Brentuximab vedotin in the treatment of a patient with refractory Hodgkin disease and Proteus syndrome – a case report and discussion

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Key Clinical Message
Treatment of patients with refractory Hodgkin lymphoma is a significant issue. We report a patient with Proteus syndrome and relapsed Hodgkin lymphoma, whose remission was finally achieved after brentuximab vedotin therapy, allowing her to receive a haploidentical stem cell transplant. The possible relationship between both disorders was discussed.

Keywords
Brentuximab vedotin, children, Hodgkin lymphoma, Proteus syndrome.

Introduction
Proteus syndrome (PS) is a rare disease caused by a somatic activating mutation in AKT1, proving the hypothesis of somatic mosaicism and implicating activation of the PI3K-AKT pathway in the characteristic clinical findings of overgrowth and tumor susceptibility in this disorder [1–3]. First described by Cohen and Heyden in 1979, Wiedemann et al. in 1983 named this distinct entity PS. The incidence of PS is approximately 1:100,000–1,000,000 [1, 2]. We report a case of PS combined with refractory Hodgkin lymphoma (HL) and discuss therapeutic options for PS patients with oncological diseases.

Case Report
A 14-year-old female was admitted to the oncohematology department with enlargement of the peripheral and mediastinal lymph nodes. Asymmetry of her scalp bones and gum hyperplasia were noted at birth. The diagnosis of PS was made phenotypically on the basis of existing scalp and oral cavity deformation, antimongoloid eyes, chest kyphosis, and intact intellect. No genetic changes were found by conventional cytogenetics, and there was no possibility to perform DNA sequencing. Since the age of 11, she had been diagnosed with hyperplasia of the myometrium, polymenorrhea, and chronic posthemorrhagic anemia. The patient had administered contraceptives and hemostatic therapy.

At 14 years of age, the patient was noted to have enlargement of the neck lymph nodes and a 37.8°C fever. Chest computed tomography (CT) imaging, lymph node biopsy were performed and nodular sclerosis HL stage IIB was diagnosed (Fig. 1). Treatment was administered according to EuroNet-PHL Trials, for risk group 2, and consisted of four courses of chemotherapy (2OEPAdac) plus radiation therapy of the involved fields (20 Gy) [4]. Complete remission was achieved and the patient was followed up according to the protocol.

After 6 months, the first early recurrence involving the supraclavicular, axillary lymph nodes, mediastinum lymph nodes, and lungs was diagnosed with biopsy confirmation.
of the same variant of the disease (Fig. 2). The patient received four courses (2IEP, 2DHAP) of second-line treatment and peripheral blood stem cell (PBSC) harvesting (Figure 3) [5, 6]. But at the end of chemotherapy, the mediastinal lymph nodes were observed to be growing again.

The patient received two more courses of VGP (vinorelbine 25 mg/m², gemcitabine 1000 mg/m², prednisolone 100 mg/m²) with a positive response, and autologous stem cell transplantation was subsequently performed. The conditioning regimen consisted of BEAM. A radiation dose of 27 Gy was then carried out on the involved fields [6, 7].

At the final examination 4 months after radiation therapy, chest CT revealed new lesions in the mediastinum and both lungs. Positron emission tomography (PET) scans showed prominent metabolic activity in these lesions. We proceeded with salvage therapy of brentuximab vedotin (1.8 mg/kg every 3 weeks). The only side effect was general deterioration and fever after first administration. No acute reactions or neuropathy were noted during further brentuximab vedotin treatment.

A CT scan after three courses of therapy revealed a remarkable response. After six courses of therapy the CT data showed good partial remission – dramatic shrinkage of mediastinal masses and lung lesions, with a small amount of contrast enhancement and calcifications.

As we did not find either related or unrelated donors the patient received a haploidentical transplant from her father with αβ T-cell receptor depletion. The conditioning regimen consisted of bendamustine, treosulfan, ifosfamide, and antithymocyte globulin. Transplant engraftment was noted at day 19 post transplant [8, 9].

In the early and late post-transplant period, the patient suffered from different complications including late acute graft-versus-host disease (GVHD) of the skin, chronic GVHD of the gut, myometrium hyperplasia with bleeding, bacterial and respiratory syncytial virus pneumonia, pericarditis, chronic heart insufficiency, chronic glomerulonephritis, and avascular necrosis of the joints. The patient received multimodal treatment.

Now, more than 1 year after transplant, according to the last examination (CT, PET), complete remission is confirmed, the girl is doing well, studying in the University.

Figure 1. Pathological and immunohistochemical findings of HL, nodular sclerosis type: At high magnification large Reed-Sternberg cells are seen (A, Hematoxylin-Eosin ×600). The large cells express CD15 (B, ×400), CD30 (C, ×400), Pax5 (D, ×400).

Figure 2. Computer tomography findings at diagnosis, showing large mediastinal mass (arrow).
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Figure 3. Treatment course: OEPA – vincristine 1.5 mg/m²/day №3, etoposide 125 mg/m²/day №5, prednisone 60 mg/m²/day №15, doxorubicin 40 mg/m²/day №2; COPDac – vincristine 1.5 mg/m²/day №2, prednisone 40 mg/m²/day №15, cyclophosphamide 500 mg/m²/day №2, dacarbazine 250 mg/m²/day №3; IEP – ifosfamide 2000 mg/m²/day №5, etoposide 125 mg/m²/day №5, prednisone 100 mg/m²/day №5, DHAP – dexamethasone 24 mg/m²/day №4, Ara-C 2000 mg/m²/twice a day №1, cisplatine 100 mg/m²/day №1; VGP – vinorelbin 25 mg/m², gemcitabine 1000 mg/m², prednisolone 100 mg/m², BEAM – etoposide 200 mg/m²/day №4, melphalan 125 mg/m²/day №1, Ara-C 200 mg/m²/twice a day №4, carmustine 100 mg/m²/day №1; BrVed – Brentuximab vedotin 1.8 mg/kg/day; BTFA, bendarmustine 240 mg/m²/day, treosulphane 42 mg/m², fludarabine 150 mg/m², antitymocyte globuline 50 mg/kg; PBSCT, peripheral blood stem cell transplantation (auto – autologous, haplo – haploidentical); 20 Gy, involved field irradiation; blue arrow – progression; green arrow – response.

Discussion

PS is a rare sporadic disease that causes mosaic overgrowth of tissues of all germ layers. The most frequently affected tissues are bone, skin, adipose, and central nervous system. One of the major clinical findings is distorting, progressive overgrowth that typically starts between ages 6–18 months, a key finding in PS is distortion of the skeletal architecture. Other clinical features include cerebriform connective tissue nevi, linear verrucous epidermal nevus, adipose dysregulation, vascular malformations, organomegaly, bilateral ovarian cystadenoma, parotid monomorphic adenoma, deep vein thrombosis, and bullous pulmonary degeneration [1, 2].

Diagnosis is established by clinical features but can be confirmed by molecular methods. In more than 90% of PS cases, the somatic mosaic mutation in gene AKT1 (c.49G>A [p. Glu17Lys]) located at 14q32.33, is detected [1–3].

The major concern for managing PS is surgical correction of skeletal deformities and bullous pulmonary disease. The prognosis of PS depends mainly on localization of lesions. The majority of manifestations are not life-threatening and do not reduce lifespan with adequate treatment. The most critical situations are deep vein thrombosis, pulmonary embolism, and bullous pulmonary disease [1].

PS can be associated with some oncological diseases. During the second decade of life, the most specific tumors are cystadenoma and parotid monomorphic adenoma. Other tumors reported are meningioma, testicular tumor, astrocytoma, optic nerve tumor, and breast intraductal papilloma. Patients may have multiple tumors simultaneously [1–3].

HL is the malignant tumor of lymphoid tissue with specific granulomatous histologic structure. HL is one of the most frequent of all childhood malignant neoplasms with morbidity of 0.7–0.9 per 100,000 children. The prognosis of HL is favorable and according to international trials data the 5-year event-free survival is approximately 88%, and the overall survival is 97% [10, 11].

Classical HL accounts for approximately 95% of cases, and is subdivided into nodular sclerosis, mixed cellularity, lymphocyte-rich classical, and lymphocyte depletion variants. Tumor cells morphologically present as Reed–Sternberg cells and the immunohistochemical profile CD30+, CD15+, and PAX5+. HL is a radiosensitive and chemosensitive tumor; however, in some cases the disease is refractory to first-line therapy. To improve the results of refractory and relapsed patients while reducing late side effects, recent approaches have been based on new agents such as monoclonal antibodies, HDAC-inhibitors, mTOR inhibitors, and others [12, 13]. One of the most promising agents is brentuximab vedotin, a CD30-directed antibody-drug conjugate. After interaction of anti-CD30 and brentuximab vedotin, a CD30-directed antitubulin drug monomethylauristatin A, and leading to apoptosis. The predominant side effect of brentuximab vedotin is neuropathy [14, 15]. To decrease cytostatic loading brentuximab vedotin seems an excellent therapeutic option, for heavy pretreated relapsed patients, who need PBSCT [16–18].

It is still not well understood which of the prognostic factors plays the main role: age, initial tumor volume, erythrocyte sedimentation rate, B-symptoms, or unknown molecular contributors. In our case, PS is speculated to play a role in such disease behavior, and only haploidentical PBSCT had curative effect. Certainly, the PI3K-AKT pathway plays the role in Proteus Syndrome, and we know that Hodgkin lymphoma is characterized by high mTOR activity in 93% of the cases, and Bcl-xL and NF-kappaB expression correlated with this mTOR activity [19]. High mTOR activity is observed in the case of both favorable and unfavorable clinical response. The presence of mTOR activity probably indicates that the inclusion of mTOR inhibition...
in the therapy of Hodgkin-lymphomas may be feasible and beneficial, especially when standard protocols are ineffective, and it may also allow dose reduction to decrease late treatment toxicity. Most likely, the combination of mTOR inhibitors with other agents will offer the highest efficiency for achieving the best clinical response.

The development of new agents for the treatment of refractory HL gives new hope to heavily pretreated patients. Further trials are needed to understand possible links pathogenesis and to establish the best treatment choice for children with HL and underlying genetic diseases.

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Conflict of Interest

Authors declare no competing financial interests in relation to the work described.

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