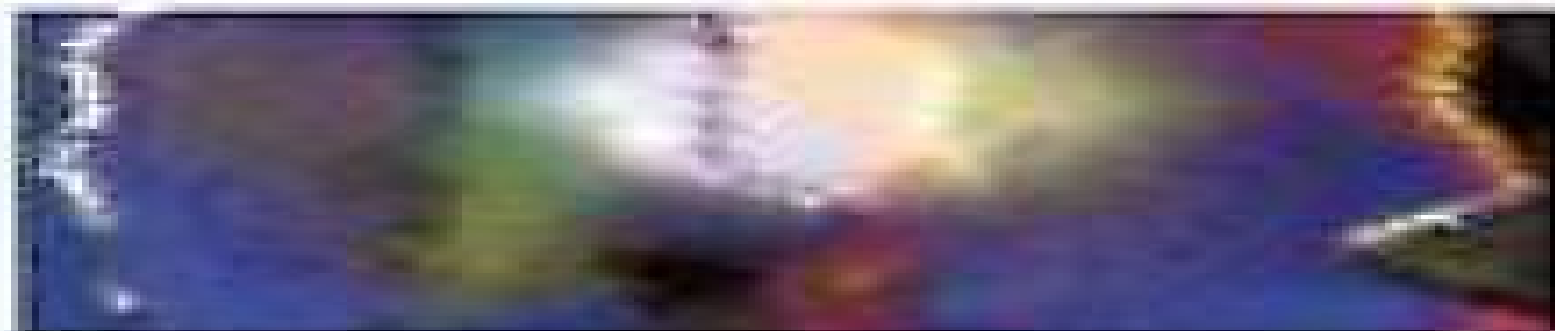




REFLECTIONS ON PALLIATIVE CARE

David Clark
Jane Seymour



Reflections On Palliative Care Sociological And Policy Perspectives

**Lieve Van Den Block,Gwenda
Albers,Sandra Martins Pereira**



Reflections On Palliative Care Sociological And Policy Perspectives:

Reflections on Palliative Care David Clark, Jane Seymour, 1999 By drawing on a wide range of sources Clark and Seymour offer a set of reflections on the development of palliative care and its place within a wider social context This work considers ethical questions and palliative care's history too

Dying: A Social Perspective on the End of Life Alex Broom, 2016-03-03 An inevitable and universal experience dying is experienced by individuals in different ways often related to the character of our relationships family structures gender identities cultural backgrounds and economic means Drawing on extensive qualitative fieldwork with patients carers and health professionals in Australia and the United Kingdom *Dying A Social Perspective on the End of Life* provides a critical examination of the different spheres of dying in social and cultural context Exploring complex issues such as the politics of assisted dying negotiating medical futility gender and dying the desire for redemption the moralities of the good fight and the lived experience of bodily disintegration this book links novel theoretical ideas within sociology to cutting edge empirical data collected in palliative and end of life care contexts A theoretically engaged understanding of the social mediation of the end of life *Dying A Social Perspective on the End of Life* also sheds light on the manner in which the end of life can be shaped by major economic cultural and socio cultural shifts including neo liberalism individualisation medicalisation professionalisation and detraditionalisation As such it will appeal to social science health and medical researchers interested in the end of life as well as those working in palliative and end of life care settings

Governing Death and Loss Stephen Conway, 2011 Political economic social cultural and technological changes have led to profound transformations in the ways that death and loss are perceived and managed in contemporary society Over the last few decades the long term shift to chronic illness as a major causal factor has significantly increased the time scale of dying Most people die in institutions and care is typically medical Many communities and ordinary citizens now relinquish control and involvement to experts in the last stages of life However at global and local levels new institutional arrangements are emerging to govern the changing face of death and a new model is being developed to counter claims of the creeping medicalisation of death and dying Emphasising the international context of the issues involved this book illustrates the interlinking nature of society death and loss and gives examples of governance that promotes the empowerment participation and the increasing need for the involvement of ordinary people and communities in differing social and cultural contexts All chapters are written at an accessible level and will appeal to a wide readership Part 1 of the book provides a sociological understanding of the governance of death and loss in international and historical contexts and the implications for practice Part 2 provides examples of good practice drawing upon a sociological understanding

To Comfort Always David Clark, 2016-10-06 Palliative medicine was first recognised as a specialist field in 1987 One hundred years earlier London based doctor William Munk published a treatise on easeful death that mapped out the principles of practical spiritual and medical support at the end of life In the intervening years a major process of development took place

which led to innovative services new approaches to the study and relief of pain and other symptoms a growing interest in holistic care and a desire to gain more recognition for care at the end of life This book traces the history of palliative medicine from its nineteenth century origins to its modern practice around the world It takes in the changing meaning of euthanasia assesses the role of religious and philanthropic organisations in the creation of homes for the dying and explores how twentieth century doctors created a special focus on end of life care To Comfort Always traces the rise of clinical studies academic programmes and international collaborations to promote palliative care It examines the continuing need to support development with evidence and assesses the dilemmas of unequal access to services and pain relieving drugs as well as the periodic accusations of creeping medicalization within the field This is the first history of its kind and the breadth of information it encompasses makes it an essential resource for those interested in the long term achievements of palliative medicine as well as the challenges that remain

Pain and Palliative Care in the Developing World and Marginalized Populations M.r. Rajagapol, 2004-03-31 Essential information for anyone involved in palliative care programs for deprived patients In this comprehensive resource leading healthcare professionals describe pioneering work on the front lines of pain and palliative care service planning and implementation for underserved populations Pain and Palliative Care in the Developing World and Marginalized Populations A Global Challenge explores the challenges and barriers preventing satisfactory pain management for patients who urgently need it This book provides you with true accounts of palliative care programs from around the world to help you meet the needs of disadvantaged clients This essential volume includes a Foreword written by a world leader in palliative care Jan Stjernsward Former Chief of the Cancer and Palliative Care Program of the World Health Organization and currently International Director of the Oxford International Centre for Palliative Care in the United Kingdom Pain and Palliative Care in the Developing World and Marginalized Populations A Global Challenge addresses issues of vital importance for the global health care community such as Why do so many people in the developing world suffer excruciating pain for months and years when simple inexpensive medication could make them comfortable They get MRI scans why don t they have access to palliative care Why do some palliative care programs fail to reach the needy How could a palliative care delivery system be adapted to local needs Why are medical and nursing students not taught the fundamentals of pain management What direction should palliative care education take Could health care resources be channeled to deliver care in a more just and equitable manner This book chronicles the efforts of ambitious pain management care professionals to confront these questions working toward an end to needless preventable pain and suffering It examines their programs and acknowledges their successes and failures to date with commentaries by international experts This indispensable manual discusses palliative care programs in developing countries such as India Chile Argentina Saudi Arabia Thailand Hong Kong Malaysia and others Pain and Palliative Care in the Developing World and Marginalized Populations also offers an important look at pain management programs geared toward several specific

underserved populations in both developing and developed countries including Native Americans and inmates in a New Zealand prison Illustrated with figures graphs and tables this book is essential for practitioners and officials in both palliative and public health care All proceeds from sales of this book will be used to support the growth of palliative care programs in India Palliative Care for Older People Lieve Van Den Block,Gwenda Albers,Sandra Martins Pereira,2015 Offering not only insights into the current state of policy work around the world it also offers examples of good practice and recommendations for the future Recommendations that can inspire support and direct healthcare policy and decision making at organisational regional national and international levels Living, Dying, Death, and Bereavement (Volume Two) David E. Balk,2020-10-21 This two volume book offers extensive interviews with persons who have made significant contributions to thanatology the study of dying death loss and grief The book s in depth conversations provide compelling life stories of interest to clinicians researchers and educated lay persons and to specialists interested in oral history as a means of gaining rich understandings of persons lives Several disciplines that contribute to thanatology are represented in this book such as psychology religious studies art literature history social work nursing theology education psychiatry sociology philosophy and anthropology The book is unique no other text offers such a comprehensive insightful and personal review of work in the thanatology field The salience of thanatology is obvious when we consider several topics including the aging demographics of most countries the leading causes of death the devastation of COVID 19 the realities of how most persons die the growth both of hospice and of efforts within medicine to ensure that a good death becomes the norm of medical practice and increases in the number of countries and states permitting physician assisted suicide This second volume includes conversations with 16 thanatologists a rich extensive bibliography an index of names and subjects and a biographical sketch of the author The experts interviewed in this volume include Danai Papadatou Holly Prigerson Jack Jordan Illene Cupit Heather Servaty Seib Irwin Sandler Simon Shimshon Rubin Carla Sofka Harold Ivan Smith and Phyllis Kosminsky **Developing Holistic Care for Long-term Conditions** Carl Margerison,Steve Trenoweth,2009-10-16 Comprising four sections this text introduces the policy and background to caring for people with chronic illness as well as the psychosocial impact of long term conditions Essential skills for practice are explored including holistic assessment symptom control and the promotion of effective partnership between client and carer in supporting coping recovery and end of life care There is an emphasis on maximising individual health potential and resilience with the role of nutrition exercise complementary therapy and spirituality considered The focus is on client centred care which addresses the whole person mind and body The extensive final section presents examples of key health issues where UK national guidelines have been published including Long term neurological conditions Diabetes Mental health Cancer Coronary heart disease Older people Palliative Care and Communication Anne-Mei The,2002 Palliative Care and Communication seeks explanations for the sense of optimism found among patients with small cell bronchial carcinoma or lung cancer Over a period of five years the author used ethnographic research methods to

monitor the illness processes of patients from the time of receiving bad news until their death from the perspectives of the various people involved the patient family and relatives the doctors and nursing staff Based on narrative descriptions interspersed with observations the author demonstrates why things happen the way they do in practice The book explores informal codes ambiguous messages the dilemma between professional detachment and personal involvement patterns of information and communication during the phases of the illness the latent realisation of approaching death and the ambivalence of patients with regard to knowing and not knowing Anne Mei The s engaging writing style and extensive use of narrative make this book accessible to a wide range of readers doctors nurses policy makers patients relatives researchers in the field of health care and social scientists It is suitable for use on various branches of medical and nursing training to illustrate the process of communication with patients during the terminal phase of a patient s illness *Abjectly Boundless* Trudy Rudge, 2016-03-16 Within a variety of practice environments health professionals often experience feelings of disgust and repulsion towards the presence of an abject object Cadaverous sick disabled bodies troubled minds wounds vomit and so forth are all part of health and care work and threaten the clean and proper bodies of those who undertake it yet this unclean side of health work is rarely accounted for in academic literature This volume employs the work of Julia Kristeva through a range of case studies drawn from care and nursing settings around the world It brings together work from researchers and practitioners within the social and health sciences the caring professions and psychotherapy to expose and highlight the important impact of the concept of abjection which historically has been silenced in the health sciences

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