Painless Mobile Nodule on the Shoulder

Preeya T. Shah, MD; Patrick M. Kupiec, MD; Allen Strickler, MD, PhD; Eric W. Hossler, MD

A 70-year-old woman presented to the outpatient dermatology clinic with an acute-onset lesion on the right shoulder. She first noticed a “cyst” developing in the area approximately 3 weeks prior but noted that it may have been present longer. The lesion was bothersome when her undergarments rubbed against it, but she otherwise denied pain, increase in size, or drainage from the site. Her medical history was remarkable for a proliferating trichilemmal tumor on the right parietal scalp treated with Mohs surgery approximately 13 years prior to presentation. She had no personal or family history of skin cancer. Physical examination revealed a 2.5-cm, mobile, nontender, flesh-colored subcutaneous nodule on the right shoulder (top); no ulceration, bleeding, or drainage was present. The surrounding skin demonstrated no clinical changes. The patient was scheduled for outpatient surgical excision of the nodule, which initially was suspected to be a lipoma. During the excision, a translucent cystlike nodule (bottom) was gently dissected and sent for histopathologic examination.

WHAT’S YOUR DIAGNOSIS?

a. cutaneous ciliated cyst
b. cutaneous metaplastic synovial cyst
c. epidermoid cyst
d. ganglion cyst
e. nodular hidradenoma

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THE **DIAGNOSIS:**

**Cutaneous Metaplastic Synovial Cyst**

Gross examination of the excised nodule revealed a 2.5×1.2×1.0-cm, intact, gray-white, thin-walled, smooth-lined nodule filled with clear mucinous-like material. Hematoxylin and eosin–stained sections demonstrated a dermal-based cystlike structure composed of a lining of connective tissue with hyalinized material and fibrin as well as spindle and epithelioid cells with a mild mixed inflammatory infiltrate (Figure). These histopathologic findings led to the diagnosis of cutaneous metaplastic synovial cyst (CMSC).

Cutaneous metaplastic synovial cyst, also known as synovial metaplasia of the skin, is an uncommon benign cystic lesion that was first reported by Gonzalez et al. in 1987. Histologically, CMSC lacks an epithelial lining and therefore is not a true cyst but rather a pseudocyst. Clinically, the lesion typically presents as a solitary subcutaneous nodule that may be tender or painless. In a literature review of CMSC cases performed by Fukuyama et al., distribution of reported cases according to body site varied; however, limbs were found to be the most commonly involved area. A PubMed search of articles indexed for MEDLINE as well as a Google Scholar search using the term *cutaneous metaplastic synovial cyst* revealed at least 37 cases reported in the English-language literature, including our present case. The pathogenesis remains uncertain; however, a majority of previously reported cases of CMSC characteristically have been associated with a pre-existing lesion, with most presentations developing at surgical scar sites secondary to operation or trauma.

Relative tissue fragility secondary to rheumatoid arthritis and Ehlers-Danlos syndrome has been linked to CMSC in some documented reports, while a minority of cases report no antecedent events triggering formation of the lesion.

As evidenced by our patient, CMSC clinically mimics several other benign entities; histopathologic examination is necessary to confirm the diagnosis. Although nodular hidradenoma also may clinically present as a solitary firm intradermal nodule, microscopy reveals a dermal-based lobulated tumor containing cystic spaces and solid areas composed of basophilic polygonal cells and round glycogen-filled clear cells. Epidermoid cysts are differentiated from CMSC by the presence of a cyst wall lining composed of stratified squamous epithelium and associated laminated keratin within the lumen, which corresponds to its pearly white appearance on gross examination. Cutaneous ciliated cysts predominantly occur on the lower extremities of young women and are lined by simple cuboidal or columnar ciliated cells that resemble müllerian epithelium. Similar to CMSC, ganglion cysts are pseudocysts that lack a true epithelial lining but differ in appearance due to their mucin-filled synovial-lined sac. Additionally, ganglion cysts most often occur on the dorsal and volar aspects of the wrist.

Excisional biopsy is indicated as the preferred treatment of CMSC, given the lesion’s benign behavior and low recurrence rate. Our case highlights this rare entity and reinforces its inclusion in the differential diagnosis of subcutaneous mobile nodules, especially in the setting of prior tissue injury secondary to trauma, surgical procedures, or conditions such as rheumatoid arthritis or Ehlers-Danlos syndrome. Unlike most previously reported cases, our patient reported no preceding tissue injury associated with formation of the lesion, and she was largely asymptomatic on presentation. Considering the limited number of CMSC cases demonstrated in the literature, it is important to continue reporting new cases to better understand characteristics and presentations of this uncommon lesion.

**REFERENCES**


