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A Regency-era romantic adventure where a Duke is ordered to assume guardianship over a bold young woman whose refuses to believe her parents lives were lost during a treasure hunt. The first in a three-book series.

This is a comprehensive, lavishly illustrated review of orthopaedic oncology that is multi-disciplinary, integrative and highly practical. Introductory chapters cover overviews of the basics of bone and soft tissue tumors - the pathology, radiology and surgical and oncologic management. These are followed by chapters covering all tumor types organized according to the 2013 WHO classification of bone and soft tissue tumors. Each class of tumor is dealt with in detail, with case presentations illustrating how they would be diagnosed and managed in a clinical setting. The range of variation of each tumor type is indicated with appropriate radiological and pathological images. Detailed tables and flow charts provide quick access to vital diagnosis and staging information for each tumor type. This book provides an expert, case-based approach to orthopaedic oncology and

patient care for trainees and practising orthopaedic surgeons, pathologists and radiologists. A practical and comprehensive review of orthopaedic oncology Covers all tumor types and their range of variation Organized according to the 2013 WHO classification of bone and soft tissue tumours Tables and flow charts provide diagnosis and staging information for each tumor type

This volume is a comprehensive guide to the diagnosis and treatment of benign and malignant tumors of the bone and soft tissues. Coverage includes: general issues in diagnosis, staging, surgery, radiation therapy, chemotherapy, and the psychological aspects of patient management. Focus is placed on benign bone tumors, malignant bone tumors, soft-tissue tumors, and carcinoma metastatic to bone. The sections on malignant bone tumors and carcinoma metastatic to bone include detailed guidelines for surgical treatment in each anatomic site. Special features include: considerations for management of malignant bone tumors in growing children; and discussions on the biology and complications of prosthetic and allograft reconstruction.

Sarcomas represent a group of rare tumors which arise from the largest tissue compartment of the body, but account only for about 1 % of all malignancies. Among this group of tumors, soft tissue sarcomas in adults are, in addition to osteosarcomas, the most important. In the diagnostic workup of soft tissue sarcomas modern radiological investigation methods such as CT and MRI techniques are being critically evaluated. While it is widely accepted that the conventional, stained microscopic specimen is still the cornerstone for properly planned treatment strategies, immunohistochemistry has come to play an indispensable role in accurate classification. Reviewing the surgical techniques, the treatment of musculo-skeletal malignancies, especially of soft tissue sarcomas, has changed considerably over the past decades. While amputation used to be the surgical treatment of choice limb-saving procedures have now become an important therapeutic modality in treating such lesions, particularly in combination with percutaneous or interstitial radiotherapy. Despite adequate local treatment, many patients with soft tissue sarcomas will ultimately develop metastatic disease, usually in the lungs. For this reason, effective adjuvant systemic treatment should be given simultaneously with or soon after treatment of local disease. The results of several randomized trials evaluating adjuvant chemotherapy have been reported in the meanwhile, indicating improved, disease-free survival with a trend towards improved overall survival. In disseminated disease, drug combinations or single agents are used as palliative treatment and might help to improve survival in selected subgroups.