Epidermal Inclusion Cyst after Carpal Tunnel Release: A Case Report

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Abstract

Background Carpal tunnel release is one of the most commonly performed procedures in upper extremity surgery, with low complication rates.

Case Description We report a case of epidermal inclusion cyst that occurred in a 44-year-old man after carpal tunnel release. The epidermal inclusion cyst formed beneath the previous operation scar and induced recurrent carpal tunnel syndrome. Symptoms resolved completely after excision of the cyst.

Literature Review Epidermal inclusion cysts are the third most common type of hand tumors. Although repeated minor trauma is the most frequent cause, surgical procedures can induce an epidermal inclusion cyst. One case report of epidermal inclusion cyst after carpal tunnel release was found in literature. Etiology, treatment, and prevention methods are reviewed.

Clinical Relevance We hope this case report alerts physicians to the possibility of development of epidermal inclusion cyst after carpal tunnel release and the importance of basic surgical skills, such as gentle handling of a surgical wound, profuse saline irrigation, and careful wound closure with precise approximation of skin edges.

Case Report

A 44-year-old, right-handed man presented with a 1-year history of bilateral (left hand greater than right hand) numbness and pain in the median nerve distribution. Symptoms had aggravated in the 3 months prior to presentation. Electromyography and nerve conduction studies indicated inclusion cyst caused by carpal tunnel release has been described only once before.

Here we describe a case of epidermal inclusion cyst that occurred after carpal tunnel release. The aim of this report is to introduce this rare complication of carpal tunnel release and emphasize the importance of basic surgical skills, such as gentle handling of a surgical wound, profuse saline irrigation, and careful wound closure with precise approximation of skin edges.

Keywords ► carpal tunnel syndrome ► carpal tunnel release ► epidermal inclusion cyst
bilateral median neuropathies at the wrist with more severe involvement of the left side. We performed mini-open carpal tunnel release on the left side as usual, and the surgical wound was closed with vertical mattress sutures of 3–0 nylon. The postoperative course was uneventful, and the patient stopped follow-up visits with complete resolution of his symptoms 4 months after the surgery. However, 20 months after surgery, he revisited with complaints of a palpable mass at the previous operation site and a tingling sensation in the medial nerve distribution. The mass grew insidiously, and he denied any traumatic episode involving the left hand or wrist after the previous operation. Physical examination revealed a mass beneath the previous operation scar (►Fig. 1). The mass was palpable and firm without signs of infection and had a positive Tinel sign. Ultrasonography demonstrated a 1.0 × 0.8-cm round, well-demarcated, irregular echogenic soft tissue mass under the dermis (►Fig. 2).

We decided to perform surgical exploration and made a 2-cm skin incision along the previous operation scar. A shiny, white, well-demarcated round mass was found in the subcutaneous tissue layer (►Fig. 3). The mass was easily pulled out. The wound was copiously irrigated and then closed with vertical mattress sutures of 3–0 nylon. Histological examination

**Fig. 1** Preoperative photographs showing an oval, firm, protruding mass on the scar of the previous carpal tunnel release.

**Fig. 2** Ultrasonography showing a round, well-demarcated, irregular echogenic mass under the dermis.

**Fig. 3** Intraoperative photographs showing a well-demarcated, round whitish yellow mass.

**Fig. 4** Photomicrographs of biopsy specimen demonstrating stratified squamous cell epithelial lining and eosinophil keratin debris in the cystic cavity. (a) Hematoxylin-eosin stain, magnification x10. (b) Magnification × 100.
confirmed the diagnosis of an epidermal inclusion cyst (Fig. 4). The patient had an uneventful postoperative course, and his preoperative symptoms disappeared. Six months following the excision, there was no evidence of recurrence and the patient remained free of symptoms.

Discussion

Epidermal inclusion cysts, also known as epidermoid inclusion cysts, epidermal cysts, epithelial cysts, keratin cysts, implantation cysts, infundibular cysts, or sebaceous cysts, are unicellular cysts without septation that are encapsulated with fibrous tissue and lined by true epithelium, which is stratified, keratinized squamous epithelium with a granular layer as found on the skin surface and in the infundibulum of hair follicles. Because these cysts do not involve sebaceous glands and do not contain sebum, the term “sebaceous cyst” is a misnomer. The cysts contain central, eosinophilic, keratinaceous material comprising accumulations of cutaneous products, the debris of keratin, proteins, cholesterol, and cell membrane lipids. The contents usually look like a chunk of cheese and have a foul smell. Cysts can be skin-colored, yellow, or white and are usually slow-growing and asymptomatic. However, polymicrobial infection of cysts with aerobic or anaerobic organisms may occur, and malignant transformation, although very rare, has been reported. Epidermal inclusion cysts are, strictly speaking, epidermoid cysts of traumatic origin. The proposed mechanisms of development of epidermod cysts include incomplete cleavage of cutaneous ectoderm at the embryonic stage, squamous metaplasia of the columnar epithelial cells within dilated ducts, downward growth of epithelial cells with inflammation after obstruction of the hair follicle, human papillomavirus infection, and growth of implanted fragments of the epidermis within deep tissue after trauma or surgical procedures. Epidermoid cysts of nontraumatic origin are commonly located on hair-bearing areas of the upper chest, upper back, neck or head, while epidermoid cysts of traumatic origin—epidermal inclusion cysts—are more common on nonfollicular areas such as the palms, soles, or buttocks.

Epidermal inclusion cysts are the third most common type of tumor of the hand, after ganglion cysts and giant cell tumors of the tendon sheath. They are more frequently seen in male manual laborers who are subject to repeated minor trauma to the hand. They are usually located in the dermis or subcutaneous tissue layer of the volar surface, especially the distal portion of the digits, as a solitary lesion. However, multiple lesions or deep tissue involvement, such as tendon and bone, have been reported also. Lincoski et al reported the largest series of epidermal inclusion cysts of the hand. They experienced 101 cysts (94 patients) among 623 hand tumors (16%) over a period of 27 years. Of these patients, 83% were male and 56% were manual laborers. Mean age was 46 years old. Forty-eight percent recalled a definite traumatic event prior to cyst development, and three cases had cysts that occurred after prior unrelated surgery. However, detailed information about the nature of previous operations was not provided. Seventy-two cysts (71%) were located on the volar surface. Four cysts (0.04%) were intraosseous, and four cases (0.04%) were multiple. There were 11 cases (11%) of recurrence. Lucas also reported 58 patients (60 cases) with epidermal inclusion cysts of the hand. In that study, 85% were male (49 patients), and 91% (53 patients) were manual laborers. Forty-seven percent (27 patients) recalled a traumatic event. Cysts were intraosseous in two patients (0.03%), and recurrence occurred in 10 patients (17%).

Although repeated minor trauma is a frequent cause of epidermal inclusion cysts in the hand, postoperative occurrence of epidermal inclusion cysts in the hand has been rarely reported. We found only two cases in the English-language medical literature. One was due to dermofasciectomy and full-thickness skin graft to treat Dupuytren’s contracture, while the other was due to mini-open carpal tunnel release. Low et al presented a case of epidermal inclusion cyst after carpal tunnel release and presumed that subcuticular suture with braided polyglactin implanted epidermal tissue into the deep tissue. In our case, we had closed the operation wound with vertical mattress sutures of 3–0 monofilament nylon. It is therefore unclear how epidermal tissue was implanted in our case.

As mentioned previously, all kinds of surgical and invasive procedures can induce epidermal inclusion cysts. To prevent development of epidermal inclusion cysts, surgeons should use a fresh, sharp scalpel and should be careful not to make invaginations of skin edges. Moreover, when sharp instruments or powered instruments are used, surgeons should be careful not to push epithelium into the deeper tissues. Profuse irrigation of the operation wound is also important to wash out epithelial debris. Although carpal tunnel release is a relatively safe procedure and is widely performed, surgeons have to keep in mind these recommendations to prevent the occurrence of epidermal inclusion cysts.

Conflict of Interest
None.

References
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