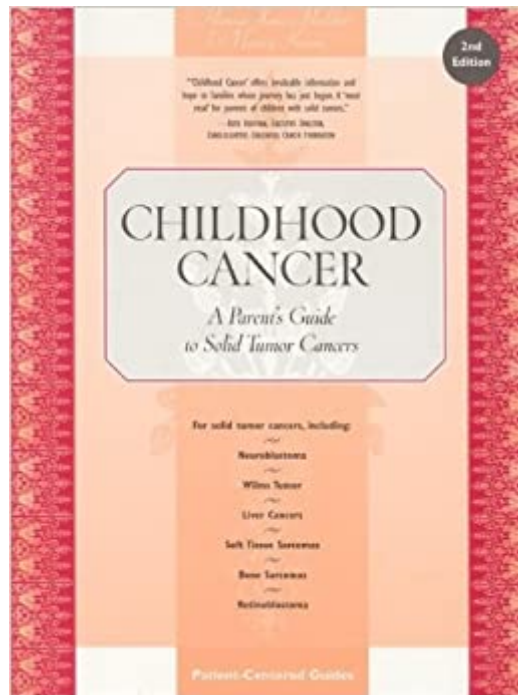


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# Childhood Cancer: A Parent's Guide To Solid Tumor Cancers, 2nd Edition



## Synopsis

This second edition of the most complete parent guide available, features detailed and precise medical information about solid tumor childhood cancers, including neuroblastoma, Wilms tumor, liver tumors, soft tissue sarcomas, and bone sarcomas. In addition, it offers day-to-day practical advice on how to cope with procedures, hospitalization, family and friends, school, social and financial issues, communication, feelings, and, if therapy is not successful, the difficult issues of death and bereavement. Woven among the medical details and the practical advice are the voices of parents and children who have lived with cancer and its treatments. As many parents have already found, advice from "veteran" parents can be a lifeline. Obtaining a basic understanding of topics such as medical terminology, how drugs work, common side effects of chemotherapy, and how to work more effectively with medical personnel can only improve the quality of life for the whole family suffering along with their child. Having parents describe their own emotional ups and downs, how they coped, and how they molded their family life around hospitalizations can be a tremendous comfort. Just knowing that there are other kids on chemotherapy who refuse to eat anything but tacos or who have frequent rages can make one feel less alone. Parents who read this book will encounter medical facts simply explained, advice to ease their daily lives, and tools to be strong advocates for their child. It also contains a personal treatment summary and long-term follow-up guide for your child to keep as a permanent record.

## Book Information

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## Customer Reviews

Each year, 11,000 children and adolescents in this country are diagnosed with cancer. Although numerous medical textbooks have been written on this subject, not many are aimed at parents and families. The authors, both mothers of children who had cancer and patient advocates in the Children's Oncology Group (Keene is also the author of *Childhood Leukemia*), fill in the gap with this guide to six solid-tumor childhood cancers: neuroblastoma, Wilms' tumor, liver tumors, soft tissue sarcomas, retinoblastoma, and bone sarcomas. The description for each tumor covers tumor origins, diagnosis, treatment, and prognosis. There are also chapters explaining clinical trials, surgery, radiation therapy, and chemotherapy in general. Emotional and psychological issues are addressed as well, with personal accounts to help parents and families cope with daily stress or in the event of death. This practical guide will empower parents to understand the disease better and participate more actively in the treatment process. Recommended for all consumer health collections.

Lily Liu, Arkansas Children's Hosp. Medical Lib., Little Rock Copyright 1999 Reed Business Information, Inc. --This text refers to an out of print or unavailable edition of this title.

Honna Janes-Hodder's youngest son Matthew was diagnosed with neuroblastoma on his third birthday, and passed away at age seven on September 3, 1997. She loved, cared for, and advocated for Matthew during his long struggle with cancer. She lives with her husband, 14-year-old son, and 11-year-old daughter in beautiful Paradise, Newfoundland, and Labrador, Canada. Honna is founder of ChildCan, The Childhood Cancer Research Association of Newfoundland and Labrador, Inc., and president of the International Pediatric Cancer Alliance, Inc. Honna manages most of the pediatric discussion groups for ACOR, the Association of Cancer Online Resources, Inc. ACOR, based in New York, provides accurate and timely medical support and information for all those affected by cancer, in both the pediatric and adult communities. She is the administrator of five online discussion groups, providing support and reliable information to families dealing with childhood cancers globally. In addition, Honna sits on the Public Issues Committee for the Newfoundland and Labrador division of the Canadian Cancer Society and writes a regular health section for a national newspaper in Canada. Honna spends a great deal of time advocating on behalf of children diagnosed with cancer. She does not view this as a job, but as an important part of her identity. Honna attends Memorial University of Newfoundland as a part-time student when time permits, working toward a Ph.D. in biochemistry. Honna and Nancy were two of the first five people appointed as patient advocates to the Children's Cancer Group (CCG), a research organization consisting of pediatric cancer specialists from Canada, the United States, and Australia. They attend CCG meetings and facilitate communication between CCG investigators and

the patient community. Nancy Keene has written or co-authored five health books for families. She is co-creator of the Patient-Centered Guides series and Editor of pediatric titles. Nancy is a tireless advocate for children's health issues, including pediatric clinical trials, childhood cancer, late effects of childhood cancer treatments, pediatric pain relief, and emotional support for families. She frequently speaks for professional and parent groups and works on/with national pediatric advocacy organizations and committees.

Helped me to understand what is going on and what to expect in the future.

Didn't receive it. discontinued

I used this book for a research paper. It had plenty of information that I needed and I donated it to the local library when class ended

Candlelighters Childhood Cancer Foundation fully endorses this book and places it on the 'must have resource list' for any parent whose child is diagnosed with any type of solid tumor cancer. This comprehensive guide provides extensive information on the diagnosis, prognosis, procedures, treatment and side effects of neuroblastoma, Wilms Tumor, retinoblastoma, rhabdomyosarcoma and bone sarcomas. Included is a thorough explanation of the various chemotherapy drugs used in treatment and their common side effects. Detailed information on radiation therapy, surgery and bone marrow transplantation is also included. Additionally, the book provides practical advice to families on related issues such as nutrition, returning to school and the end of treatment. The book addresses the emotional impact that such a cancer diagnosis places upon the child, their family and friends and offers practical insight on how to cope. A comprehensive list of childhood cancer websites, support organizations and additional reading materials is listed at the back of the book. The authors' extensive interviewing of families who have 'walked down this road' offers hope to those whose 'journey' has just begun.

This book is filled with loads of good information in lay terms to help family members, friends, teachers, social workers, etc. understand the various childhood cancers and their often complex treatments. Every family having a child diagnosed with cancer should automatically be given this book, because it contains so much information on diagnosis, procedures, tests, treatments and side effects. It also realistically discusses the potential effects on a whole family when a child is

diagnosed with cancer, and provides tips on how to have an active and constructive relationship with doctors and nurses. The authors apparently interviewed a lot of parents who've been down this road and you can read their ideas and suggestions about how to deal with all sorts of issues from the mundane (how to get your child to take a certain medication) to the tremendously difficult (how to deal with relapse or talk to a child about death). I wish this had been available when my child was first diagnosed!

If you have found yourself catapulted into the nightmare of having a child with cancer, this book will help you to understand much of the journey to come. No matter what stage your child is in treatment...beginning, middle, finished treatment, relapsed, or palliative...this book is an indispensable resource. Personal stories peppered throughout the book help you come to realize that you are not alone in this battle. This book also has a wonderful list of resources to help families through this hard time. Buy one for yourself, family members (especially in-laws!), your child's pediatrician, your child's teacher, and your closest friends so they can have a glimpse into what you and your child are facing and can help you along the way. This book should be given to EVERY parent when they hear those horrible words, "Your child has cancer..."

This book is amazing. I keep it handy at home to help answer my questions or learn more about the disease. My daughter was diagnosed with retinoblastoma right before Christmas and this book was given to us by a social worker in the hospital. There is an entire chapter about retinoblastoma as well as chapters on many other solid tumor cancers. There are real stories from parents and it is written in language that is informative but easy to understand. It is especially helpful right after diagnosis when all of the jargon is still new. I would caution people to take the advice given in the book and only read sections that are relevant to your situation at the time.

I cannot overstate how helpful this book was to me. My daughter was diagnosed with Wilms (kidney) cancer 14 months ago. This book was given to me several days after her initial diagnosis and I turned to it many, many times over the long journey to recovery. Even today, with my daughter happily in remission, I turn to it for help in dealing with late side effects and the emotional consequences of the rollercoaster that is childhood cancer. Of all the books I have read over the past 14 months (and there have been many - I'm an avid reader), this one was absolutely the best. If I could, I would buy up all the copies and give it to every parent of a child with a solid tumor.

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Fighting Foods And How The Spices Turmeric, Ginger And Garlic Can Reduce Cancer Risk  
(Essential Spices and Herbs Book 4) Childhood Leukemia: A Guide for Families, Friends &  
Caregivers (Childhood Cancer Guides)

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