

Pediatric Palliative And Hospice Care Pdf Pdf

Perinatal Palliative Care

2020-02-05 Erin M. Denney-Koelsch This unique book is a first-of-its-kind resource that comprehensively covers each facet and challenge of providing optimal perinatal palliative care. Designed for a wide and multi-disciplinary audience, the subjects covered range from theoretical to the clinical and the practically relevant, and all chapters include case studies that provide real-world scenarios as additional teaching tools for the reader. *Perinatal Palliative Care: A Clinical Guide* is divided into four sections. Part One provides the foundation, covering an overview of the field, key theories that guide the practice of perinatal palliative care, and includes a discussion of perinatal ethics and parental experiences and needs upon receiving a life-limiting fetal diagnosis. Part Two delves further into practical clinical care, guiding readers through issues of obstetrical management, genetic counseling, neonatal pain management, non-pain symptom management, spiritual care, and perinatal bereavement care. Part Three discusses models of perinatal palliative care, closely examining evidence for different types of PPC programs: from hospital-based programs, to community-based care, and examines issues of interdisciplinary PPC care coordination, birth planning, and team support. Finally, Part Four concludes the book with a close look at special considerations in the field. In this section, racial, ethnic, and cultural perspectives and implications for PPC are discussed, along with lessons in how to provide PPC for a wide-range of clinical and other healthcare workers. The book closes with a look to the future of the field of perinatal palliative care. Thorough and practical, *Perinatal Palliative Care: A Clinical Guide* is an ideal resource for any healthcare practitioner

working with these vulnerable patient populations, from palliative care specialists, to obstetricians, midwives, neonatologists, hospice providers, nurses, doulas, social workers, chaplains, therapists, ethicists, and child life specialists.

Pediatric Palliative Care

2015-08-31 Betty R. Ferrell 'Pediatric Palliative Care', the fourth volume in the 'HPNA Palliative Nursing Manuals' series, addresses paediatric hospice, symptom management, paediatric pain, the neonatal intensive care unit, transitioning goals of care between the emergency department and intensive care unit, and grief and bereavement in paediatric palliative care.

Oxford Textbook of Palliative Care for Children

2012-05-24 Ann Goldman Comprehensive in scope and definitive in authority, this second edition has been thoroughly updated to cover new practices, current epidemiological data and the evolving models that support the delivery of palliative medicine to children. This book is an essential resource for anyone who works with children worldwide.

Interdisciplinary Pediatric Palliative Care

2022 Joanne Wolfe "We hope that the lives of all children will be filled with possibility, with open horizons and rainbows into the future. Children with serious illnesses, their families, and those who care for them, confront the realization that "not everything is possible," that despite dramatic scientific and medical advances, the lifespan of some children will be shortened. This threat of premature loss heightens the sense of time for children and families alike, and challenges clinicians to create new pathways of hope for them"--

Pediatric Palliative Care

2021 Betty Davies "Pediatric Palliative Care: A Model for Exemplary Practice lays out a road map for healthcare providers interested in optimizing care for seriously ill children and their families. Grounded in clinical practice and the study of positive rather than problematic encounters between providers and parents, the book presents an evidence-based model of exemplary interaction. Chapters offer a clear understanding of the complex, holistic process of interaction between providers and parents, as well as the personal and professional knowledge and skills needed to interact in optimal ways. This is a one-of-a-kind guidebook for healthcare providers interested in (re)discovering how to maximize positive outcomes for both families and providers. It's also a valuable source of inspiration for educators, supervisors, and hospital administrators who want to facilitate personal and professional development and create supportive environments for students, providers, seriously ill children, and their families"--

When Children Die

2003-02-09 Institute of Medicine The death of a child is a special sorrow. No matter the circumstances, a child's death is a life-altering experience. Except for the child who dies suddenly and without forewarning, physicians, nurses, and other medical personnel usually play a central role in the lives of children who die and their families. At best, these professionals will exemplify "medicine with a heart." At worst, families' encounters with the health care system will leave them with enduring painful memories, anger, and regrets. *When Children Die* examines what we know about the needs of these children and their families, the extent to which such needs are—and are not—being met, and what can be done to provide more competent, compassionate, and consistent care. The book offers recommendations for involving child patients in treatment decisions, communicating with parents, strengthening the organization and delivery of services, developing support programs for bereaved families, improving public and private insurance, training health professionals, and more. It argues that taking these steps will improve the care of children who

survive as well as those who do not—and will likewise help all families who suffer with their seriously ill or injured child. Featuring illustrative case histories, the book discusses patterns of childhood death and explores the basic elements of physical, emotional, spiritual, and practical care for children and families experiencing a child's life-threatening illness or injury.

Dying in America

2015-03-19 Institute of Medicine 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.

Comprehensive Cancer Care for Children and Their Families

2015-08-31 National Academies of Sciences, Engineering, and Medicine Childhood cancer is an area of oncology that has seen both remarkable progress as well as substantial continuing challenges. While survival rates for some pediatric cancers present a story of success, for many types of pediatric cancers, little progress has been made. Many cancer treatments are known to cause not only significant acute side effects, but also lead to numerous long-term health risks and reduced quality of life. Even in cases where the cancer is considered curable, the consequences of treatment present substantial long-term health and psychosocial concerns for children, their families, their communities, and our health system. To examine specific opportunities and suggestions for driving optimal care delivery supporting survival with high quality of life, the National Cancer Policy Forum of the Institute of Medicine and the American Cancer Society co-hosted a workshop which convened experts and members of the public on March 9 and 10, 2015. At this workshop, clinicians and researchers in pediatric oncology, palliative, and psychosocial care, along with representatives from the U.S. Food and Drug Administration, National Cancer Institute, Children's Oncology Group, pharmaceutical companies, and patient advocacy organizations, discussed and developed a menu of options for action to improve research, quality of care, and outcomes for pediatric cancer patients and their families. In addition, parents of children with cancer and pediatric cancer survivors shared their experiences with care and provided poignant personal perspectives on specific quality of life concerns and support needs for children and families across the life spectrum. This report summarizes the presentations and discussion of the workshop.

Palliative Care in Pediatric Oncology

2017-12-20 Joanne Wolfe This textbook is the first to focus on comprehensive interdisciplinary care approaches aimed at enhancing the

wellbeing of children with cancer and their families throughout the illness experience. Among the topics addressed are the epidemiology of pediatric cancer distress, including physical, emotional, social, and spiritual dimensions; the role of the interdisciplinary team; communication and advance care planning; symptom prevention and management; care at the end of life; family bereavement care; and approaches to ease clinician distress. The contributing authors are true experts and provide guidance based on the highest available level of evidence in the field. The book has not only an interdisciplinary but also an international perspective; it will appeal globally to all clinicians caring for children with cancer, including physicians, nurses, psychosocial clinicians, and chaplains, among others.

Hospice Care for Children

2001 Ann Armstrong-Dailey Children with life-threatening and terminal illnesses--and their families-- require a unique kind of care to meet a wide variety of needs. This book, the first edition of which won the 1993 Pediatric Nursing Book of the Year Award, provides an authoritative source for the many people involved in caring for dying children. It draws together contributions from leading authorities in a comprehensive, fully up-to-date resource, with an emphasis on practical topics that can be put to immediate use. The book covers the entire range of issues related to the hospice environment and is intended for all those who participate in the hospice-care process: physicians, nurses, social workers, teachers, clergy, family therapists, parents, and community service volunteers.

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