Long-Term Survival in Adult Neuroblastoma with Multiple Recurrences

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Abstract
Neuroblastoma (NB) rarely occurs in adults, and less than 10% of the cases occur in patients older than 10 years. Currently, there are no standard treatment guidelines for adult NB patients. We report the case of a young man suffering from NB in adulthood with multiple recurrences. Treatment included multiple resections, chemotherapy, and radiotherapy. This patient remains free of clinical disease more than 7 years after diagnosis.

Introduction
Neuroblastoma (NB) is a tumor derived from the neural crest that usually originates in the adrenal medulla, but it may arise anywhere within the sympathetic nervous system. NB is a common malignancy in children but rarely occurs in adults; less than 10% of all cases are diagnosed after the age of 10 years. The incidence for patients between 30 and 39 years of age is approximately 0.2 cases per million person-years [1, 2]. NB has different biological features in adolescents and adults, and a longer course in both of these groups than in children. However, the outcome is poor regardless of stage [3].

We report the effective use of multimodal therapy to achieve long-term survival in adult retroperitoneal NB with multiple recurrences.

Case Report
In February 2002, a previously healthy 25-year-old man presented with a supraclavicular left node that was completely resected with a diagnosis of neurofibrosarcoma. Physical examination revealed an abdominal mass. Computed tomography (CT) showed a multinodular 13.5 × 10-cm tumor in the
retroperitoneal region, covering the aorta and repulsing the inferior vena cava. No distant metastases were detected. Guided needle biopsy of the lesion revealed adult NB upon histopathological examination at an outside institution. Histologically, the tumor consisted of small and round-shaped pseudorosette-forming cells with hyperchromatic nuclei. Immunohistochemistry showed cytoplasmic positivity for neuronal specific enolase (NSE), chromogranin A, and synaptophysin. A bone scan was negative. Metaiodobenzylguanidine (MIBG) scintigraphy demonstrated an increased uptake by the mass described above. The serum levels of NSE and catecholamine metabolites, including dopamine, were in the normal range. Catecholamine metabolites were not increased in a 24-hour urine collection.

The patient was diagnosed as having stage IV NB. Combination chemotherapy with a CADO/CDDP-VP16 regimen according to the pediatric protocol was started with cyclophosphamide (CPM 1,500 mg/m² per cycle), doxorubicin (AD 60 mg/m² per cycle), vincristine (2 mg per cycle), cisplatin (CDDP 100 mg/m² per cycle), and etoposide (VP16 500 mg/m² per cycle) with G-CSF.

After 5 courses of chemotherapy, the mass was surgically removed and a lomboaortic lymphadenectomy performed. The mass was NB with 70% necrosis. Fluorescence in situ hybridization analysis of the MYCN locus showed no MYCN oncogene amplification. External irradiation of paraaortic lymph nodes was performed postoperatively, delivering 35 Gray (Gy) with 1.8 Gy per day, 5 days per week.

Postoperative MIBG scintigraphy showed no abnormal uptake.

Semestrial follow-up studies, including MIBG scintigraphy, CT, and catecholamine values in the urine and blood were performed.

The first recurrence occurred in 2005, 30 months after the initial surgery. The patient was asymptomatic when he presented for the visit. Physical examination and routine blood levels were normal, but the serum dopamine level was elevated (655 μg/g creatinine for normal range 96–463). MIBG scintigraphy demonstrated an increased uptake in the upper mediastinal region, and CT scans of the chest revealed a 25-mm mediastinal mass. The patient underwent thoracotomy and the mass measuring 40 × 60 mm was removed completely. Histological and immunohistochemical examination confirmed the diagnosis of NB metastasis.

One month later, the postoperative scintigraphy showed abnormal uptake in the left supraclavicular area, corresponding to a metastatic lymph node. Because MIBG scintigraphy confirmed disease progression, we administrated carboplatin/VP16 (carboplatin 800 mg/m² per cycle and VP16 500 mg/m² per cycle with G-CSF) therapy. After 2 courses of chemotherapy, ¹⁸F-DG scintigraphy showed a partial response with a diminishing SUVₘₐₓ from 5.2 to 3.3. A 3rd surgical resection was performed, guided by a perioperative detection probe 24 h after the injection of 5.2 mCi of iodine-123-labeled MIBG.

Postoperative metabolic imaging performed at 1, 3, 6, 12, 15, 21, and 24 months did not reveal any abnormality in connection with the disease. However, the postoperative FDG-TEP in the 15th month showed an increased uptake in the left popliteal area, corresponding to a less than 1-cm nodule localized in the lateral part of the superior extremity of the left leg. Surgical biopsy revealed a hemangioma with schwannoma.

In August 2008, ¹⁸F-DG scintigraphy showed abnormal uptake in the left jugular carotid and nodular pleural uptake on the right, but MIBG scintigraphy was normal. Physical examination was normal and did not find any catecholaminergic manifestations.

In January 2009, FDG-TEP was carried out (33 months after the last surgery) and showed a single lesion, measuring 25 mm at its greatest diameter, localized in the pleural area with close contact at the 6th rib and involved the intercostal muscle on the right. An MRI of the chest confirmed the metabolic imaging data, but MIBG scintigraphy was still normal. A thoracotomy was performed and permitted a complete macroscopic resection of the tumor. The histological findings determined that the lesion was a parietal and visceral pleuritic localization of the NB demonstrated with R1 resection. No complications were encountered during the postoperative period. The patient underwent concomitant postoperative radiochemotherapy with cisplatin regimen. The patient is still alive and has no new metastases more than 8 years after diagnosis.
Discussion

Surgical resection, histology, CT, and MIBG scintigraphy are the methods of choice for diagnosis.

In general, catecholamine metabolites in urine or serum, NSE, and lactate dehydrogenase are useful parameters for diagnosis and follow-up. However, an abnormal tumor marker leads to further assessment with detection of relapse or progression in only 14% of patients [4]. For our patient, the serum dopamine level was elevated only in the first recurrence.

Currently, a reported specificity of 99% and cumulative sensitivity of roughly 90% in paraganglioma make $^{123}$I-MIBG the most important nuclear imaging method [5]. However, false negatives have been reported and may be explained by the absence of catecholamine reuptake and/or dedifferentiated features of malignant tumors [6, 7]. In our case, MIBG scintigraphy was negative in the last recurrence.

For paraganglioma, FDG-PET diagnoses malignancy better than MIBG scintigraphy and reveals more metastases. However, FDG-PET has lower sensitivity and, in particular, lower specificity than MIBG scintigraphy for detecting paragangliomas [8, 9]. Moreover, there are some false positive cases with FDG-TEP, as we reported for our patient.

The perioperative detection probe is a supplementary diagnostic tool for guiding the exeresis of recurrent tumors, which are sometimes difficult to access surgically. Several publications have reported the use of a perioperative detection probe after the injection of MIBG-labeled with iodine-125, or iodine-123, or $^{111}$In-pentetrotide [10, 11].

Stage III and IV tumors are often initially treated with chemotherapy followed by tumor resection, as was done in our patient [12]. Because of a lack of experience in adults, the treatment followed pediatric protocols. The most frequently used cytostatic agents, either alone or in combination, include alkylating agents (cyclophosphamide/ifosfamide), platinum-based agents (cisplatin/carboplatin), etoposide, adriamycin, and vinca-alkaloids.

Clinical data on the survival outcomes of adult patients (defined as 20 years of age and older) with NB are scarce due to the rarity of the disease. Small, single institution reports have described worse outcomes for adults than pediatric patients [13, 14]. Data from the publicly accessible Surveillance Epidemiology and End Results confirm these findings, but there has been a trend of improved survival for the past decade, which may reflect advancements in treatment [15]. NB in adolescents and adults exhibits different biological characteristics and has a longer course than in children, with multiple recurrences and/or chronic persistent disease [3]. Moreover, MYCN amplification was rarely found.

Conclusion

Currently, there are no standard treatment guidelines for patients with adult NB. In patients with metastatic disease and at late stages of disease, multimodal therapy should include surgical resection, radiotherapy, and outpatient chemotherapy. Follow-up of patients with NB should include metabolic imaging studies and tumor marker determination. The collection and evaluation of data on adult patients with this rare tumor are warranted in order to optimize treatment strategies. Whether more intensive treatment or myeloablative therapy with stem cell support will improve the outcome of NB in adults is unclear.
References


