Research Article

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An unusual evolution of a case of Klippel-Trenaunay syndrome

DOI 10.1515/med-2015-0084 Received October 27, 2015; accepted November 2, 2015

Abstract: Klippel–Trenaunay syndrome (KTS) is a rare congenital disorder. KTS can be diagnosed on the basis of any 2 of 3 features: cutaneous capillary malformations, soft tissue or bony hypertrophy and varicose veins. We present an unusual case of KTS complicated by an infection of venous ulcers of the lower limb by larvae. The treatment of infection was a complete debridement; however baseline treatment of KTS is still in evaluation.

Keywords: Klippel-Trenaunay syndrome, maggots, larvae therapy

1 Introduction

Klippel–Trenaunay syndrome (KTS) is a rare congenital disorder due to a sporadic, autosomal dominant or mosaic homozygosity mutation. KTS syndrome is characterized by a triad of varicose veins, cutaneous capillary malformation with dermatomal distribution, and hypertrophy of bone and soft tissue [1]. The prevalence of KTS is about 1:100000 livebirths [2]. Cutaneous capillary malformations are usually red/purple "port wine type" vascular nevus. Limb hypertrophy is due to bone hypertrophy and secondary soft tissue overgrowth. Varicosities are due the presence of a large incompetent vein on the lateral aspect of the affected leg. The classic triad is hallmark, however patients may not always present with all three symptoms simultaneously.

KTS was first described in 1900 by the French physicians Klippel and Trenaunay [3] and was classified by You in 1983 into 5 levels of severity [4]. There is no documented gender or racial predilection but cases of similar demographic [6]. Complete pattern of KTS has not yet been demonstrated but alterations in in the VG5Q gene on chromosome 5 may result in the vascular defects. In some patients the mutation of chromosomes 8q22.3 and 14q13 have been observed. Life expectancy depends on the severity of the malformation.

Suspected patients should be evaluated with imaging studies in order to differentiate hemangiomas from vascular malformations: CT and MRI should be used for visualizing the extent of the lesions and the deep of tissue infiltration [7].

Additional imaging studies can be used to improve the diagnosis (color-doppler ultrasound, radiography, ascending phlebogram with or without contrast material).

The complicated lesions are the most common causes of debilitating pain in patients with KTS.

Frequent complications related to venous abnormalities are: chronic venous insufficiency, cellulitis, infections, superficial thrombophlebitis and deep vein thrombosis [8].

Other complication, described by Maari et al [9] in 2004, were: leg length discrepancy, cellulitis, severe pain, thrombophlebitis, pulmonary embolus, coagulopathy, mental retardation, necessity of limb amputation, gastrointestinal bleeding and death.

Patients with KTS should receive multidisciplinary medical care. Treatment of KTS patients has consisted mainly of conservative medical management, including compressive stockings and anti-inflammatory medications for pain relief. Operative treatment has been controversial and surgery on the superficial venous system has been reserved for patients with intact deep systems only. Patients with patent deep veins can be considered for excision of symptomatic varicose veins and VMs venous malformations [10].

Frasier et al [11] describe an alternative treatment of superficial venous varices utilizing endovenous thermal

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ablation via radiofrequency (RFA) complimented by ultrasound guided sclerotherapy in KTS patients as a minimally invasive and potentially efficacious treatment.

Open technique or subfascial endoscopic perforator surgery are also described for incompetent perforator veins surgery.

Excision of varicosities is often incomplete and VMs may recur in 50% of patients. For this reason the preferred treatment in uncomplicated cases is, in first instance, conservative approach [10].

2 Case Report

We present a case of a 77-year old woman that was referred to our Emergency Department in December 2014 because of intense pain due to ulcers of the lower limbs and asthenia. Her medical history was significant for diabetes, arterial hypertension, chronic gastropathy and previous cholecystectomy, hysterectomy and left hip prosthesis. The patient was seen by our deparment 5 months prior in order to evaluate varicose veins (Fig. 1). She was affected by Klippel-Trenaunay Syndrome (diagnosed 30 years before). The patient's right shoulder displayed a port wine stain and a slight deformity of the right leg; the right leg of the patient was longer than the other one from birth. We proposed another screening with color-doppler ultrasound, ascending phlebogram and vascular-RM to confirm the diagnosis of KTS but the patient refused it. From then, we have not had any news of patient up to the access at the Emergency Department.

On admission, the patient presented with dirty bandages, upon removal of the dressings, multiple ulcerations were present with larval overlay that consumed the entire leg extending proximally to the pubis.

The ulcers were were clean. We cleansed the leg in a solution of povidone-iodine and physiological solution for 15 minutes. Than we rinse the leg with plenty of water. The procedure was repeated three times and parasites were completely removed.

Due to the patient's obvious failure to thrive, she was admitted to the Medicine Department. She was treated with broad-spectrum antibiotics and supportive therapy. Larvae were analyzed and were classified in: houseflies, cheeseflies and sarcophagaflies.

One day after admission, we observed additional larval presence in her wounds. We repeated our cleaning regimen performed one day priod. Complete drainage required 10 days of treatment. No sign of bacterial infection was observed. Currently the patient has been assigned to a social worker and she has agreed to perform the complete screening for KTS to confirm the diagnosis.

Ethical approval: The research related to human use has been complied with all the relevant national regulations, institutional policies and in accordance the tenets of the Helsinki Declaration, and has been approved by the authors' institutional review board or equivalent committee.

Informed consent: Informed consent has been obtained from all individuals included in this study.

3 Discussion

KTS can be diagnosed on the basis of any 2 of 3 features: cutaneous capillary malformations, soft tissue or bony hypertrophy and varicose veins [12].

You [4] in 1983 described 4 levels of severity in KTS respect to varicose veins: in the I Class the features are venous dysplasia and phlebectasic dysplasia; the peculiarity of II Class is Arterial dysplasia. In the III Class described arterial and associated venous dysplasias, without arteriovenous phlebectasia shunts and angiodysloaisas with shunt (Kllippel-Treaunay-Weber syndrome). The most serious Class is the IV with mixed angiodysplasias (atypical KTS) [4].

Our patient belongs to the first class. The diagnosis more over are those made in children who have a deformity of a limb; in our case the diagnosis was made when the patient was 40 years old.

After diagnosis, the patient had not been followed by the specialist and no therapies was performed.

Some authors described an association with parent but without a statistical significance [13]. In our case we had no information in regards to her family medical history. Ulcerations associated with diabetes are the most common cause of foot ulcers and approximately 15% of indvidulas with diabetes will develop foot ulceration during their lifetime [14]. In literature there is no evidence of association between diabetes and KTS but diabetes is a worsening factor in development of venous ulcers. Usually, the chronic wound microenvironment is complex and contaminated by more than one species of bacteria. In our report the patient had a colonization of the bandage of her leg from fly eggs. In a second time, the development of larvae have colonized ulcerating tissue. Finally, necrotic tissue allowed the larvae propagate. Bacteria produee biofilm as a protective mechanism which subsequently leads to an increase in an resistance from their hosts and their defenses, antiseptics, and topical or systemic antibiotics [15]. In our study, we propose that the larvae were beneficial to the integrity of the patient's wounds. A complete debridement of necrosis and devitalized

and infected tissue was carried out by the larvae. The beneficial effects of using larvae in wounds were first noticed by Ambrose Paré in 1557 [16]. The first clinical application of larvae therapy was performed by Zacharias and Jones during the American Civil War [17].

Scientists first postulated that the debriding action of larvae was due to their mechanical wriggling using a pair of mandibles/hooks for movement and attachment [18].

Recently Chamers et al. described three proteolytic enzyme classes have been identified in the maggot excretions that can degrade extracellular matrix components, including laminin and fibronectin [19-25].

Also in our case the maggots were the responsible factor of complete debridement of necrosis and sovrainfected tissue. After this event the patient is back to being completely asymptomatic.

Usually the patients with KTS are successfully treated with surgery or radiofrequency ablation of incompetent veins; probably, in our case, because of asintomaticity after conservative treatment and because patients' history. Unusually, in our case, the therapy of infective complication of venous ulcers in KTS, was larvae.

4 Conclusion

KTS is a rare syndrome that involves multiple clinical presentations. The most complex cases require a surgical therapy while simple cases only a conservative therapy. In the case of complications due to venous ulcers linked to the syndrome, classic loco-regional treatment and constant monitoring is preferred.

The report we analyzed is certainly a strange case of superinfection and therapy at the same time in a patient with complicated venous ulcerations in KTS.

Conflict of interest statement: Authors state no conflict of interest

References

- Kihiczak GG, Meine JG, Schwartz RA, Janniger CK. Klippel– Trenaunay syndrome: A multisystem disorder possibly resulting from a pathogenic gene for vascular and tissue overgrowth. Int J Dermat 2006;45:883-90.
- [2] Lorda-Sanchez I, Prieto L, Rodriguez-Pinilla E, Martinez-Frias ML. Increased parental age and number of pregnancies in Klippel-Trenaunay-Weber syndrome. Ann Hum Genet 1998;62:235-39.
- [3] Klippel M, Trenaunay P. Du naevus variqueux osteohypertrophique. Arch General Med 1900;185:641-72.

- [4] You CK, Rees J, Gillis DA, Steeves J. Klippel–Trenaunay syndrome: a review. Can J Surg 1983;26:399-403.
- [5] Wang Q, Timur AA, Szafranski P, Sadgephour A, Jurecic V, Cowell J, et al. Identification and molecular characterization of de novo translocation t(8;14)(q22.3;q13) associated with a vascular and tissue overgrowth syndrome. Cytogenet Cell Genet 2001;95:183–8.
- [6] Craven N, Wright AL. Familial Klippel-Trenaunay syndrome: a case report. Clin Exp Dermatol 1995;20:76–9.
- [7] Dubois J, Garel L, Grignon A, David M, Laberge L, Filiatrault D, et al. Imaging of hemangiomas and vascular malformations in children. Acad Radiol 1998;5:390–400.
- [8] Lee A, Driscoll D, Gloviczki P, Clay R, Shaughnessy W, Stans A. Evaluation and management of pain in patients with Klippel-Trenaunay syndrome: a review. Pediatrics 2005;115:744-9.
- [9] Maari C, Frieden IJ. Klippel-Trénaunay syndrome: the importance of "geographic stains" in identifying lymphatic disease and risk of complications. J Am Acad Dermatol 2004;51:391-8.
- [10] Noel AA, Gloviczki P, Cherry KJ Jr, Rooke TW, Stanson AW, Driscoll DJ. Surgical treatment of venous malformations in Klippel-Trénaunay syndrome. J Vasc Surg 2000;32:840-7.
- [11] Frasier K1, Giangola G, Rosen R, Ginat DT. Endovascular radiofrequency ablation: a novel treatment of venous insufficiency in Klippel-Trenaunay patients. J Vasc Surg 2008;47:1339-45.
- [12] Capraro PA, Fisher J, Hammond DC, Grossman JA. Klippel-Trenaunay syndrome. Plast Reconstr Surg 2002;109:2052-9.
- [13] Delis KT, Gloviczki P, Wennberg PW, Rooke TW, Driscoll DJ. Hemodynamic impairment, venous segmental disease, and clinical severity scoring in limbs with Klippel-Trenaunay syndrome. J Vasc Surg 2007;45:561-7.
- [14] Pecoraro RE, Reiber GE, Burgess EM. Pathways to diabetic limb amputation: Basis for prevention. Diabetes Care 1990,13: 513-21.
- [15] O'Meara S, Al-Kurdi D, Ologun Y, Ovington LG. Antibiotics and antiseptics for venous leg ulcers. Cochrane Database Syst Rev. 2010;20:CD003557.
- [16] Paré A. The battell of S. Quintin (1557). In: Keynes G, editor. The Apologie and Treatise of Ambroise Paré. Chicago: The University of Chicago Press; 1952:68-70.
- [17] Mumcuoglu KY. Clinical applications for maggots in wound care. Am J Clin Dermatol 2001;2:219-27.
- [18] Barnard DR. Skeletal-muscular mechanisms of the larva of Lucilia sericata (Meigen) in relation to feeding habit. Pan-Pac Entomol 1977;53:223-9.
- [19] Chambers L, Woodrow S, Brown AP, Harris PD, Phillips D, Hall M, et al. Degradation of extracellular matrix components by defined proteinases from the greenbottle larva Lucilia sericata used for the clinical debridement of non-healing wounds. Br J Dermatol 2003;148:14-23.
- [20] Serra, R., Buffone, G., Molinari, V., et al.: Low molecular weight heparin improves healing of chronic venous ulcers especially in the elderly. *International Wound Journal*, 2013, Article in press
- [21] Persico, G., Amato, B., Aprea, G., et al. : The early effects of intravenous L-propionyl carnitine on ulcerative trophic lesions of the lower limbs in arteriopathic patients: A controlled randomized study. *Drugs under Experimental and Clinical Research*, 1995, 21: 5, 187-198

- [22] Amato B., Coretti G., Compagna R., et al: Role of matrix metalloproteinases in non-healing venous ulcers. *International Wound Journal*, 2013, Article in press
- [23] Serra R., Grande R., Buffone G., et al.: Extracellular matrix assessment of infected chronic venous leg ulcers: Role of metalloproteinases and inflammatory cytokines. *International Wound Journal*, 2014 Article in press
- [24] De Franciscis S., Gasbarro V., Amato B., Buffone G., Grande R., Serra, R.: Hemodynamic surgery versus conventional surgery in chronic venous disease: A multicenter retrospective study . Acta Phlebologica, 2013, 14: 3, 109-114
- [25] Serra R., Gallelli L., Conti A., et al. : The effects of sulodexide on both clinical and molecular parameters in patients with mixed arterial and venous ulcers of lower limbs. *Drug Design*, *Development and Therapy*, 2014, 8: 13, 519-527