

Desmoplastic Fibroblastoma of the Foot: A Case Report¹발에서 발생한 결합조직형성 섬유모세포종: 증례 보고¹Ji-Ye Lee, MD¹, Noh-Hyuck Park, MD¹, Chan-Sub Park, MD¹, Su Ok Seong, MD¹, Tae Jung Kwon, MD²Departments of ¹Radiology, ²Pathology, Myongji Hospital, Goyang, Korea

The desmoplastic fibroblastoma is a unique fibrous soft tissue tumor which shows clinically and morphologically distinct characteristics. Most of the desmoplastic fibroblastomas arise from the extremities, shoulder girdle, posterior neck, upper back or abdominal wall. This tumor resembles histological a desmoid tumor. However, it is much better microscopic circumscribed. The clinical outcome after a surgical resection is better than that of a desmoid tumor. We will report about imaging findings of a desmoplastic fibroblastoma which raised in the foot with a review of the relevant literature.

Index terms

Desmoplastic Fibroblastoma

Collagenous Fibroma

Foot

Magnetic Resonance Imaging

Soft Tissue Tumor

Received October 22, 2013; Accepted December 13, 2013

Corresponding author: Noh-Hyuck Park, MD

Department of Radiology, Myongji Hospital, 55 Hwasu-ro 14beon-gil, Deogyang-gu, Goyang 412-270, Korea.

Tel. 82-31-810-7164 Fax. 82-31-810-6537

E-mail: nhpark904@gmail.com

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/3.0>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

INTRODUCTION

In 1995, Evans (1) first described a desmoplastic fibroblastoma (DF), a fibrous soft tissue tumor comprising spindle shaped cells to stellate fibroblastic cells sparsely distributed in a dense fibrous background. This tumor, an alternative also called collagenous fibroma (2), was clinically and morphologically distinct, as well as completely benign in previously reported series. Details of magnetic resonance imaging (MRI) findings for such a tumor have been described for only a few cases (3, 4).

In this article, we present the case of a DF of the foot and we will discuss about the MRI findings of this tumor.

CASE REPORT

A 44-year-old man presented a palpable mass in the plantar aspect of the left foot. He was otherwise feeling well with no reports in the medical history or a previous foot trauma. The laboratory samples results were normal.

The foot radiography showed an about 3.2 × 2.0 cm sized slightly dense soft tissue mass at the plantar aspect of his left foot (Fig. 1A).

In a MRI, a 2 × 2.8 × 2.1 cm sized ovoid shaped mass lesion in the medial plantar aspect of the left foot was shown. On T1-weighted images [repetition time (TR) = 436 ms, echo time (TE) = 12 ms] the mass was well circumscribed, homogenous and had slightly low signal intensity (SI) compared with adjacent muscles (Fig. 1B). The mass was located beneath the abductor hallucis muscles and medial to the flexor digitorum brevis muscle. The lesion had a mass effect upon the adjacent muscles. On the T2-weighted images (TR = 3179 ms, TE = 60 ms), the mass had mixed SI as high SI within low SI (Fig. 1C). The post contrast T1-weighted images (TR = 548 ms, TE = 12 ms) did not show any enhancement (Fig. 1D).

An excisional biopsy was performed which revealed a firm encapsulated mass (Fig. 1E, F). This mass was totally removed. As a pathological result the tumor had well circumscribed margins and was composed of a proliferation of spindle and stellate

fibroblast-like cells sparsely distributed in a collagenous matrix. The final diagnosis was desmoplastic fibroblastoma (collagenous fibroma) (Fig. 1G, H).

DISCUSSION

The desmoplastic fibroblastoma (collagenous fibroma) are more present in adult male patients with a mean age 50 years at diagnosis. Most of the DFs arise from the extremities, shoulder girdle, posterior neck, upper back or the abdominal wall. The tumors usually are variable in size (1.0–20.0 cm). Clinically, the DF displays clinical typically as a painless, mobile, firm and well-cir-

cumscribed subcutaneous mass which grows slowly with an enlargement duration of more than 6 months. In general, the tumors appear oval, disc shaped or fusiform and have a firm homogenous pearl-gray colored consistency on cut-section (5).

Microscopic most of the DFs show low a cellularity and consist of spindle and stellate shaped fibroblastic cells without atypical or hyperchromatic nuclei (3). Also, in case of doubts, immunohistochemistry aids the correct diagnosis. Especially, the S-100 protein, CD34 and EMA are useful markers for an exclusion of neurofibroma, solitary fibrous tumor and perineuroma (5).

There are only a few previous case reports of DF with MRI features present. Beggs et al. (6) described a single case of syno-

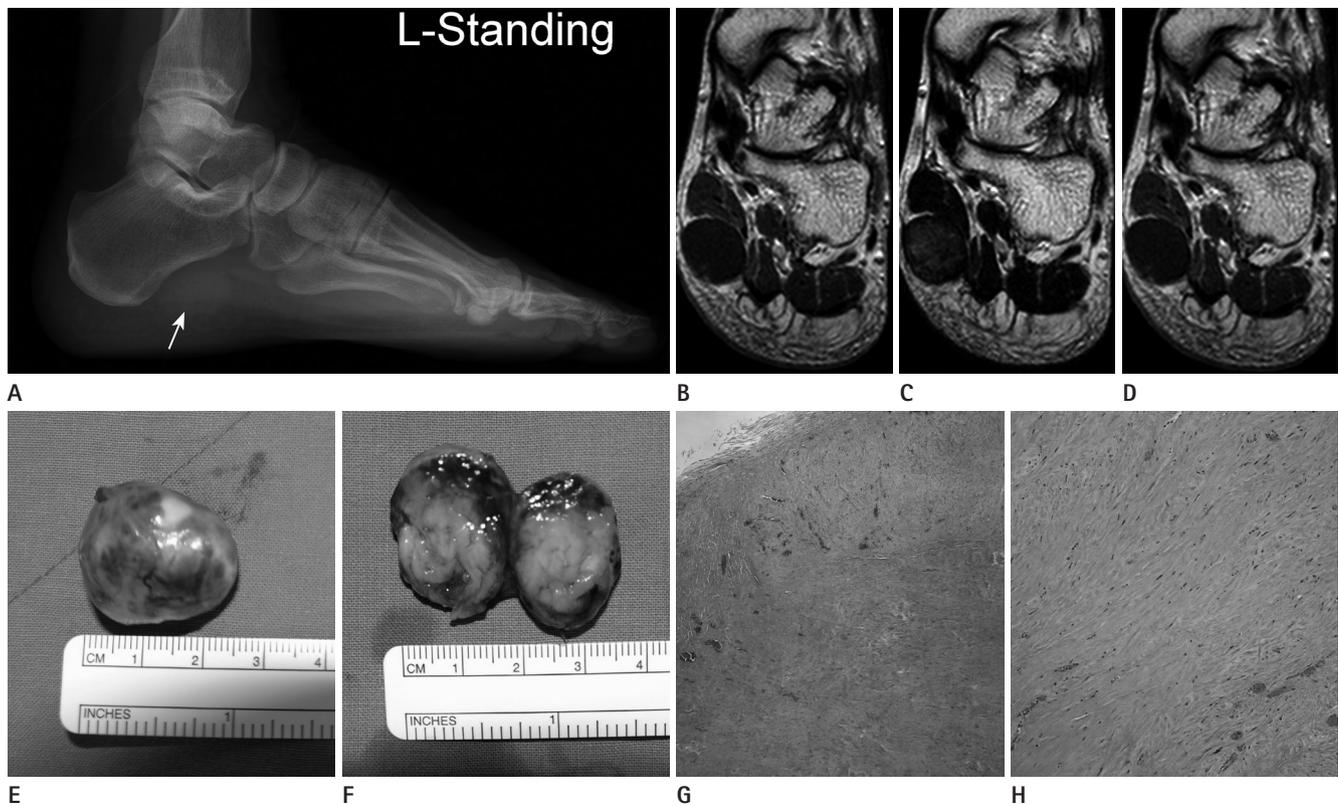


Fig. 1. A 44-year-old man presenting with palpable mass in his left foot.

- A.** The plain lateral radiographs of the foot shows a well margined soft tissue density lesion (arrow). There is no evidence of internal calcification.
- B.** The axial T1-weighted image shows an hypointense lesion compared to the adjacent normal muscle. It is well circumscribed, and was located beneath abductor hallucis muscles and medial to flexor digitorum brevis muscle. The lesion had mass effect upon adjacent muscles.
- C.** The axial T2-weighted image shows a hypointense lesion with internal high signal intensity foci.
- D.** The axial T1-weighted image after gadolinium contrast administration shows minimal or no enhancement.
- E.** Surgical specimen shows a well margined encapsulated grey-purplish firm mass measured as about 2.5 cm in its longest diameter.
- F.** The cut surface appears hemorrhagic at the periphery and glistening gray-to-white aspect and bulging at the center.
- G.** Low-power microscopic specimen shows fibrous and myxocollagenous areas. The tumor border is well defined. Fibrous area in the center of the tumor demonstrates wavy and dense collagen fibers. Cellularity is very low, consisting of spindle-shaped and stellate tumor cells. The periphery appears as a capsule-like covering of loose fibrous tissue (H&E stain, × 100).
- H.** High power microscopic specimen of the fibrous center of the tumor shows monotonous spindle shaped fibroblasts with slender nuclei interspersed in the hyaline stroma of thick, wavy bands of collagen fiber with low cellularity (H&E stain, × 400).

vial DF of the hip joint. Magnetic resonance imaging showed a mass with a minimally high signal intensity on T2-weighted images. Gadolinium enhanced images were not performed.

In our case, the tumor had a smooth well defined margin, inhomogeneous low signal intensity in the T2-weighted images. The area showed slightly high SI on T2-weighted images which corresponded to the hypercellular area within the lesion that consisted of a tumor with loose collagen fibers. The low T2 signal intensity is attributed to the thick abundant collagen background (5).

The most of the soft tissue masses have a high signal intensity on T2-weighted images. Soft-tissue masses with a low SI on T2-weighted images include neurofibroma, cicatricial fibroma, malignant fibrous histiocytoma, aggressive fibromatosis and calcified masses (myositis ossificans, extraskeletal osteosarcoma or chondrosarcoma and synovial sarcoma) (7).

Abundant collagen and marked hypocellularity in a soft-tissue tumor in the absence of calcification result in a decreased signal on the T2-weighted pulse sequence.

Other malignant neoplasms, such as malignant fibrous histiocytoma and fibrosarcoma may also have a low SI within the lesions. Because these malignant neoplasms infiltrate to the surrounding tissue and have usually a necrotic/cystic change, the majority of these tumors can be differentiated from DF due to a MRI (8).

The desmoid tumor is the most important differential diagnosis which should be considered (8, 9). The desmoid tumor is histological more cellular, vascular, and infiltrative at its periphery than the DF. On imaging studies, a desmoid tumor is not as clear circumscribed as a DF. The desmoid tumor may also have larger areas which show a very high SI on T2-weighted images and represents more cellularity than the DF.

The treatment of DF is surgical resection. In our case neither local recurrence nor metastasis occurred after resection which also differed from the high local recurrence rate of a desmoid tumor.

In summary, we described a patient with a DF arising from the foot. This disease should be considered when low SI-areas with poor contrast enhancement in a mass with well-defined margins are seen on both T1- and T2-weighted images.

REFERENCES

1. Evans HL. Desmoplastic fibroblastoma. A report of seven cases. *Am J Surg Pathol* 1995;19:1077-1081
2. Nielsen GP, O'Connell JX, Dickersin GR, Rosenberg AE. Collagenous fibroma (desmoplastic fibroblastoma): a report of seven cases. *Mod Pathol* 1996;9:781-785
3. Ogose A, Hotta T, Emura I, Higuchi T, Kusano N, Saito H. Collagenous fibroma of the arm: a report of two cases. *Skeletal Radiol* 2000;29:417-420
4. Shuto R, Kiyosue H, Hori Y, Miyake H, Kawano K, Mori H. CT and MR imaging of desmoplastic fibroblastoma. *Eur Radiol* 2002;12:2474-2476
5. Miettinen M, Fetsch JF. Collagenous fibroma (desmoplastic fibroblastoma): a clinicopathologic analysis of 63 cases of a distinctive soft tissue lesion with stellate-shaped fibroblasts. *Hum Pathol* 1998;29:676-682
6. Beggs I, Salter DS, Dorfman HD. Synovial desmoplastic fibroblastoma of hip joint with bone erosion. *Skeletal Radiol* 1999;28:402-406
7. Wu JS, Hochman MG. Soft-tissue tumors and tumorlike lesions: a systematic imaging approach. *Radiology* 2009; 253:297-316
8. Hartman TE, Berquist TH, Fetsch JF. MR imaging of extraabdominal desmoids: differentiation from other neoplasms. *AJR Am J Roentgenol* 1992;158:581-585
9. Quinn SF, Erickson SJ, Dee PM, Walling A, Hackbarth DA, Knudson GJ, et al. MR imaging in fibromatosis: results in 26 patients with pathologic correlation. *AJR Am J Roentgenol* 1991;156:539-542

발에서 발생한 결합조직형성 섬유모세포종: 증례 보고¹

이지예¹ · 박노혁¹ · 박찬섭¹ · 성수옥¹ · 권태정²

결합조직형성 섬유모세포종은 드문 섬유성 연조직 종양으로, 임상적, 형태학적으로 특징적인 소견을 보인다. 대부분의 결합조직형성 섬유모세포종은 사지, 견관절, 경부나 복벽에서 많이 발생한다. 이 종양은 조직학적으로 데스모이드 종양과 유사한 특성을 보이나, 주변 조직과 더 경계가 잘 지어진 것이 특징이다. 수술적 절제 후 임상 결과 또한 데스모이드 종양에 비하여 좋다. 저자들은 발에서 발생한 결합조직형성 섬유모세포종의 영상소견 보고 및 관련 문헌의 고찰을 하고자 한다.

명지병원 ¹영상의학과, ²병리과