



Multiple Primary Malignancies (Updates in Surgery)

By Andrea Renda

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In 2006, when my colleague Andrea Renda proposed multiple primary malignancies (MPM) as the subject of the Biennial Report to the 2008 Congress of the Italian Society of Surgery, I, together with the Steering Committee, quickly agreed. Recent progress in our understanding of the etiopathology of these neoplasms has led to innovative and significant progress on the clinical level. Importantly, the incidence of the onset of two or more tumors in the same patient suggests a more than casual relationship. Furthermore, the occurrence of MPM derives from several different mechanisms: viral, iatrogenic, immunologic, environmental, and hereditary, such that any form of treatment must take into account the etiology of these tumors. After an epidemiological introduction, this monograph analyzes various aspects of multitumoral syndromes based on the experience of the Department of Surgical Sciences, along with that of other clinical departments of the University Federico II of Naples. In the discussion of inherited tumors, reference is made to the series of patients treated at the Department of Surgery at the University of Siena. The many topics that comprise this volume range from carcinogenesis to diagnostic strategies, and from epidemiology to innovations in imaging and endoscopic techniques. Among the clinical aspects, particular emphasis is given to sporadic and hereditary syndromes, as these patients are frequently treated by general surgery departments.

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Editorial Review

Review

From the reviews:

"This volume ... focus casual associations rather than chance in multiple primary malignancies. ... Practitioners and researchers in surgery, oncology, and allied fields are the intended audience. ... It will be of most interest to those with a research or clinical interest in multiple primary tumors occurring through circumstance ... rather than chance. ... This is an excellent reference with particular strength in the area of hereditary syndromes. ... Other workers in the field will find it of interest." (Carol Scott-Conner, Doody's Review Service, July, 2009)

From the Back Cover

Two to three decades ago, multiple primary malignancies (MPM) were considered to be simply a clinical curiosity, with sporadic reports, mostly single case descriptions, found in the literature. While in the last few years, analyses of larger series have been published, the number of cases has still been relatively small and most of the reports have addressed a single type of primary tumor and its associations.

With the improved prognosis and survival of cancer patients, MPM is becoming increasingly prevalent in this population, necessitating a better understanding of the characteristics and associations of the malignancies involved. Thus, as a prerequisite, a universal definition and an internationally accepted classification system, based on chronological, pathological, clinical, and other parameters, are needed. Moreover, much remains to be learned about the etiology of MPM, whether genetic, iatrogenic, or environmental.

Several of the hereditary syndromes, such as familial adenomatous polyposis, hereditary non-polyposis colorectal cancer, hereditary breast-ovarian cancer, and multiple endocrine neoplasia, are already well-known and their characteristics in relation to MPM must be kept in mind. Nonetheless, along with these syndromes, there are sporadic and apparently casual associations between primary neoplasms that can involve almost any part of the body.

This volume points out the clinical aspects of MPM and discusses the diagnostic and therapeutic problems that are encountered in treating these patients. "DNA-guided" surgery, currently confined to the treatment of patients with hereditary syndromes, will, along with other novel treatment strategies, no doubt play an increasingly greater role in the therapy of MPM.

This work presents state-of-the-art information about MPM that is aimed at a broad range of medical specialists, including surgeons, endoscopists, oncologists, and geneticists,. The goal is to improve our understanding of this group of diseases as well the treatment of these patients.

Users Review

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