A case report of a thymic neuroblastoma associated with syndrome of inappropriate secretion of antidiuretic hormone: Ten-year follow-up results after surgical treatment

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1. Introduction

In 2009, we reported an extremely rare case of thymic neuroblastoma in an adult with syndrome of inappropriate secretion of antidiuretic hormone (SIADH) [1]. Till date, few cases of thymic neuroblastoma with SIADH have been reported in the English literature [2,3]. Although neuroblastomas are known to be malignant, grow rapidly, and have a poor prognosis, the etiology of thymic neuroblastoma is still uncertain. Moreover, limited long-term prognostic data on the disease in adults are available, and standard therapy is yet to be established [1]. We describe here the clinical course of neuroblastoma with SIADH over a period of 10 years; the patient achieved a successful outcome and had no recurrence after surgery. We focus on the management and surgical assessment. This work has been reported in line with the SCARE criteria [4].

2. Presentation of case

In 2008, a 60-year-old Japanese male patient was admitted to our hospital for further examination and treatment of an abnormal shadow seen on his chest x-ray during a regular health check-up. An enhanced chest computed tomography scan revealed a 47-mm solid mass with a clear rim in the anterior mediastinum (Fig. 1). Magnetic resonance imaging using intravenous contrast showed iso-intensity and high intensity of the mass on T1- and T2-weighted images, respectively. Laboratory findings showed a serum sodium concentration of 119 mEq/L, plasma osmolality of 261 mOsm/L, and an elevated plasma antidiuretic hormone level of 6.4/L. A diagnosis of thymoma with SIADH was suspected, and the patient underwent total thymectomy. Based on the microscopic findings with immunohistochemistry, the final diagnosis was thymic neuroblastoma. The histologic pattern confirmed it as a poorly differentiating neuroblastoma based on the classification according to the new World Health Organization system [5]. The tumor had a thin, fibrous capsule and showed no invasion into the surrounding atrophic thymic tissue. His serum sodium levels and blood osmolality quickly returned to normal after surgical excision of the tumor. The patient was provided detailed information about the disease. We recommended adjuvant therapy since the disease is considered malignant with a poor prognosis; however, the patient did not wish to undergo therapy after surgery. Therefore, he was regularly followed up on an outpatient basis with periodic computed tomography scans and laboratory studies including blood analysis and urinalysis once a year. At the 10-year follow-up after surgery, the 70-year-old patient remains free of the disease and is asymptomatic. Written informed consent was obtained from this patient for the publication of this report.

3. Discussion

Neuroblastoma is considered a pediatric malignancy since over 95% of the cases are diagnosed in children aged less than 10 years [6]. These tumors have a very broad spectrum of clinical manifes-
4. Conclusion

In the present case, complete surgical removal is considered to be an option for the treatment of this tumor when it is clearly separated from the surrounding tissues and with no invasion. However, further investigations are necessary to confirm whether it could indeed be a treatment strategy in this disease.

Please state any conflict of interest

The author declares no conflicts of interests.

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Ethical approval

This case report is exempted from ethical approval by our institution.

Consent

The author states that they have written and signed consent from the patient to publish this report.

Author contribution

Yukitoshi Satoh: Conceptualization, Methodology, Validation, Investigation, Writing – original draft, Writing – review and editing, Visualization, Supervision.

Registration of research studies

The author does not need to register this work.

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