Microbiological Profile of Prosthetic Joint Infections in Orthopaedic Oncology

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INTRODUCTION: Periprosthetic joint infections (PJI) are a devastating complication after reconstruction with megaprosthesis in orthopaedic oncology. PJI severity is further compounded by the unique immunologic considerations in oncology–chronic immunodeficiency, systemic and radiation therapy, increased surgical complexity–as well as tumor-microbiome interactions that may put patients at increased risk. Despite significant efforts to characterize the microbiologic profile of PJI in traditional joint arthroplasty, data is lacking for oncologic patients with megaprosthesis. The objective of this study was to characterize causative microorganisms and time-to-positivity (TTP) of PJI in oncologic patients at our institution, with subanalyses for cancer type and primary tumor characteristics.

METHODS: A retrospective review of our institutional megaprosthesis database was conducted. We reviewed tissue and fluid cultures for patients diagnosed with PJI-defined using the 2011 Musculoskeletal Infection Society criteria-between the years 2000 and 2022. All positive causative microorganisms, including the number of samples, specimen type, identified pathogen, and TTP, were recorded and assessed, along with basic oncologic data. Median and interquartile ranges were used to describe continuous variables, and comparisons were made using the Mann-Whitney U test.

RESULTS: We included 75 patients diagnosed with megaprosthesis PJIs (Table 1). Patients primarily consisted of those with a McPherson host grade B (84%) and metastatic bone disease (85%), and half of patients received prior radiation therapy (Table 1). PJI classification of acute versus chronic PJI was relatively balanced (42% vs. 57%), and in most cases, patients underwent proximal (31%) or distal (41%) femur replacement (Table 2). The six most prevalent pathogens isolated in cultures were *Staphylococcus epidermidis* (n=63), *Staphylococcus aureus* (n=48), *Enterococcus spp.* (n=35), *Streptococcus spp.* (n=32), *Coagulase negative staphylococcus* (n=21) and *Gram-negative rods* (n=20) (Figure 1). Differences in microorganism prevalence were observed between primary vs. metastatic bone disease (Figure 2A), bone vs. soft tissue sarcoma (Figure 2B), and bone tumor histology (Figure 2C). TTP varied among microorganisms and differed significantly (p <0.05) between gram-positive vs. gram-negative bacteria (Figure 3A), culture origin (bone, soft tissue, or synovial fluid) (Figure 3B), and culture method (broth only vs. broth and solid media) (Figure 3C). Of the more common pathogens, *Streptococcus epidermidis, Streptococcus lugdunensis, Finegoldia magna,* and *Corynebacterium* had the longest TTPs (Figure 4). TTP varied widely among microorganism species (Table 3).

DISCUSSION AND CONCLUSION: While the causative microorganisms in oncologic PJI were broadly similar to those in traditional joint arthroplasty literature, there are pointed differences in prevalence and TTP by subpopulation. These differences may be the result of cancer-driven microbiome disruptions in oncologic patients. Cultures in megaprosthesis PJI should be held for at least two weeks if positivity is not achieved sooner. Further investigation is needed to understand the mechanisms behind these differences to provide prompt, individualized, and targeted antimicrobial therapy for this atrisk

