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ONCOLOGIST**

**FAMILY
DOCTOR**

**EXERCISE
THERAPIST**

**PHYSIOTHERAPIST/
MASSAGE THERAPIST**

The Cancer Patient Social And Medical Aspects Of Care

National Cancer Institute (U.S.)



The Cancer Patient Social And Medical Aspects Of Care:

The Cancer Patient Barrie R. Cassileth, 1979

The Cancer Patient and Supportive Care Donald J. Higby, 2012-12-06

During the course of editing *Supportive Care in Cancer Therapy* Martinus Nijhoff Publishers 1983 it became apparent that several topics would have to await a second volume. Furthermore development of new information and evolution of ideas continues. This volume continues the intent of the first to present reviews of issues relating to supportive care and to identify areas where further definition and further research is needed. The physician reading this volume will find the contents thought provoking. In addition to reviews authored by physicians there are chapters authored by non physicians who present a different perspective and a different style of writing. Reverend Bigler writes from his long experience as a chaplain for cancer patients and tries to identify the changes that take place in the personality of the chaplain who works with dying patients. Ms. Kilgallon and Ms. Powell try to describe what an Oncology Nurse is rather than what one does. Attorneys Reese and Price present a very practical summation of issues which face the cancer patient and their chapter could easily be copied and offered to one's patients. Furthermore this chapter gives direction to the physician who is frequently called upon for advice regarding issues which lie entirely outside his/her formal training. Excellent reviews on more medical subjects are here. Cancer in the Elderly should provoke oncologists to re-examine their approach to the geriatric patient. [NCI](#)

Publications for Cancer Patients and the Public National Cancer Institute (U.S.), 1994

Supportive Care in Cancer

Patients Hans-Jörg Senn, Agnes Glaus, Luzius Schmid, 2012-12-06. The symposium on supportive care in cancer patients which took place in St. Gallen, Switzerland on February 18-21, 1987, welcomed renowned experts in the field and more than 600 participants from 25 countries with the aim of stimulating discussion on how to improve our professional skills and personal attitudes toward cancer patients in all stages of their disease. Why did we organize such a symposium on supportive care in cancer patients? Recent decades have witnessed remarkable success in cancer treatment and we have learned how to cure a finite number of neoplastic diseases. Some malignant tumors that previously entailed high fatality rates such as leukemias, lymphomas, and testicular cancers can now be cured even when at an advanced stage. Yet it seems to many that our struggle to improve results and to fight death from cancer has also imposed greater toxicity on patients. Conventional scientifically based oncology has only recently made adequate efforts to improve the subjective quality of life of cancer patients, for example through prophylaxis against emesis, nausea, and scalp hypothermia, pain control, and the development of psychosocial support structures. The search for less toxic and yet equally effective treatment measures has not been one of our primary goals in the past. Supportive care has always been part of nurses' professional aim even though many have not known how best to offer it. **Cancer Caregiving in the United States** Ronda C. Talley, Ruth

McCorkle, Walter F. Baile, 2012-04-24. Despite advances in detection and treatment, cancer remains a source of pain and distress to patients and of complex challenges to the loved ones caring for them. The trend toward shorter hospital stays in

particular has increased the physical psychological and financial burden on caregivers often leading to adverse effects on patients Cancer Caregiving in the United States illuminates these complex concerns with authoritative detail This wide ranging volume provides a comprehensive survey of cancer related issues including those affecting the care triad patients family members professionals and quality of care as well as the numerous physical emotional and financial challenges that caregivers may need to confront Sources of caregiver difficulty at each stage of the disease from diagnosis to end of life are explored Each chapter analyzes its topic in terms of practice research education and policy providing a wealth of literature reviews assessment and care models interventions and recommendations for future study and practice Coverage includes Caregiving issues for cancer patients with long term short term and intermittent needs Family caregivers as members of the treatment team The impact of health disparities on caregivers Cancer care policy and advocacy End of life issues for cancer caregivers Legal financial and ethical issues Cancer Caregiving in the United States is a core reference for researchers professionals scientist practitioners and graduate students in such caregiving fields as clinical psychology social work nursing public health and medicine social policy and educational policy

The Complete Guide to Complementary Therapies in Cancer Care Barrie R. Cassileth, 2011 This book guides the way through the often confusing maze of complementary and alternative therapies promoted to cancer patients and survivors The functions benefits backgrounds and risks are clearly presented from publisher description

Complete Guide To Complementary Therapies In Cancer Care, The: Essential Information For Patients, Survivors And Health Professionals Barrie R Cassileth, 2011-07-18 With recent clinical advances millions of people survive many years after a cancer diagnosis And while their physicians deliver conventional evidence based therapies to treat the cancer sometimes the cancer patient and his or her symptoms are not treated with equal skill To manage their physical and emotional symptoms and promote health and well being most cancer patients and survivors use complementary therapies Naturopathy Ayurveda herbalism homeopathy hypnosis yoga acupuncture music therapy macrobiotics chelation therapy colonics hydrotherapy and many many more But are they safe Are they effective What problems do they address What are the risks When can they help This unique and comprehensive book guides the way through the often confusing maze of complementary and alternative therapies promoted to cancer patients and survivors The functions benefits backgrounds and risks are clearly presented Learning when if and how to use them provides medical professionals cancer patients and survivors with the information they need to better control the symptoms and side effects of cancer and its treatment Unfortunately using some of these therapies without this expert guidance can lead to medical complications or worse The Complete Guide to Complementary Therapies in Cancer Care is an invaluable resource in making educated health care decisions for managing life during and after cancer Through the wise and informed use of these approaches cancer patients whether just diagnosed during treatment or throughout Survivors are better able to manage the physical and emotional stresses that accompany cancer leading to symptom control and improved quality of

life *Current Catalog* National Library of Medicine (U.S.),1979 First multi year cumulation covers six years 1965 70

Patient and professional education materials for ostomates Kathy Jursik,1981 *Source Book of Educational Materials for Nuclear Medicine* ,1981 **Cancer and Self-help** Mark A. Chesler,Barbara K. Chesney,1995 Written for and about the special population of parents of children with cancer this book explores the remarkable effectiveness of self help groups and profiles their rapid rise as a resource complementing traditional health care Mark A Chesler and Barbara K Chesney draw on their own experience as members of such groups and on a combined thirty years of research on self help They provide essential information for families of children with cancer and other chronic life threatening illnesses for health care professionals working with them and for scholars of self help and psychosocial processes in health care including explanations of how self help groups function why they are effective and how they can be created and maintained The authors show that through self help groups parents can learn coping skills find personal affirmation and mutual support and share the wisdom gained from their experiences Chesler and Chesney find that group participation improves parents coping capabilities in the face of terrible odds and fosters an increased sense of empowerment as they care and advocate for their children in an increasingly complex health care system *Cancer and Self Help* distills the experiences of more than fifty self help groups and their members over twelve years It also places cancer self help groups in a larger context comparing them to other social movement organizations and to other strategies for personal coping or change The book includes the voices of individual parents and professionals recounting their experiences detailed examples of group activities programs operating procedures and organizational structures fundamental how to information on forming a self help group comments on the roles and dilemmas of health care professionals in these groups and on the medical care system as a whole and interpretations of these individual and organizational dynamics **Handbook of Oncology Social Work** Grace Hyslop Christ,Carolyn Messner,Lynn C. Behar,2015 This inaugural Handbook of Oncology Social Work Psychosocial Care for People with Cancer provides for the first time a repository of the breadth and scope art and science of oncology social workers practice education research policy and program leadership in the psychosocial care of people with cancer and their families *Topics in Palliative Care* Russell K. Portenoy,Eduardo Bruera,1997 **Reconstitution Of Social Work: Towards A Moral Conception Of Social Work Practice** Yuk-ying Ho,Sun-pong Yuen,2010-04-06 This volume is the first book length explication of hermeneutics in social work It attempts to clarify and reconstruct the moral basis of social work Against the mainstream current of doing social work with the technical rational outlook this book argues that value constitutes the very core of social work It is with this solid foundation of moral concepts that social work techniques are reconstituted This volume seamlessly combines theoretical discussions with empirical studies It starts with a theoretical inquiry that combines hermeneutics and critical theory and examines the moral nature of social work It then extends the theoretical analysis to empirical research on mental illness cancer community development and social work management Redefining the

relationship between theory and practice the discussion on first person value involvement and dialogical mode of understanding will inspire social workers to develop their professional practice in a new light This volume will capture the attention of both social work scholars and frontline social work practitioners The hermeneutic point of view will also be of interest to readers students of social theory and social research **An Anthropology of Lying** Sylvie Fainzang,2016-03-09

In the era of health democracy where a patient s right to be informed is not only widely advocated but also guaranteed by law what is the real situation regarding patient information Do patients receive the information that they request with regard to their diagnosis prognosis or treatments And what information do patients themselves give to their doctors Drawing on observational research in hospitals and covering the exchanges between doctors and patients on the subject of cancer treatment and that of other pathologies this book reveals that the practice of telling lies is widespread amongst parties on both sides of the medical relationship With attention to the manner in which information of various types is withheld and the truth concealed on either side of the doctor patient relationship the author explores the boundaries between what is said and what is left unsaid and between those who are given information and those who are lied to Considering the misunderstandings that occur in the course of medical exchanges and the differences between the lies told by doctors and patients **An Anthropology of Lying Information in the Doctor Patient Relationship** analyses the role of mendacity in the exercise of and resistance to power A fascinating study of the mechanisms at work and social conditions surrounding the accomplishment of lying in medical settings this book casts fresh light on a subject that has so far been overlooked As such it will appeal not only to sociologists and anthropologists of health and medicine but also to medical professionals **Library of Congress Subject Headings** Library of Congress. Cataloging Policy and Support Office,2009 Library of Congress Subject Headings Library of Congress,2003 *Social Aspects of Care* Nessa Coyle,2016 *Social Aspects of Care* provides an overview of financial and mental stress illness places not just on the patient but on the family as well This volume contains information on how to support families in palliative care cultural considerations important in end of life care sexuality and the impact of illness planning for the actual death and bereavement **Library of Congress Subject Headings: P-Z** Library of Congress. Subject Cataloging Division,1989 Library of Congress Subject Headings Library of Congress. Office for Subject Cataloging Policy,1992

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