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Hepatic angiosarcoma with clinical and histological features of Kasabach-Merritt Syndrome

Sanya Wadhwa, Tae Hun Kim, Leah Lin, Gary Kanel, Takeshi Saito

Abstract

Hepatic angiosarcoma is a mesenchymal tumor originating from liver sinusoidal endothelial cells. It is an extremely rare malignant neoplasm accounting for less than 1% of primary malignant liver tumors. The deregulated coagulopathy that can be seen in hepatic angiosarcoma fulfills the clinical diagnostic criteria of disseminated intravascular coagulation. However, the mechanism that governs this coagulopathy has been poorly understood. This case report provides histological evidence of the

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