

THE CLINICAL PICTURE

Morteza Khodaei, MD, MPH
Associate Professor, Department of Family
Medicine, University of Colorado School
of Medicine, Denver, CO

Eric Kim, MD, PhD
House Officer, Department of Family
Medicine, University of Colorado School
of Medicine, Denver, CO

Firm lesion on the lateral thigh



Figure 1. A hyperpigmented nodule on the lateral thigh. **Figure 2.** Retraction with lateral compression.

A 41-year-old man presented with concern about a lesion he had first noticed 6 years earlier

A 41-YEAR-OLD MAN presented with concern about a lesion on the right lateral thigh that he had first noticed 6 years earlier. The lesion had not changed since that time.

There was no history of trauma or infection to the area. He reported no weight loss, and he said he had not noticed similar lesions elsewhere.

Physical examination revealed a firm, nontender, hyperpigmented nodule (7 mm × 9 mm) on the lateral aspect of his right thigh (**Figure 1**). There was no surrounding erythema or warmth. Lateral pressure on the sides of the lesion produced a depression (“dimple” sign) (**Figure 2**).

Based on the clinical picture and physical examination, the diagnosis of dermatofibroma was made.

THE DIFFERENTIAL DIAGNOSIS

Firm, hyperpigmented, macular or nodular skin lesions are prevalent and seen in a number of conditions (**Table 1**).

Dermatofibroma (fibrous histiocytoma) is a benign proliferation of collagen fiber and

doi:10.3949/ccjm.87a.19107

other mesenchymal cell lines, likely in response to local inflammation or trauma.¹

Dermatofibromas are more common in women and typically develop between ages 20 and 50.^{1,2} They often present as smooth, slow-growing, firm, tan to reddish brown papules or nodules less than 1 cm in diameter that classically dent on compression.¹⁻³ They are mostly asymptomatic and may appear anywhere on the body, though 20% are on extremities.¹⁻³ Lesions are typically darker in color in the center and lighter toward the perimeter.¹⁻³

Lentigo maligna is a premalignant melanocytic nevus that may be considered melanoma in situ in its most advanced stages.⁴ It has a high risk of progression to invasive melanoma.⁴ Lesions typically present on sun-exposed skin such as the head or neck.⁴ They appear as heterogeneous asymmetric macules with irregular borders that grow centrifugally.⁴ Ultraviolet light examination with a Wood lamp can show extension of the lesion far beyond the pigmented borders.⁴

Treatment is typically by surgical excision with borders greater than 7 mm and

TABLE 1

Differential diagnosis of a firm, hyperpigmented, macular skin lesion

| Condition | Characteristics |
|---------------------------------|--|
| Lentigo maligna | Irregular asymmetric pigmented macules that grow centrifugally |
| Dermatofibrosarcoma protuberans | Similar to dermatofibroma, but larger, irregular border, deeper skin invasion |
| Seborrheic keratosis | Shiny (“oily”), well-demarcated macule or papule, “stuck-on” appearance |
| Epidermoid inclusion cyst | Flat or raised flesh-colored cystic lesion; often has dark central punctum; size varies; may spontaneously drain |
| Dermatofibroma | Slow-growing, firm, tan to reddish-brown papules, < 1 cm, that “dimple” to lateral compression |

histopathologic examination of the margins, though radiation and topical imiquimod may be used in specific circumstances.⁴

Dermatofibrosarcoma protuberans is a malignant neoplastic lesion, more common in women and darker-skinned individuals age 30 to 50.⁵ It can present as an asymptomatic, slow-growing, violaceous nodule or plaque, more often on the trunk or upper extremities.⁵ It is typically larger than a dermatofibroma, with an irregular border and deeper palpable skin invasion.³ Diagnosis is typically by excisional biopsy.⁵ Though it has a low metastatic potential, it can have a great capacity for local invasion and destruction.⁵

Treatment requires excision with exhaustive histopathologic examination of boundaries for tumor cells, either by Mohs micrographic surgery or wide local excision.⁵ Adjuvant and neoadjuvant therapies such as radiation and imatinib may also be used in cases refractory to excision or with extensive invasion.⁵

Seborrheic keratosis is a common benign skin tumor that becomes increasingly common with age, though lesions can present at any age.⁶ They can be pigmented and so may be mistaken for dermatofibroma, but they do not dimple to lateral compression.⁶ They

typically present as sharply demarcated ovoid macules or papules 1 cm in diameter and with a shiny (“oily”) appearance.⁶ They classically appear raised and “stuck on” to the skin.^{1,6} Obvious seborrheic keratoses may be monitored, but questionable lesions should be diagnosed with shave excision or curettage and histopathology.⁶

Epidermoid inclusion (sebaceous) cyst is a common cystic lesion that can be flat or raised, with size ranging from a few millimeters to a few centimeters.¹ They often have a dark central punctum, which occasionally drains.¹ They are benign and should be removed only if they cause symptoms such as frequent infection or for cosmetic reasons.¹

■ MANAGEMENT OF DERMATOFIBROMA

Most dermatofibromas do not require treatment unless they show signs of malignant progression such as a change in quality or rapid growth.^{1,2} It is essential to distinguish them from the far more malignant dermatofibrosarcoma protuberans, as well as melanoma and other malignant lesions. Irregular borders or substantial palpable depth of invasion through skin should prompt excisional biopsy for definitive diagnosis.³

Most dermatofibromas do not require treatment unless they show signs of malignant progression

■ REFERENCES

- Higgins JC, Maher MH, Douglas MS. Diagnosing common benign skin tumors. *Am Fam Physician* 2015; 92(7):601–607. PMID:26447443
- Myers DJ, Fillman EP. *Dermatofibroma*. Tampa, FL: StatPearls Publishing; 2019.
- Zelger B, Zelger BG, Burgdorf WH. Dermatofibroma—a critical evaluation. *Int J Surg Pathol* 2004; 12(4):333–344. doi:10.1177/106689690401200406
- Kallini JR, Jain SK, Khachemoune A. Lentigo maligna: review of salient characteristics and management. *Am J Clin Dermatol* 2013; 14(6):473–480. doi:10.1007/s40257-013-0044-6
- Acosta AE, Velez CS. Dermatofibrosarcoma protuberans. *Curr Treat Options Oncol* 2017; 18(9):56. doi:10.1007/s11864-017-0498-5
- Hafner C, Vogt T. Seborrheic keratosis. *J Dtsch Dermatol Ges* 2008; 6(8):664–677. doi:10.1111/j.1610-0387.2008.06788.x

Address: Morteza Khodae, MD, MPH, Department of Family Medicine, University of Colorado School of Medicine, AFW Clinic, 3055 Roslyn Street, Denver, CO 80238; morteza.khodae@cuanschutz.edu