Sarcomas in children under 5 years old: general characteristic and oncologic outcomes.


Objective: Sarcomas affecting children in the first years of life have been rarely reported in literature and represents a challenging disease. We propose to analyse a group of patients under de age of 5 years with primary sarcomas and report its oncologic outcomes.

Material and methods: Multicenter study design. A retrospective review was performed and all paediatric patients with a primary sarcoma (soft tissue sarcoma –STS- or primary bone sarcoma –PBS-) under the age of 5 years, and treated with surgery in three different sarcoma centres were analysed. Overall survival rates and oncologic prognostic factors were analyzed through Kaplan-Meir and log rank test.

Results: One-hundred twenty patients were included in the study, 41 STS and 79 PBS. The mean age of diagnosis was 3.5 years (range: 0-5) and mean follow-up 100 months (range 3-325). Ninety-six tumors were located in the extremities and 24 were classified as apendicular tumors. Sarcomas under the age of 5 year comprised 2% of all paediatric sarcomas treated in the 3 institutions involved in the study (1% for STS and 6% for PBS). The most prevalent soft tissue sarcoma was rhabdomiosarcoma (n: 23) follow by PNET (n:5) and fibrosarcoma (n:3). Ewing sarcoma (n:56) was the most common primary bone tumor follow by osteosarcoma (n:19). Limb salvage procedures were possible in 95 (79%) patients. Five year Overall Survival was 72% (95%IC: 64-84). Survival was not affected in this selected group by age at diagnosis (p=0.6), gender (p=0.7), tumor location (p=0.2) or type of tumor (STS vs PBT) (p=0.7). Tumor size over 5 cm at time of diagnosis (p=0.01) and local recurrence (p=0.002) affected significantly the oncologic prognosis.
**Conclusion:** Sarcomas in paediatric population under the age of 5 are very rare (4 referrals per year), being Rhabdomyosarcoma and Ewing sarcoma the most prevalent histological diagnosis. Five year overall survival in this particular group is over 70% and significantly affected by the size of tumor and local recurrence. Limb salvage procedure is possible in nearly 80% of the patients.