

[DATE]

Dear Representative:

As a lymphoma advocate with the Lymphoma Research Foundation and as your constituent, I support efforts to ensure that all cancer patients have access to cancer care of high quality. Expert reports have identified critical elements of a high quality cancer care system. At the core of such a system is the coordination of active treatment with symptom management. Patients who receive this kind of care experience the appropriate management of the symptoms of cancer and its treatment at the same time they are getting the best possible therapy for their disease.

Cancer treatment is often a very difficult experience, but steps can be taken to be sure that the side effects of treatment are well managed. We are fortunate that Representatives Lois Capps (D-CA) and Tom Davis (R-VA) have introduced legislation that takes important steps toward reforming the cancer care system. The Capps-Davis bill would authorize Medicare payment for services to plan and summarize cancer care and would also establish pilot programs of coordinated care and systems for training cancer care specialists in coordinating cancer care.

I urge you to lend your support to the Comprehensive Cancer Care Improvement Act, H.R. 5465. This bill represents a critical first step toward a better system of cancer care, and your support of this measure is important to me and other cancer survivors.

Thank you in advance for considering this bill. I look forward to hearing that you will support the Comprehensive Cancer Care Improvement Act.

Sincerely,

# Sample Letter For Cancer Support

**Robert DeBellis**



## **Sample Letter For Cancer Support:**

**Cancer Care for the Whole Patient** Institute of Medicine, Board on Health Care Services, Committee on Psychosocial Services to Cancer Patients/Families in a Community Setting, 2008-03-19 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 *A Physician's Guide to Pain and Symptom Management in Cancer Patients* Janet L. Abrahm, 2015-01-01 This highly regarded handbook provides clinicians with the information they need to treat their cancer patients effectively and compassionately This comprehensive guide to managing pain and other symptoms for people with cancer has helped tens of thousands of patients and families Designed for busy practicing clinicians *A Physician's Guide to Pain and Symptom Management in Cancer Patients* provides primary care physicians advanced practice nurses internists and oncologists with detailed information and advice for alleviating the stress and pain of patients and family members alike Drawing on the work of experts who have developed revolutionary approaches to symptom management and palliative care as well as on the lessons learned from patients and their families during her thirty years as a teacher and clinician Dr Janet L Abrahm shows how physicians and other caregivers can help patients and families heal emotionally even as the disease progresses The third edition includes updates to medications and clinical stories and features two new chapters Working with Patients Families and Sexuality Intimacy and Cancer New lessons from palliative care and hospice care can help patients their professional caregivers and their families support each other every step of the way Cancer Control J. Mark Elwood, Simon B. Sutcliffe, 2010-01-07 Cancer control is the term applied to the development of integrated population based approaches to reduce the incidence and mortality from cancer and to minimize its impact on affected individuals and on the community It covers a spectrum of prevention early diagnosis optimal treatment and supportive and palliative care It emphasizes the application of new knowledge gained through research to achieve current best practice Cancer control has become a political priority in many countries in recent years with the evolution of both national and regional cancer control strategic plans The integrated nature of cancer control involving a wide spectrum

of health care professionals researchers and health managers and planners is reflected in this multi disciplinary text which is the first in this rapidly developing field      Evidence-Based Cancer Care and Prevention Charles W. Given, PhD, Barbara Given, RN, PhD, FAAN, Sharon Kozachik, MSN, RN, MS, Danielle Nicole DeVoss, PhD, 2003-03-01 The goal of behavioral oncology is to understand and explain the role and impact of behaviors at all phases in the cancer trajectory from prevention and detection to diagnosis and early treatment to survivorship recurrence and or death Each chapter includes summaries of recent research on cancer related behavioral interventions discussions of the studies summarized and suggestions for future research The book is a product of collaboration among members of the Behavioral Cooperative Oncology Group of the Mary Margaret Walther Program for Cancer Care Research at the Walther Cancer Institute in Indianapolis Indiana      Handbook of Cancer Survivorship Michael Feuerstein, 2007-10-18 Not long ago a cancer diagnosis was regarded as an automatic death sentence today there are ten million survivors Equally impressive is the growing number of clinicians and researchers dedicated to improving the quality of survivors lives and care Yet despite this encouraging picture there has never been a reliable central source for relevant clinical information till now The Handbook on Cancer Survivorship responds to the diverse needs of survivors and their support communities by comprehensively addressing the major issues in the field from the burden of survivorship to secondary prevention Editor Michael Feuerstein himself a cancer survivor and sixty other top scientist practitioners analyze in depth how survivors meet and manage the challenges of life after cancer and what clinicians researchers and public health systems can do to ease the transition The Handbook s 27 comprehensive chapters include the latest research and practice related to Survivors quality of life and how it can be assessed Managing everyday and chronic stress Depression anxiety pain disorders and cognitive changes Coping adaptation and resilience Behavior change strategies exercise weight control smoking cessation Cancer survivorship centers and other models for follow up care Survivor clinician and international perspectives New frontiers in practice research and policy Such wide ranging coverage benefits everyone involved in cancer survival primary care providers oncologists behavioral health specialists physical and occupational therapists nutritionists epidemiologists health systems professionals and policymakers and of course survivors themselves and their families      **The Politics of Breast Cancer** Maureen Hogan Casamayou, 2001-03-26 Between 1990 and 1993 breast cancer activism became a significant political movement The issue began to receive extensive media attention and federal funding for breast cancer research jumped dramatically Describing the origins of this surge in interest Maureen Hogan Casamayou attributes it to the emergence of politically potent activism among breast cancer survivors and their supporters Exploring the creation and development of the National Breast Cancer Coalition NBCC she shows how many of its key leaders were mobilized by their own traumatic experiences with the disease and its treatments Casamayou details the NBCC s meteoric rise and impressive lobbying efforts explaining how in contrast to grassroots movements founded by dedicated individuals the coalition grew from the simultaneous efforts of a network of women who invested their time energy

money and professional skills in the fight for increased funding for breast cancer research This multiple leadership or collective entrepreneurialism says Casamayou was crucial to the NBCC's success framing the issue in the minds of the public and policymakers alike *Cumulated Index Medicus* ,1995 *Index Medicus* ,2004 Vols for 1963 include as pt 2 of the Jan issue Medical subject headings *Effect of Cancer On Quality of Life* David Osoba,2021-11-01 This book is comprised of extensive reviews and instructional chapters that discuss the quality of life in several aspects of cancer The first six chapters deal with conceptual issues relating to measuring quality of life in adult and pediatric populations with cancer The next five chapters provide practical information on how to select quality of life measures the statistical analysis of trials economic evaluations to be considered and some possible abuses of quality of life measures Five chapters review the results of studies using selected quality of life measures and provide recent information on their performance These are followed by three chapters dealing with specific issues relating to nausea and vomiting associated with cancer therapy Three chapters are devoted to the problems of assessing and controlling pain in patients who have cancer There are also two chapters that deal with the quality of life in palliative care *Effect of Cancer on Quality of Life* is intended for all who have an interest in measuring the quality of life in patients with cancer This includes investigators who are just entering the field and can benefit from instructions on how to conduct quality of life research as well as those who are experienced in conducting this kind of research *Psychosocial Aspects of Chemotherapy in Cancer Care* Robert DeBellis,1987 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

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