An atypical Klippel–Treanunay–Weber syndrome

Abstract

A 44 year old female was referred for hyperbaric oxygen treatment of leg ulcers. The patient had port-wine stain, hand vitiligo, hypertrophy of one extremity, collateral abdominal, and varicose leg veins. Calcium was 12 mg/dL and PTH was 309 ng/L. Hepatomegaly, deep vein thrombosis of the right leg, transudative ascites, hypersplenism, parathyroid adenoma, mesenteric cyst, and papillary thyroid carcinoma were identified. The final clinical diagnosis was Klippel-Treanunay-Weber syndrome.

Several theories such as intrauterine damage to sympathetic ganglia, deep vein abnormalities, mesodermal and ectodermal dysplasia exist for the pathogenesis of this syndrome. This is the first case of Klippel-Treanunay-Weber syndrome with coexisting vitiligo, hypersplenism, non-chirrotic hepatic fibrosis, a mesenteric cyst, a parathyroid adenoma, and a papillary thyroid carcinoma. The vitiligo, mesenteric cyst, parathyroid adenoma, and the papillary thyroid carcinoma are associated with the genetic abnormalities of the syndrome while hypersplenism and hepatic fibrosis are the late complications the disease.

Case Report

A 44 year old female was admitted for hyperbaric oxygen treatment of leg ulcers. On inspection, port-wine stains (Figure 1), vitiligo of the left hand (Figure 2), varicose leg and abdominal collateral veins were observed. Serum biochemistry, and urine analysis were normal but Ca was 12 mg/dL and PTH was 309 ng/L. Chest x-ray showed bilateral hilar lymphadenopathy.

Figure 1: Port-wine stains at the back.
Tuberculin test was negative. ECG revealed a sinus rhythm of 84/minute. Abdominal CT showed vena hepatica thrombosis, ascites, and hepatosplenomegaly. Histopathologic examination of the liver biopsy identified non-chirotic hepatic fibrosis due to portal hypertension. Heterogenous prothrombine gen mutation was positive. Bone marrow biopsy depicted normal cellular elements. The low platelet count of 60x10^3/mm^3 was compatible with hypersplenism. The ascites was transudative due to portal hypertension caused by vena hepatica thrombosis. MRI revealed a 10 cm mesenteric cyst (Figure 3). Ultrasonography showed a thyroid and a parathyroid nodule. Histopathology of the resected thyroid gland was compatible with papillary thyroid carcinoma and parathyroid adenoma. Deep venous thrombosis of the right leg was detected by Doppler ultrasound and enoxaparin 6000 U/day was started. Abdominal lymph node biopsy pathology revealed non-necrotizing granulomatous inflammation. Transbronchial lung and lip biopsy were negative. Serum ACE was 20 U/L. The final diagnosis was Klippel–Treaunay–Weber syndrome with coexisting vitiligo of the hand, hypersplenism, non-chirotic hepatic fibrosis, parathyroid adenoma, and thyroid papillary carcinoma. The patient is currently under follow-up with ongoing enoxaparin treatment.

Discussion

Klippel–Treaunay–Weber syndrome is a triad of port-wine stains, varicose veins, and bony and soft tissue hypertrophy involving an extremity. Klippel and Treaunay first described the syndrome in two patients presenting with a port-wine stain and varicosities of an extremity associated with bone and soft tissue of limb [1,2]. Parkes Weber reported a patient with these symptoms and an arteriovenous malformation of an extremity [3,4]. Currently, it is conflicting how to designate these two syndromes as arteriovenous malformation can lead to a significant prognostic outcome. Exact cause of Klippel–Treaunay–Weber syndrome is unknown. The syndrome may be due to intrauterine damage to the sympathetic ganglia or intermediolateral tract leading to dilated microscopic arteriovenous anastomoses [12], to deep vein abnormalities with resultant obstruction of venous flow leading to venous hypertension and the development of varicese with limb hypertrophy [13], to a mesodermal defect during fetal development that causes maintenance of microscopic arteriovenous communications [14], or to an underlying mixed mesodermal and ectodermal dysplasia [15].

Most cases are sporadic but may carry an autosomal dominant pattern of inheritance [16]. A case KTW syndrome with an unaffected twin supports the paradigmatic genetic pattern [17]. The affiliation amid AGGF1 [angiogenic factor with G patch and FHA domains 1 (AGGF1)] and KTW appears to be significant while common AGGF1 versions may lead to enhanced risk for this syndrome [18]. Kihiczak has reported that KTW syndrome may result from a pathogenic vascular and tissue overgrowth gene [19]. The syndrome has a complex disease pattern and different genetic mechanisms play a role in the development of this genetic disorder leading to variable clinical manifestations that may create a diagnostic challenge for the clinician. Some of these findings may appear as the natural evolution of the disease while others may come out as complications of the syndrome itself. The classical features of the syndrome were present in our case while vitiligo of the hand, parathyroid adenoma, mesenteric cyst, and the papillary thyroid carcinoma may have arisen due to the distinct genetic defects or mutations. These findings may be coincidental but the distinction is not possible in our patient currently. The other clinical manifestations such as hypersplenism, venous thrombosis, and hepatic fibrosis, on the other hand, may have occurred as the expected long-term complications of the disease. Our patient was a sporadic case without a previous family history.

Nearly all the patients demonstrate all three signs of the clinical syndrome: port-wine stain, varicose veins, and bony and soft tissue hypertrophies. In a series of 252 patients at the Mayo Clinic, 63% of patients had all three features and 37% had two of the three features. Port-wine stain was seen in 98% of patients, varicosities or venous malformations in 72%, and limb hypertrophy in 67% of the patients [20]. Brain abnormalities such as hemorrhage, infarction, hemimegalencephaly, venous malformation, arteriovenous malformations, cavernoma, aneurysm, hydrocephalus, choroid plexus abnormalities, atrophy, calcifications, leptomeningeal enhancement, cortical dysplasia, and seizures may occur while cerebral infarctions and brain tumors are encountered rarely [21–23]. Pulmonary emboli may develop secondary to venous limb thrombosis [24]. Consequently, KTW syndrome may present with many different manifestations due to the genetic structure and the anatomic location or the type of vascular malformations as it is the case in our patient.

Our case is unique for the manifestations such as the vitiligo of the hand, parathyroid adenoma, mesenteric cyst,
and the papillary thyroid carcinoma that have not been described in the literature previously. On the other hand, vena hepatica thrombosis, hypersplenism, hepatic fibrosis, and deep vein thrombosis are expected long term complications of this syndrome. Clinicians should bear in mind that KTW syndrome may present with an unusual clinical profile that may be either due to the primary genetic profile of the patient or the secondary to the long-term complications of the disease. Complex genetic pattern or genetic mutations may lead to an atypical presentation in patients with the KTW syndrome. The case was sporadic in nature. An atypical presentation due to the variable genetic profile of the syndrome with coexisting disease complications was considered in this patient.

Conclusions

Klippel-Trenaunay-Weber syndrome is a rare and a sporadic genetic disease. The syndrome may present with atypical manifestations that are probably related to the genetic differences or mutations of the adenoma. The vitiligo of the hand, mesenteric cyst, parathyroid adenoma, and the papillary thyroid carcinoma may result from chromosomal abnormalities, in utero mesodermal and ectodermal dysplasia. On the other hand, long term complications of the syndrome accompany the clinical profile thereby leading to a diagnostic challenge. Clinicians should bear in mind that the atypical presentation may be related to the genetic features along with the expected complications of this syndrome. Differential diagnosis of Klippel-Trenaunay-Weber syndrome from other capillary malformation—overgrowth syndromes such as isolated capillary malformation, diffuse capillary malformation with overgrowth (DCMO), and other overgrowth syndromes is necessary. The PROS term can encompass different overlapping conditions of this syndrome and thereby may better designate such patients.

Author contributions

Muammer Bilir has contributed to patient evaluation and follow-up.

Cuneyt Tetikkurt has performed case report preparation and written the case report.

Halil Yanardag has contributed to patient data evaluation and references.

References


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DOI: http://dx.doi.org/10.17352/aprc.000033