**Introduction** Chronic nonbacterial osteomyelitis is a rare autoimmune inflammatory disease of unknown etiology that primarily affects bone and presents with sterile inflammation of the most common metaphyseal areas of the long bones, clavicle, spine, and pelvis. Affected bones are prone to pathologic fractures and slow growth. Laboratory workup results are nonspecific, and skeletal scintigraphy with technetium and magnetic resonance imaging are used as diagnostic imaging modalities. Differentially, malignant events must be considered, such as leukemia and malignant bone tumors, chronic infectious osteomyelitis, avascular necrosis, etc. If the diagnosis is unclear, a bone biopsy with bacteriological analysis of the bone sample is required to exclude infectious osteomyelitis.

**Case Report** An 8-year-old girl was treated for acute lymphoblastic leukemia of T immunophenotype, high risk, according to ALL IC-BFM 2009 protocol, early complete remission was achieved. Intensive chemotherapy treatment was characterized by a number of complications, including frequent febrile neutropenia, invasive fungal lung disease, and acute neurotoxicity in the form of cerebral ischemia. After completion of reinduction therapy, a period of persistent bone marrow aplasia followed by recurrent and prolonged sepsis caused by a highly resistant strain of Pseudomonas aeruginosa of unknown origin preceded maintenance therapy. Because the girl complained of foot pain with inability to walk during maintenance therapy, extensive diagnostic workup was performed, including technetium bone scintigraphy and magnetic resonance imaging, to verify an osteomyelitis focus in the left calcaneus with intense bone remodeling of SI and shoulder joint. After completion of treatment for sepsis in the control laboratory findings continued slightly elevated C-reactive protein with normal sedimentation of erythrocytes and procalcitonin. Pathohistological analysis of bone biopsy showed chronic inflammatory reaction, while microbiological analysis was negative. Skin lesions were not observed, HLA typing was negative, anamnestic data of maternal inflammatory bowel disease was known. In collaboration with an immunoneumatologist and an orthopedist, chronic nonbacterial osteomyelitis was treated and treatment with bisphosphonates was started. According to a standard regimen, with an excellent clinical response. The girl was pain free six months after starting treatment and physical therapy. The underlying malignancy is still in remission, inflammatory parameters are normal, while control imaging still continues.

**Conclusion** Chronic nonbacterial osteomyelitis should be considered as a differential diagnosis in chronic inflammatory bone lesions without an isolated microbiologic agent and in patients with acute lymphocytic leukemia. Therapeutic options include nonsteroidal anti-inflammatory drugs, disease-modifying antirheumatic drugs (DMARDs), corticosteroids, and bisphosphonates. However, given the possibility of influencing hematologic findings, bisphosphate therapy is the treatment of choice if oncologic disease requires further treatment.

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**Abstracts**

**323** **TREATMENT RESULTS OF T-LYMPHOBLASTIC LYMPHOMA IN CHILDREN – A SINGLE-CENTER EXPERIENCE**

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Aim Our study aimed to determine the overall survival of children with non-Hodgkin T-lymphoblastic lymphoma (T-LBL), analyze and present patient characteristics and initial presentation of their disease.

**Methods** Our study included all patients with T-LBL non-Hodgkin lymphoma treated in the Department of Pediatrics, Division of Hematology and Oncology, University Hospital Centre Zagreb between January 1st, 2002 and December 31st, 2017. Relevant information including general patient data (gender, age at the time of diagnosis), initial clinical presentation (the presence of mediastinal masses, pericardial and pleural effusion, the presence of bone marrow and central nervous system disease) and treatment information (used therapeutic protocol and mortality data) was collected from the available medical documentation. The Kaplan-Meier curve shows patients' survival.

**Results** Overall, 19 patients were included, 14 of which were male (73.68%) and 5 were female (26.32%). The median age at diagnosis was 10 years. 17 patients (94.44%) presented with mediastinal mass and 9 of them (52.94%) had pleural effusion. At the time of diagnosis, pericardial effusion and superior vena cava syndrome were present in 6 (33.33%) and 3 (16.67%) patients, respectively. Bone marrow involvement was detected in 10 patients (52.63%), whereas CNS involvement in only one (5.26%). 16 out of 19 patients survived (84.21%); 95% CI 67.81–100). Survival was higher for boys (85.71%) than for girls (80%), but the difference was not statistically significant (p = 0.764).

**Conclusion** Comparing general epidemiologic data with the one in the available literature we haven’t found a significant deviation of the prevalence of the disease between the sex or the median age of onset of the disease. The result of 90% five-year survival of children with T-LBL in the BFM group study was not repeated in later studies and the overall survival of 84% in our group was consistent with the results reported in the available literature.