Iatrogenic Intraspinal Epidermoid Cyst

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Epidermoid cyst is generally regarded as congenital disease, and commonly related to other congenital spinal anomalies. However, it also develops iatrogenically. We report one rare case of epidermoid cyst that we experienced among intradural extramedullary tumors. A 21-year-old female patient was admitted to the hospital due to low back pain with radiating pain into the right lower extremity that initiated about a month ago. She complained sensory loss and motor weakness (grade 4+) on her right extremity as well as urinary dysfunction and sphincter dysfunction. She had a lumbar puncture three times due to Pneumococcal meningitis when she was 13 years old. The well-circumscribed intradural extramedullary mass of 1.8×1.6×4 cm size was found on the L4-5 in a magnetic resonance image. Gross total tumor removal was carried out after the total laminectomy L5 and partial laminectomy L4. The tumor mass was gray-colored and so fragile that it was easy to be removed. The histological diagnosis confirmed epidermoid cyst without malignancy. After the operation, the patient progressively showed remarkable neurological recovery. In this case, the cause of epidermoid cyst is considered iatrogenic concerning history of several times of lumbar puncture as meningitis.

Key Words: Epidermoid cyst • Lumbar puncture • Iatrogenic

INTRODUCTION

Epidermoid cyst, also called “pearly tumor” because of the smooth, glistening, white capsule, is a rare, and slow growing benign tumor, which represents less than 1% of all intraspinal tumors. Most intraspinal epidermoid tumors are intradural extramedullary and their origin may be congenital or acquired. Congenital epidermoid tumors may develop from displaced cutaneous ectoderm during neural tube formation, while the common cause of an acquired epidermoid tumor is implantation of epidermal tissue in the subdural space during surgery or lumbar puncture.

We report one rare case of an iatrogenic epidermoid cyst occurring intradural extramedullary portion.

CASE REPORT

A 21-year-old female patient was admitted to the hospital...
without malignancy (Fig. 4). After the operation, the patient showed remarkable neurological recovery.

**DISCUSSION**

Epidermoid tumors are represented histologically by a mass of desquamated cells containing keratohyalin encased within a laminated capsule of well-differentiated stratified squamous epithelium. Pathogenetically, these tumors may be secondary to congenital development or iatrogenic implantation. Congenital tumors are associated with spinal malformations, such as spina bifida and other dysraphic states, with a greater tendency to develop in thoracic spines.

Many cases of epidermoid cyst development have since been described in patients with a history of lumbar puncture, and Korea is no exception. In 1962, Manno reported 41% of patients with spinal epidermoid cyst had a history of lumbar puncture. Gibson and Norris mentioned that skin fragments had been left in needles in 70% of cases when skin puncture had been done without a stylet. According to the Van Gilder and Schwartz study in rats, intraspinal epidermoid tumors after skin implantation had occurred in 89% of young rats. Through this study, it became widely known that epidermoid cysts can be acquired when using needles without a stylet or with an ill-fitting stylet for lumbar puncture because parts of epithelial layers could be infiltrated to spinal cavities.

The most frequent site of congenital spinal epidermoid tumor is the intramedullary thoracic segment, while the tumors acquired during lumbar puncture are generally located poste-
riorly in the lumbar inter space and intradural extramedullary space of the conus medullaris the cauda equina segment. MRI can show a typical homogeneous or heterogeneous hypointense signal on T1-weighted image and hyperintense signal on T2-weighted image. However, epidermoid tumors showing atypical signal intensity changes have been reported, such as hyperintense signal on T1-weighted image and hypointense signal on T2-weighted image, as seen in our case. This variability in signal characteristics, which might be related to the chemical state of cholesterol or the relative composition of cholesterol and keratin, makes preoperative diagnosis difficult. Additionally, no contrast enhancement is seen in most cases, but enhancement of a thin peripheral rim has been reported.

Our patient had no other congenital malformations and had a history of 3 times lumbar punctures for pneumococcal meningitis. Although she had no imaging before, such as myelography or MRI, a pre-existing tumor could be excluded. Lumbar puncture levels were also not reported, so their locations could not be verified. However, it can be suggested that this spinal epidermoid cyst had developed as a complication after lumbar puncture because no other anomalies had been found in the patient's 21 years, symptoms had occurred in the 8 years after lumbar puncture, and the cyst's location at L4-5 level corresponded with a frequent puncture site. Time between puncture and onset of symptoms was 2-23 years in one study.

There is a possibility that use of a hollow needle without stylet, an ill-fitting stylet, and young age may be associated with development of the epidermoid cyst.

**CONCLUSION**

The present case emphasizes the fact that epidermoid tumors can occur after lumbar puncture. Patient without congenital defects of the spinal column but with history of lumbar puncture should be aware of potential adverse complications, such as epidermoid tumors, although their incidence may be low. In cases of intradural tumors or tumor-like lesions showing high signal on T1-weighted image without central enhancement, spinal epidermoid cysts should be considered.

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