Case Report

Glomangiosarcoma *de novo* in a child: A case report and review of literature

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ABSTRACT

Glomus tumors (GT) are cutaneous and soft tissue mesenchymal neoplasms. The malignant variant that is, glomangiosarcomas (GS) are rare and account for <1% of all GT. GS are usually seen in adults with a wide age range of 20-89 years. GS usually have an indolent course with local aggressiveness. The treatment is complete surgical excision with excellent prognosis though metastasis workup is advised. Herein, we report an unusually large GS *de novo* in a 6-year-old girl. Complete surgical excision was performed. There was no evidence of recurrence or metastasis over the last 15 months of follow-up postoperatively. GS should be considered as one of the differential diagnosis of cutaneous and soft tissue neoplasms.

Key Words: Glomangiosarcoma, glomus tumor, immunohistochemistry

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