised edition of *Crossing Over* is a book of stories narratives of giving and receiving palliative care in the context of end of life It is not a textbook that portrays ideal palliative care or that prescribes specific management techniques Instead it presents stories of actual patients and families who have experienced terminal illness with the support of hospice or palliative care teams The narratives are derived from a three year qualitative ethnographic study of the experiences of patients families and caregivers

Crossing Over David Barnard, Anna Towers, Patricia Boston, Yanna Lambrinidou, 2023 Twenty richly detailed narratives vividly bring to life the experiences of dying and bereavement in *Crossing Over* weaving together emotions physical symptoms spiritual concerns and the stresses of family life as well as the professional and personal challenges of providing hospice and palliative care

**Crossing Over**, 2000 This volume presents a series of case narratives following individual patients and families throughout the course of illness and death in the context of hospice and palliative care

**Transforming the Culture of Dying** David Clark, 2013-06-13 *Transforming the Culture of Dying* assesses the establishment of the Project on Death in America and evaluates its contributions to the development of the palliative care field and end of life care in American society

**The New Testament and Bioethics** R. Dennis Macaleer, 2014-01-01 Beauchamp and Childress's *Principles of Biomedical Ethics* is a well accepted approach to

contemporary bioethics Those principles are based on what Beauchamp and Childress call the common morality This book employs New Testament theological themes to enhance the meaning of those principles of bioethics The primary New Testament text for this study is the twin commands from Jesus to love God and love one's neighbor The three theological themes developed from this study the image of God the covenant and the pursuit of healing are deeply embedded in the New Testament and in the ministry of Jesus Three contemporary bioethics principles are used for this dissertation based on The Belmont Report They are the principles of respect for persons justice and beneficence In each case the theological themes are shown to enhance the meaning of these bioethics principles Each of the three principles as understood through the three theological themes is applied to a current bioethics issue to demonstrate the efficacy of this approach The three current issues addressed are the withdrawal or withholding of life sustaining treatment the distribution of health care in the United States and the use of palliative care

**Handbook of Psychiatry in Palliative Medicine** Harvey Max Chochinov, William Breitbart, 2009 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

*Communication as Comfort* Sandra L. Ragan, Elaine M. Wittenberg-Lyles, Joy Goldsmith, Sandra Sanchez Reilly, 2008-05-15 This exceptional work explores the complexities of communication at one of the most critical stages of the life experience during advanced serious illness and at the end of life Challenging the predominantly biomedical model that informs much communication between seriously ill and or dying patients and their physicians caregivers and families Sandra L Ragan Elaine M Wittenberg Lyles Joy Goldsmith and Sandra Sanchez Reilly pose palliative care medical care designed to comfort rather than to cure patients as an antidote to the experience of most Americans at the most vulnerable juncture of their lives With an author team comprised of three health communication scholars and one physician certified in geriatrics and palliative medicine this volume integrates the medical literature on palliative care with that of health communication researchers who advocate a biopsychosocial approach to health care Applying communication theories and insights to illuminate problems and to explain their complexities the authors advocate a patient centered approach to care that recognizes and seeks to lessen patients suffering and the many types of pain they may experience physical psychological social and spiritual during life threatening illness

*Oxford Textbook of Palliative Medicine* Geoffrey W. C. Hanks, 2011-07-21 Emphasising the multi disciplinary nature of palliative care the fourth edition of this text also looks at the individual professional roles that contribute to the best quality palliative care

*Death and Dying* John F. Szabo, 2009-11-02 While it can be traced to the 1700s the term thanatology has gained traction in recent decades to refer to the study of death. Because of the breadth of the published material and the challenges of discovery in large bibliographic databases and catalogs finding and identifying monographic material on death, dying, grief and bereavement can be extremely challenging. Although there have been several bibliographical resources published on thanatology, those titles were incomplete, limited in scope or leaned heavily toward one area of the field. In *Death and Dying: An Annotated Bibliography of the Thanatological Literature*, John F. Szabo provides more than 2,200 citations of monographs on the science and study of death and dying. Among the areas this volume addresses are the psychological, philosophical and attitudinal aspects of coping and dealing with the burdens of caregiving and working in the helping professions; instructional and educational topics for practitioners, primarily health care providers; cultural differences in bereavement, rituals and grief; mourning and loss. In addition, notable titles on or relating to death and dying in popular culture, death themes in literature, methods of death or specific ethical policy or public issues are also included when they contribute important information to subject areas on which the book focuses. This book will be helpful to students, researchers, academics, caregivers, health care professionals, psychologists, social workers and anyone with an interest in death, dying, bereavement or the care of the terminally ill.

*Making Sense Of: Health, Illness and Disease*, 2021-10-25 Health, illness and disease are topics well suited to interdisciplinary inquiry. This book brings together scholars from around the world who share an interest in and a commitment to bridging the traditional boundaries of inquiry. We hope that this book begins new conversations that will situate health in broader socio-cultural contexts and establish connections between health, illness and disease and other socio-political issues. This book is the outcome of the first global conference on Making Sense of Health, Illness and Disease held at St Catherine's College, Oxford in June 2002. The selected papers pursue a range of topics from the cultural significance of narratives of health, illness and disease to healing practices in contemporary society as well as patients' illness experiences. Researchers and health care practitioners now live in the age of interdisciplinarity which has transformed both health care delivery and research on health. The essays in this collection transcend the traditional boundaries of biomedicine and draw attention to the many ways in which health is embedded in socio-cultural norms and how these norms in turn shape health practices and health care. This volume is of interest not only to researchers but also to those delivering health care.

*Participatory Action Research in Health Care* Tina Koch, Debbie Kralik, 2009-02-18 Participatory Action Research in Healthcare provides a guide to participatory action research in the community health setting. It draws upon the authors' experiences working, researching and engaging with people utilising collaborative participatory approaches. The authors position participatory action research as a vital, dynamic and relevant approach that can be engaged by practitioners and health service providers. It is argued that participating with people is the way to move forward towards sustainable services that evoke human flourishing. *Participatory Action Research in Healthcare* explores the key issues surrounding participatory

action research and examines the benefits of this approach for community development and health promotion. It includes detailed guidelines on data generation and analysis.

**Communicating at the End of Life** Elissa Foster, 2014-04-08 This enlightening volume provides first hand perspectives and ethnographic research on communication at the end of life a topic that has gone largely understudied in communication literature. Author Elissa Foster's own experiences as a volunteer hospice caregiver form the basis of the book. *Communicating at the End of Life* recounts the stories of Foster and six other volunteers and their communicative experiences with dying patients using communication theory and research findings to identify insights on the relationships they form throughout the process. What unfolds is a scholarly examination of a subject that is significant to every individual at some point in the life process. Organized chronologically to follow the course of Foster's involvement with hospice and the phases of the study, the book opens with Part 1 providing background and contextual information to help readers understand subsequent stories about communication between volunteers and patients. Part 2 of the volume emphasizes the adjustments required by the volunteers as they entered the world of hospice and the worlds of the patients. Part 3 underscores the importance of improvisation and finding balance within the role of volunteer in particular how to be fully present for patients as well as their family members. The volume concludes with Part 4 which addresses how volunteers coped with the death of their patients and what they learned from the experience of volunteering. *Communicating at the End of Life* is appropriate for scholars and advanced students studying personal relationships, health communication, gerontology, interpersonal communication, lifespan communication and communication aging. Its unique content offers precious and meaningful insights on the communication processes at a critical point in the life process.

**The Edge of Medicine** David J. Bearison, Linda Granowetter, 2012-09-27 *The Edge of Medicine* tells the stories of dying children and their families capturing the full range of uncertainties, hopes and disappointments and ups and downs of children near the end of life. Dr. Bearison relies on narrative to bridge the disconnect among abstract theories, medical technologies and clinical realities.

**Eros and Illness** David B. Morris, 2017-02-27 When we or our loved ones fall ill, our world is thrown into disarray, our routines are interrupted, our beliefs shaken. David Morris offers an unconventional, deeply human exploration of what it means to live with and live through disease. He shows how desire, emotions, dreams, stories, romance, even eroticism plays a crucial part in illness.

Death, Society, and Human Experience Robert Kastenbaum, Christopher M. Moreman, 2018-03-29 Providing an overview of the myriad ways that we are touched by death and dying both as an individual and as a member of society, this book will help readers understand our relationship with death. Kastenbaum and Moreman show how various ways that individual and societal attitudes influence both how and when we die and how we live and deal with the knowledge of death and loss. This landmark text draws on contributions from the social and behavioral sciences as well as the humanities such as history, religion, philosophy, literature and the arts to provide thorough coverage of understanding death and the dying process. *Death, Society and Human Experience* was originally written by Robert

Kastenbaum a renowned scholar who developed one of the world's first death education courses Christopher Moreman who has worked in the field of death studies for almost two decades specializing in afterlife beliefs and experiences has updated this edition     Caring for the Elderly in Japan and the US Susan Orpett Long, 2013-01-11 In an era of changing demographics and values this volume provides a cross national and interdisciplinary perspective on the question of who cares for and about the elderly The contributors reflect on research studies experimental programmes and personal experience in Japan and the United States to explicitly compare how policies practices and interpretations of elder care are evolving at the turn of the century     *Textbook of Palliative Medicine* Eduardo Bruera, Irene Higginson, Charles F von Gunten, 2009-01-30 Textbook of Palliative Medicine provides an alternative truly international approach to this rapidly growing specialty This textbook fills a niche with its evidence based multi professional approach and global perspective ensured by the international team of editors and contributing authors In the absence of an international curriculum for the study of palliative medicine this textbook provides essential guidance for those both embarking upon a career in palliative medicine or already established in the field and the structure and content have been constructed very much with this in mind With an emphasis on providing a service anywhere in the world including the important issue of palliative care in the developing nations Textbook of Palliative Medicine offers a genuine alternative to the narrative approach of its competitors and is an ideal complement to them It is essential reading for all palliative care physicians in training and in practice as well as palliative care nurses and other health professionals in the palliative care team

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