Abstract. Myxoma is rare in the upper leg. The current study presents the case of a large tumor in the right upper leg. The tumor extended to pelvic cavity and was found to be connected with a cystic and solid neoplasm that was adjacent to the ascending colon in the right lower quadrant. The large tumor of the upper leg had existed for 15 years and had ulcerated through the skin 4 days prior to admittance. Palliative surgery was performed, with histological findings of a myxoma. Since appendiceal mucinous neoplasms may invade organizations outside of the mucous layer of the appendix and cause secondary peritoneal myxoma, this myxoma of the upper leg probably originated from an appendiceal mucinous neoplasm.

Introduction

Myxomas are benign neoplasms that are characterized by slow growth (1). The most frequent location of these neoplasms is the cardiac muscle, where they account for 50% of all benign lesions (2). Myxoma is rare in the leg, but can occur within its muscles (1). To date, <10 cases of myxoma located in the leg have been documented (1,3-5). Computed tomography usually shows a well-delimited mass with an absorption density between that of muscle and water. The most precise examination technique that enables diagnosis is magnetic resonance imaging (6,7). Similarly to previous studies (1,6,9), the myxoma in the present study had a sharply defined border. In addition, it exhibited a signal intensity lower than that of the skeletal muscle.

Case report

An 88-year-old female was hospitalized at Yantai Affiliated Hospital of Binzhou Medical University (Yantai, China) in January 2014 with a large tumor in the right upper leg. The tumor had existed for 15 years and had ulcerated through the skin 4 days prior to admittance. The function and appearance of the right lower limb were seriously affected. Laboratory examinations showed no abnormalities and no other symptoms were observed. The MR studies were performed by a SIEMENS Avanto 1.5T MR scanner. A large tumor (45x15x20 cm) in the right thigh was found with a slight high signal on T1-weighted imaging (Fig. 1A) and an appreciable high signal on T2-weighted imaging (Fig. 1B). The tumor extended to the pelvic cavity and was found to be connected with a cystic and solid neoplasm (Fig. 1C). Due to the older age of the patient, total resection was considered to be too traumatic. Instead, the patient underwent palliative surgery for the tumor in the upper leg, and the tumor was found to be full of brown and yellow gelatinous material. Myxoma was finally diagnosed via histological examination (Fig. 2). Following surgery, the patient recovered well and was subsequently discharged. The patient rejected further treatment and no follow-up examination was planned. At the time of writing, the patient was alive.

This study was approved by the Ethics Committee of Binzhou Medical University (Yantai, Shandong, China) and written informed consent was obtained from the patient.

Discussion

Myxoma is rare in the upper leg (3). Similarly to a previous study (1), the growth of the myxoma in the present case was slow. A plain X-ray may be normal or rarely visualize the lesion within the soft tissues as shading with calcifications inside (9). An ultrasonographic examination usually visualizes hypoechoic lesions with fluid compartments located within the muscles. Computed tomography usually shows a well-delimited mass with an absorption density between that of muscle and water. The most precise examination technique that enables diagnosis is magnetic resonance imaging (6,7).

Similarly to previous studies (1,6,9), the myxoma in the present study had a sharply defined border. In addition, it exhibited a signal intensity lower than that of the skeletal muscle.
muscles on T1-weighted images and brighter than adipose tissue on T2-weighted images.

In the present case, the large tumor of the thigh was interlinked with the pelvic cavity. It is important to observe the association between the lesion and pelvic organs, such as the appendix or ovaries, as appendiceal mucinous neoplasm may invade organizations outside of the mucous layer of the appendix and cause secondary peritoneal myxoma (10). Inside the mucus, epithelial cells rich in secretory functions can be found (11). In the present case, no epithelial cells were found through pathological examination. The reason for this may be associated with an inadequate local palliative surgery.

The patient rejected further treatment and no follow-up examination was planned. Since the tumor extended to the pelvic cavity and was found to be connected with a cystic and solid neoplasm that was adjacent to the ascending colon in the right lower quadrant, this case of myxoma probably originated from an appendiceal mucinous neoplasm.

In conclusion, the present study showed that the growth of myxoma is slow, with a long disease course. Furthermore, appendiceal mucinous neoplasms may invade outside of the mucosal layer of the appendix, leading to secondary myxoma, which may extend to the leg, as observed in the present study.

References