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Kyoto University
SOLITARY RETROPERITONEAL NEUROFIBROMA: A CASE REPORT

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A solitary neurofibroma arising in the retroperitoneal space without any other stigma of von Recklinghausen's disease is reported. Confusion with another nerve sheath tumor, a schwannoma is a diagnostic pitfall. Histochemical and immunohistochemical stainings of the tumor are useful for the diagnosis of solitary neurofibroma.

Key words: Solitary neurofibroma, Retroperitoneal space, Alcian blue staining, S-100, NSE

INTRODUCTION

A neurofibroma is a benign nerve sheath tumor that occurs as a solitary tumor or as a partial manifestation of von Recklinghausen's disease: an inherited disease characterized by multiple cutaneous pigmented lesions (cafe-au-lait spots) and subcutaneous neurofibromas. Although genitourinary involvement of neurofibromas is rare, such cases have been reported in the bladder, kidney, ureter, penis, and retroperitoneal space\(^2\)\(^-\)\(^6\). Previous literature shows confusion relating to the terminology of the peripheral nerve tumors\(^7\). We report a case of a solitary retroperitoneal neurofibroma diagnosed by histochemical and immunohistochemical stainings as well as light microscopic appearance of the tumor.

CASE REPORT

A 56-year-old Japanese woman presented with a 2-year history of dysuria which fluctuated with periods of exacerbations and remissions. Significant medical history included total abdominal hysterectomy for leiomyomas 22 years earlier. Family history was negative for von Recklinghausen's disease. Physical examination reveal-
Fig. 1. DIP shows the displacement of the bladder and left ureter medially, suggesting a large extravesical mass.

Fig. 2. CT demonstrates a retrovesical mass.

Fig. 3. Histologic appearance of the tumor shows fusiform tumor cells in a matrix (H & E stain).

(Fig. 4, A and B). Thus, the tumor was diagnosed as a solitary neurofibroma. The patient remains well without any signs of tumor recurrence.

**DISCUSSION**

A neurofibroma and a schwannoma (neurilemoma) are benign nerve sheath tumors, originating from the Schwann cells. Although there is confusion relating to differentiation between a neurofibroma and a schwannoma, a neurofibroma is distinct from a schwannoma in the following points. (i) The former lacks the compact arrangement of cells in the Antoni type A tissue and delicate single cell thick fascicles of the Antoni type B tissue of a schwannoma\(^7\) (ii) Alcian blue stains for acid mucopolysaccharides are negative in a schwannoma but positive in a neurofibroma\(^7\). (iii) A neurofibroma contains fewer S-100 protein positive cells than a schwannoma\(^8\).

S-100, a dimetric calcium binding protein, composed of different combination of \(\alpha\) and \(\beta\) subunits, mainly locates in glial and Schwann cells\(^9\). NSE, a glycolytic dimetric enzyme, locates in neurons and neuroendocrine cells\(^10\). Immunohistochemical demonstration of S-100 and NSE is valuable in the diagnosis of
a nerve sheath tumor. Unbalanced expression of S-100 subunits is observed in malignant nerve sheath tumors. Hayashi et al. observed an increased expression of the S-100 a subunit in the malignant part of neurofibromas. They concluded that the S-100 subunit immunoreactivity is a good marker for malignant schwannomas.

As described above, we could not distinguish between a neurofibroma and a schwannoma in the previous literature. Das Gupta et al. considered solitary neurofibromas and schwannomas (neurilemomas) as the same entity, and they reviewed 303 such cases. In that report, most tumors occurred in the head and neck, and extremities. On the other hand, only two of these cases (0.66%) were retroperitoneal origin. Surgical excision of the tumor is the choice of treatment.

Careful search for the coexistence of a malignant tumor is necessary, because 16% of the patients with a solitary neurofibroma were reported to have an associated malignant tumor.

It remains to be elucidated whether the clinical course of a solitary neurofibroma differs from that of a schwannoma.

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和文抄録

孤立性後腹膜神経線維腫の1例

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後腹膜に原発し，von Recklinghausen氏病の徴候のない孤立性神経線維腫の1例を報告する。診断上，もう一つの神経線維腫を併発するのを問題にした。腫瘍の組織化学染色および免疫組織化学染色が孤立性神経線維腫の診断に有用であった。

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