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Ethical Issues in Cancer Patient Care

Springer Science & Business Media This book addresses a variety of ethical issues that arise in the care of oncology patients. Many volumes have been written on medical ethics in the past 30 years. However, few have focused on ethical issues specific to the care of cancer patients. This book brings together such a focused examination. The contributors are experienced clinicians, ethicists, medical humanists, and medical educators. The issues raised have direct relevance to the care of oncology patients in treatment as well as research settings. The chapters address issues that are central to contemporary medical practice and medical ethics inquiry. Any practicing clinician will be well aware of the problems of communication and how uncertainty, cross-cultural issues, and religious influences can impact patient care. The limits of care and the role of advance directives and palliative care are common issues that must be addressed in treating patients at the end of life. For oncologists and oncology patients, participation in clinical trials may be a thorny topic, especially when phase I clinical trials are being considered. The impact of managed care and reimbursement issues cannot be avoided in the contemporary patient care and similarly cannot be neglected when considering the ethical ramifications raised. No discussion of ethics in oncology can be complete without attention to the specific challenges raised by the pediatric patient with cancer. All of these topics are explored by the contributors to this book.

Ethical Challenges in Oncology

Patient Care, Research, Education, and Economics

Academic Press Ethical Challenges in Oncology: Patient Care, Research, Education, and Economics covers a wide variety of topics and viewpoints about ethical issues that arise in oncology throughout the full cancer care continuum. This book provides a holistic view on oncology ethics, incorporating the knowledge and expertise of authors from various departments and oncology specialties within the University of Texas MD Anderson Cancer Center. The first section focuses on the ethical issues associated with treating cancer patients. Next, the ethical challenges associated with oncology research, including funding, regulation, subject selection and the big picture are explored. The third section covers ethical issues associated with education in oncology as it reflects both past and future trends in developing proficient health care providers, patients and even executives. Final sections concentrate on the ethical dilemmas associated with the economic repercussions of oncology, offering thoughts on how to alleviate the ethical consequences that can arise from the global effects of cancer and cancer treatment. Each chapter includes discussion topics, answers pertinent questions and provides an ethical framework for problem-solving in each scenario. The topics uncover the ethical apprehensions and problems associated with oncology research and practice in order to determine best practices as well as provide guidance for all parties involved. Offers a diverse range of topics and viewpoints about ethical issues that arise in oncology Provides direction by bringing real issues to the forefront Contains a truly ethical approach on problems as broad as research, finance, appropriateness of care, and professional education

Assessing and Improving Value in Cancer Care

Workshop Summary

National Academies Press Unlike many other areas in health care, the practice of oncology presents unique challenges that make assessing and improving value especially complex. First, patients and professionals feel a well-justified sense of urgency to treat for cure, and if cure is not possible, to extend life and reduce the burden of disease. Second, treatments are often both life sparing and highly toxic. Third, distinctive payment structures for cancer medicines are intertwined with practice. Fourth, providers often face tremendous pressure to apply the newest technologies to patients who fail to respond to established treatments, even when the evidence supporting those technologies is incomplete or uncertain, and providers may be reluctant to stop toxic treatments and move to palliation, even at the end of life. Finally, the newest and most novel treatments in oncology are among the most costly in medicine. This volume summarizes the results of a workshop that addressed these issues from multiple perspectives, including those of patients and patient advocates, providers, insurers, health care researchers, federal agencies, and industry. Its broad goal was to describe value in oncology in a complete and nuanced way, to better inform decisions regarding developing, evaluating, prescribing, and paying for cancer therapeutics.

Ethics and Cancer

An Annotated Bibliography

776 references that deal with the ethical issues involved in cancer care and research. Monographic and serial literature cited is multidisciplinary in nature, covering such areas as medicine, philosophy, ethics, law, psychology, psychiatry, social work, and nursing. Alphabetical arrangement by authors. Contains an outline of ethical problem areas. Author, subject, and ethical positions indexes.

Structural Racism and Community Building

Ethical Issues in Pediatric Hematology/Oncology

Springer Nature This book identifies the various ethical challenges that arise in pediatric hematology/oncology and provides the necessary tools to overcome these challenges. Aiming to expand upon and strengthen providers' knowledge and experience in pediatric health care ethical issues, the text positions providers to be beneficial resources to faculty, staff, patients, and families within their institution. It presents a multidisciplinary approach to sound ethical practices that is necessary to effectively care for these patients and their families. The book reviews the principles of ethical decision-making, the unique difficulties in using children as research subjects, common ethical conundrums involved in providing end-of-life care, and general moralities of professional practice. Written by experts in their fields, Ethical Issues in Pediatric Hematology/Oncology is an innovative and valuable resource for clinicians, practitioners, and trainees who work in the field of pediatric hematology/oncology.

Issues in Palliative Care Research

Oxford University Press Symptom control, management of psychosocial and spiritual concerns, decision-making consistent with values and goals, and care of the imminently dying that is appropriate and sensitive are among the critical issues in palliative care. This book explores progress made and future goals.

Cancer Caregiving in the United States

Research, Practice, Policy

Springer Science & Business Media 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.

Pediatric Psycho-Oncology

A Quick Reference on the Psychosocial Dimensions of Cancer Symptom Management

Oxford University Press, USA Preceded by: Quick reference for pediatric oncology clinicians / senior editors, Lori S. Wiener, Maryland Pao. c2009.

The Ethics of Personalised Medicine

Critical Perspectives

Routledge In recent times, the phrase 'personalised medicine' has become the symbol of medical progress and a label for better health care in the future. However, a controversial debate has developed around whether these promises of better, more personal and more cost-efficient medicine are realistic. This book brings together leading researchers from across Europe and North America, from both normative and empirical disciplines, who take a more critical view of the often encountered hype associated with personalised medicine. Partially drawing on a four year collaborative research project funded by the German Ministry for Education and Research, the book presents a multidisciplinary debate on the current state of research on the ethical, legal and social implications of personalised medicine. At a time when future health care is a topic of much discussion, this book provides valuable policy recommendations for the way forward. This study will be of interest to researchers from various disciplines including philosophy, bioethics, law and social sciences.

Cancer Care for the Whole Patient

Meeting Psychosocial Health Needs

National Academies Press Cancer care today often provides state-of-the-science biomedical treatment, but fails to address the psychological and social (psychosocial) problems associated with the illness. This failure can compromise the effectiveness of health care and thereby adversely affect the health of cancer patients. Psychological and social problems created or exacerbated by cancer--including depression and other emotional problems; lack of information or skills needed to manage the illness; lack of transportation or other resources; and disruptions in work, school, and family life--cause additional suffering, weaken adherence to prescribed treatments, and threaten patients' return to health. Today, it is not possible to deliver high-quality cancer care without using existing approaches, tools, and resources to address patients' psychosocial health needs. All patients with cancer and their families should expect and receive cancer care that ensures the provision of appropriate psychosocial health services. Cancer Care for the Whole Patient recommends actions that oncology providers, health policy makers, educators, health insurers, health planners, researchers and research sponsors, and consumer advocates should undertake to ensure that this standard is met.

Ethical Dilemmas at the End of Life

Explores a range of issues--including pediatric hospice, historical, religious, spiritual and cultural perspectives on the end of life, hospice in nursing homes, surrogate decision making, physician assisted suicide, organ donation, and our society's legal tenants of end-of-life care. Includes an index.

Quality of Life Among Cancer Survivors

Challenges and Strategies for Oncology Professionals and Researchers

Springer This multidisciplinary reference explores the concepts and realities of quality of life among cancer survivors in its physical, psychological, cognitive, social, and familial dimensions. Informed by a broad range of fields including genetics, psychiatry, nursing, dentistry, rehabilitation, and ethics, it addresses daily challenges of living for this population, from self-care to cultural concerns and from social interactions to experiences with providers. Family issues of pediatric, young adult, and elder survivors, caregiving parents, and siblings are a major area of concern. And contributors describe interventions for survivors as individuals, in family content, and as part of integrated care across primary and specialty settings. Included among the topics: Play, leisure activities, and cognitive health among older cancer survivors. Genetic mutations in cancer susceptibility genes: a family history of cancer. Cancer patients in a pediatric intensive care unit: a single center experience. The impact of childhood cancer on the quality of life among healthy siblings. When cancer returns: family caregivers and the hospice team. Experiencing cancer services: a story of survival and dissatisfaction. A significant addition to the cancer survivorship literature, Quality of Life Among Cancer Survivors is a practice-building resource for oncology and allied health professionals, health psychologists, and social workers, as well as researchers in these fields.

Ethical Challenges in Cancer Diagnosis and Therapy

Springer This book presents in detail the problems and ethical challenges in daily oncological practice. In western industrialized countries, roughly 25 percent of all citizens still die from cancer. Despite significant progress in basic science and in individual areas of clinical care, even in the 21st century, being diagnosed with cancer has lost none of its dread and can still be a death sentence. This situation raises many problems and challenges for medical ethics, e.g., the question of the benefits and risks of prevention programs, or the right to know and not to know. Clinical trials with cancer patients and quality assurance for surgery, radiotherapy and medication also pose a series of ethical dilemmas. Furthermore, cancer treatment is a psychological challenge not only for patients but also for physicians and caregivers. The issues of adequate pain management and good palliative care, of treatment limiting and the question of assisted suicide at the end of life also have to be considered. In order to reflect the subject's diverse and multifaceted nature, the book incorporates legal, ethnographic, historical and literary perspectives into ethical considerations.

Principles and Practice of Fertility Preservation

Cambridge University Press The specialty of fertility preservation offers patients with cancer, who are rendered infertile by chemo- and radiotherapy, the opportunity to realize their reproductive potential. This gold-standard publication defines the specialty. The full range of techniques and scientific concepts is covered in detail, and the author team includes many of the world's leading experts in the field. The book opens with introductions to fertility preservation in both cancer and non-cancer patients, followed by cancer biology, epidemiology and treatment, and reproductive biology and cryobiology. Subsequent sections cover fertility preservation strategies in males and females, including medical/surgical procedures, ART, cryopreservation and transplantation of both ovarian tissue and the whole ovary, and in-vitro follicle growth and maturation. Concluding chapters address future technologies, as well as ethical, legal and religious issues. Richly illustrated throughout, this is a key resource for all clinicians specializing in reproductive medicine, gynecology, oncology, hematology, endocrinology and infertility.

Palliative Care

Psychosocial and Ethical Considerations

A great number of cancer patients will suffer some form of social, emotional or psychological distress and challenges as a result of the disease and its treatment. Unattended, psychosocial issues can leave patients and families ill-equipped to cope and manage their cancer diagnosis and treatment. When psychosocial care is properly integrated into clinical care, it has a direct impact on the patients quality of life. Psychosocial aspects must be integrated into routine cancer care. The patient should be screened at their initial visit for psychosocial needs, and survivors should have a treatment plan that includes attention to possible increased anxiety on completing treatment, development of posttraumatic stress symptoms, mixed anxiety and depressive symptoms. In this book, the authors cover clinical, psychosocial and end-of-life aspects.

The Immortal Life of Henrietta Lacks

Pan Macmillan Her name was Henrietta Lacks, but scientists know her as HeLa. She was a poor black tobacco farmer who worked the same land as her slave ancestors yet her cells - taken without her knowledge - become one of the most important tools in modern medicine.

Mastering Communication with Seriously Ill Patients

Balancing Honesty with Empathy and Hope

Cambridge University Press Physicians who care for patients with life-threatening illnesses face daunting communication challenges. Patients and family members can react to difficult news with sadness, distress, anger, or denial. This book defines the specific communication tasks involved in talking with patients with life-threatening illnesses and their families. Topics include delivering bad news, transition to palliative care, discussing goals of advance-care planning and do-not-resuscitate orders, existential and spiritual issues, family conferences, medical futility, and other conflicts at the end of life. Drs Anthony Back, Robert Arnold, and James Tulsky bring together empirical research as well as their own experience to provide a roadmap through difficult conversations about life-threatening issues. The

book offers both a theoretical framework and practical conversational tools that the practising physician and clinician can use to improve communication skills, increase satisfaction, and protect themselves from burnout.

Delivering High-Quality Cancer Care

Charting a New Course for a System in Crisis

National Academies Press In the United States, approximately 14 million people have had cancer and more than 1.6 million new cases are diagnosed each year. However, more than a decade after the Institute of Medicine (IOM) first studied the quality of cancer care, the barriers to achieving excellent care for all cancer patients remain daunting. Care often is not patient-centered, many patients do not receive palliative care to manage their symptoms and side effects from treatment, and decisions about care often are not based on the latest scientific evidence. The cost of cancer care also is rising faster than many sectors of medicine--having increased to \$125 billion in 2010 from \$72 billion in 2004--and is projected to reach \$173 billion by 2020. Rising costs are making cancer care less affordable for patients and their families and are creating disparities in patients' access to high-quality cancer care. There also are growing shortages of health professionals skilled in providing cancer care, and the number of adults age 65 and older--the group most susceptible to cancer--is expected to double by 2030, contributing to a 45 percent increase in the number of people developing cancer. The current care delivery system is poorly prepared to address the care needs of this population, which are complex due to altered physiology, functional and cognitive impairment, multiple coexisting diseases, increased side effects from treatment, and greater need for social support. *Delivering High-Quality Cancer Care: Charting a New Course for a System in Crisis* presents a conceptual framework for improving the quality of cancer care. This study proposes improvements to six interconnected components of care: (1) engaged patients; (2) an adequately staffed, trained, and coordinated workforce; (3) evidence-based care; (4) learning health care information technology (IT); (5) translation of evidence into clinical practice, quality measurement and performance improvement; and (6) accessible and affordable care. This report recommends changes across the board in these areas to improve the quality of care. *Delivering High-Quality Cancer Care: Charting a New Course for a System in Crisis* provides information for cancer care teams, patients and their families, researchers, quality metrics developers, and payers, as well as HHS, other federal agencies, and industry to reevaluate their current roles and responsibilities in cancer care and work together to develop a higher quality care delivery system. By working toward this shared goal, the cancer care community can improve the quality of life and outcomes for people facing a cancer diagnosis.

Ethics in Oncology Nursing

Nursing Practice

The Ethical Issues

Prentice Hall

Radiation Oncology

Management Decisions

Lippincott Williams & Wilkins Ideal for on-the-spot consultation, this pocket manual, *Radiation Oncology: Management Decisions*, provides easily accessible information for residents and practitioners in radiation oncology. It presents the most essential information that is immediately required in the clinical setting. The first eight chapters of the book focus on key basic concepts; the remaining 46 chapters describe treatment regimens for all cancer sites and tumor types. Includes coverage of pain and palliation, and covers all latest therapeutic techniques. This edition includes expanded information on image-guided therapy, 3D techniques, and 4D protocols. The updated cancer staging guidelines have been used throughout the manual. In addition, there is a brand-new chapter devoted to QUANTEC dosage recommendations.

Legal and Ethical Issues in Cancer Care in the United States

Charles C Thomas Pub Limited

To Err Is Human

Building a Safer Health System

National Academies Press Experts estimate that as many as 98,000 people die in any given year from medical errors that occur in hospitals. That's more than die from motor vehicle accidents, breast cancer, or AIDS--three causes that receive far more public attention. Indeed, more people die annually from medication errors than from workplace injuries. Add the financial cost to the human tragedy, and medical error easily rises to the top ranks of urgent, widespread public problems. *To Err Is Human* breaks the silence that has surrounded medical errors and their consequence--but not by pointing fingers at caring health care professionals who make honest mistakes. After all, to err is human. Instead, this book sets forth a national agenda--with state and local implications--for reducing medical errors and improving patient safety through the design of a safer health system. This volume reveals the often startling statistics of medical error and the disparity between the incidence of error and public perception of it, given many patients' expectations that the medical profession always performs perfectly. A careful examination is made of how the surrounding forces of legislation, regulation, and market activity influence the quality of care provided by health care organizations and then looks at their handling of medical mistakes. Using a detailed case study, the book reviews the current understanding of why these mistakes happen. A key theme is that legitimate liability concerns discourage reporting of errors--which begs the question, "How can we learn from our mistakes?" Balancing regulatory versus market-based initiatives and public versus private efforts, the Institute of Medicine presents wide-ranging recommendations for improving patient safety, in the areas of leadership, improved data collection and analysis, and development of effective systems at the level of direct patient care. *To Err Is Human* asserts that the problem is not bad people in health care--it is that good people are working in bad systems that need to be made safer. Comprehensive and straightforward, this book offers a clear prescription for raising the level of patient safety in American health care. It also explains how patients themselves can influence the quality of care that they receive once they check into the hospital. This book will be vitally important to federal, state, and local health policy makers and regulators, health professional licensing officials, hospital administrators, medical educators and students, health caregivers, health journalists, patient advocates--as well as patients themselves. First in a series of publications from the Quality of Health Care in America, a project initiated by the Institute of Medicine

Patient-Centred Ethics and Communication at the End of Life

CRC Press This book provides the best information available on the ways priorities are currently set for health care around the world. It describes the methods now used in the six countries leading the process, and contrasts the differences between them. It shows how, except in the UK, frameworks have now been developed to set priorities. *Making Choices for Health Care* sets forth the key issues that need to be tackled in the years ahead. Descriptions of the leading trends are accompanied by suggestions to resolve outstanding difficulties. Topics include: the need for national research and development funding for new treatments, ways to shift resources permanently towards prevention and chronic care, and how DALYs may replace QALYs. While the concepts and values underlying priority setting have been discussed elsewhere, *Making Choices for Health Care* highlights real current practice. It is a vital tool for policy-makers, health care managers, clinicians, patient organizations, academics, and executives in pharmaceutical and medical supply industries.

Palliative Care for Non-cancer Patients

The specialty of palliative care has traditionally grown out of oncology and there has been little research into the needs of patients dying from causes other than cancer. Few non-cancer patients receive hospice in-patient, home care or day care although a good proportion of hospices say that their services are available to non-cancer patients. As a result, the importance of palliative care for non-cancer patients is now being increasingly recognized internationally, and in the UK a committee reporting to the Department of Health recommended that palliative care should be accessible to all patients who need such care. *Palliative Care of the Non-cancer Patient* considers the needs and experiences of patients dying from, for instance, stroke, heart disease or dementia by drawing on a range of disciplines and specialties in medicine. The provision of palliative care for patients dying from causes other than cancer raises a number of important questions for policy makers and purchasers. This book summarises what is known about the needs of and appropriate service provision for people dying of causes other than cancer and begins to set a research agenda.

Clinical Pain Management : Cancer Pain

CRC Press Now divided into four parts, the second edition of *Cancer Pain* delivers broad coverage of the issues that arise in the management of malignancy-related pain, from basic science, through end of life care and associated ethical issues, to therapies, both medical and complementary. Part One reviews basis considerations in cancer pain management, includ

Geriatric Psycho-Oncology

A Quick Reference on the Psychosocial Dimensions of Cancer Symptom Management

Apos Clinical Reference Handbo Geriatric Psycho-Oncology is a comprehensive handbook that provides best practice models for the management of psychological, cognitive, and social outcomes of older adults living with cancer and their families. Chapters cover a wide range of topics including screening tools and interventions, psychiatric emergencies and disorders, physical symptom management, communication issues, and issues specific to common cancer sites. A resource section is appended to provide information on national services and programs. This book features contributions from experts designed to help clinicians review, anticipate and respond to emotional issues that often arise in the context of treating older cancer patients. Numerous cross-references and succinct tables and figures make this concise reference easy to use. *Geriatric Psycho-Oncology* is an ideal resource for helping oncologists and nurses recognize when it may be best to refer patients to their mental health colleagues and for those who are establishing or adding psychosocial components to existing clinics.

Treating Survivors of Childhood Abuse and Interpersonal Trauma, Second Edition

STAIR Narrative Therapy

Guilford Publications Now revised and expanded with 50% new content reflecting important clinical refinements, this manual presents a widely used evidence-based therapy approach for adult survivors of chronic trauma. *Skills Training in Affective and Interpersonal Regulation (STAIR) Narrative Therapy* helps clients to build crucial social and emotional resources for living in the present and to break the hold of traumatic memories. Highly clinician friendly, the book provides everything needed to implement STAIR--including 68 reproducible handouts and session plans--and explains the approach's theoretical and empirical bases. The large-size format facilitates photocopying; purchasers also get access to a Web page where they can download and print the reproducible materials. First edition title: *Treating Survivors of Childhood Abuse: Psychotherapy for the Interrupted Life*. New to This Edition *Reorganized, simplified sessions make implementation easier. *Additional session on emotion regulation, with a focus on body-based strategies. *Sessions on self-compassion and on intimacy and closeness in relationships. *Chapter on emerging applications, such as group and adolescent STAIR, and clinical contexts, such as primary care and telemental health. *Many new or revised handouts--now downloadable. *Updated for DSM-5 and ICD-11.

Handbook of Communication in Oncology and Palliative Care

OUP Oxford This comprehensive text provides clinicians with practical and evidence-based guidelines to achieve effective, patient-centered communication in the areas of cancer and palliative care. Written by an outstanding panel of international experts, it integrates empirical findings with clinical wisdom, draws on historical approaches and presents a state-of-the-art curriculum for applied communication skills training for the specialist oncologist, surgeon, nurse and other multi-disciplinary team members involved in cancer care today. In this book communication is broken down into key modules that cover the life-cycle of cancer care. They include coverage of diagnosis and treatment including clinical trials, empathic support in response to distress, transition to survivorship or palliative therapies, discussion of prognosis, conduct of family meetings, and care of the dying. Complementary training of patients in their communication with the doctor completes the interactive dyad. The art of teaching, impact of gender and power in the consultation and the ethical context are carefully considered. Special communication challenges include discussion of genetic risk, rehabilitative and salvage surgery, promotion of treatment adherence, unanticipated adverse outcomes, intercultural issues, fertility and sexuality. The value of decision aides, question prompt lists, audio-recording of consultations and use of the internet is illustrated. By looking across the full spectrum of disciplines involved in the multidisciplinary team, discipline-specific issues are considered by experts in each field. In this manner, the needs of patients and their relatives are evaluated, including paediatric and geriatric populations. To achieve all of this, theoretical models are examined from the medical school to the highly specialized practice, facilitation training and actor training are made explicit, and international approaches to communication skills training are compared and contrasted. Finally, research tools that assist in coding cancer consultations, evaluating training courses, and employing mixed methods in studies aid the reader in providing clear and sensitive communication when handling challenging situations whilst treating cancer sufferers and palliative care patients.

Nutrition Management of the Cancer Patient

Jones & Bartlett Learning Abstract: This book provides nutrition guidelines and management techniques for cancer patients. Written by dietitians, nutritionists, and physicians for health professionals who are providing nutrition support for cancer patients, this publication describes skills and techniques acquired by these experts through years of experience. Topics include: nutrition needs of cancer patients; cancer's impact on the nutrition status of patients; screening, assessing, and monitoring; nutrition concerns for specific patient populations; nutrition concerns of treatment modalities; methods of management; tube feeding; parenteral nutrition; home care training; ethical and psychologic issues relating to the cancer patient; and cancer quackery.

Cancer Care

Oxford University Press This book provides the practising GP with all the information needed to help in the management of patients with cancer.

Oxford Handbook of Respiratory Medicine

Oxford University Press, USA Respiratory ailments are the most common reason for emergency admission to hospital, the most common reason to visit the GP, and cost the NHS more than any other disease area. This pocket-sized handbook allows instant access to a wealth of information needed in the day-to-day practice of respiratory medicine.

Cancer Biomarkers

Ethics, Economics and Society

Megaloceros Forlag Cancer care is undergoing a shift from a 'one-size-fits all' approach to more personalised medicine. One way of personalising cancer treatments is through biomarkers: molecules or biochemical changes found in the patient's tissues and body fluids. This book reflects upon the promise of cancer biomarkers and asks questions such as: How may the complexity of cancer biology impede the robustness of biomarkers in the clinic? How should one draw the line between the various sub-groups of patients for personalised treatment? How can one evaluate the cost-effectiveness and fairness of personalised cancer treatments? By bringing together authors from the fields of science and technology studies, medical ethics and philosophy, health economics and oncology, the book aims to give a critical yet accessible overview of some of the key social, ethical and economic issues that surround cancer biomarkers. "The book should be required reading for oncologists, medical students, graduate students and especially for those who make policy decisions regarding the use and reimbursement of cancer biomarkers." - Bruce Zetter, Charles Nowiszewski Professor of Cancer Biology in the Department of Surgery, Harvard Medical School

100 Questions & Answers About Advanced & Metastatic Breast Cancer

Jones & Bartlett Publishers Companion to: 100 questions & answers about advanced and metastatic breast cancer / by Lillie D. Shockney and Gary R. Shapiro. 2nd ed. c2012.

Ethical Challenges for Healthcare Practices at the End of Life: Interdisciplinary

Perspectives

Springer Nature This book presents a collection of exclusively selected manuscripts on current ethical controversies related to professional practices from an interprofessional perspective. Insights are provided into the diversity of practices and viewpoints from different countries are merged in a unique way. The book contributes to the debate on social and legal issues regarding end-of-life practices such as organ donation, medically assisted dying and advance care planning. In addition, joint international author groups contributed exclusive chapters about European comparisons on end-of-life topics. The focus on country- and culture-specific aspects broadens the view on key issues and makes the book attractive for an international readership. The variety of approaches and methods used informs and inspires the development of new research and best-practice projects.

Prognosis in Advanced Cancer

Oxford University Press Predicting survival and other outcomes is increasingly being recognized as an important skill for palliative care physicians, internists, and other health care professionals who care for patients with advanced cancer at the end of life. There is much prognostic information available that is scattered throughout the palliative care and oncological literature but this is the first time it has been gathered systematically in one place. This book has 36 chapters divided into three sections. The first is an introductory section that deals with the principles of prognostication, including formulating the prediction and then communicating it. Topics such as statistical issues, evidence based medicine, and the ethics of prognostication are also covered. The second section addresses prognostication in fifteen specific cancers when they have reached the advanced stage. The third section deals with prognostication in patients with a variety of common clinical conditions at the end of life, such as bowel obstruction, hypercalcemia, and brain metastases.

Psychosocial Aspects of Chemotherapy in Cancer Care

The Patient, Family, and Staff

Psychology Press Leading medical professionals--physicians, nurses, social workers--who treat cancer patients receiving chemotherapy address vital areas of concern: physician/patient relationships, the psychosocial issues of being a patient, the pediatric patient, and new frontiers. Valuable and readable for health professionals and cancer patients and their families, this book deals honestly with the relevant, often painful subjects inherent in the diagnosis and treatment of cancer with chemotherapy. Contributing authors emphasize the importance of establishing a positive, trusting relationship between patient and doctor and patient's family and of treating the psychological and social aspects of patients, as well as the medical problems. The eleven patients'rights described in one chapter should be available to all cancer patients and their families who are faced with the decision of choosing chemotherapy as the course of treatment. The problems particular to treating pediatric patients are examined, along with sound advice for school personnel in dealing with cancer patients in the classroom.

Cancer Pain Management

Jones & Bartlett Learning Cancer Pain Management, Second Edition will substantially advance pain education. The unique combination of authors -- an educator, a leading practitioner and administrator, and a research scientist -- provides comprehensive, authoritative coverage in addressing this important aspect of cancer care. The contributors, acknowledged experts in their areas, address a wide scope of issues. Educating health care providers to better assess and manage pain and improve patients' and families' coping strategies are primary goals of this book. Developing research-based clinical guidelines and increasing funding for research is also covered. Ethical issues surrounding pain management and health policy implications are also explored.

Individualized Medicine Between Hype and Hope

Exploring Ethical and Societal Challenges for Healthcare

LIT Verlag Münster "Individualized medicine" is a catchphrase currently used to denote efforts in medical research and practice to establish tailored healthcare. The vision of "personalized" medicine has proved to be highly ambivalent, reflecting hype and hope - compared to the great expectations only very few applications have been realized up to now. The contributions to this volume discuss the challenges for patients, doctors, and the healthcare system and examine ethical and societal issues arising from one the most promising and most controversial developments in medical science and biotechnology. (Series: Medizin und Gesellschaft - Vol. 19)