CASE REPORT

KLIPPEL TRENAUNAY SYNDROME: A CASE REPORT
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ABSTRACT: INTRODUCTION: Klippel Trenaunay Syndrome is a rare congenital disorder, but it is the most common condition involving combined vascular malformation. KTS was first described by two French Physicians, Klippel and Trenaunay in 1900.¹ Incidence of KTS reported is approximately 2 to 5 cases per 100,000 live births.²,³ KTS generally affects a single extremity, although cases of multiple affected limbs have been reported. The leg is most common site followed by the arms, the trunk, and rarely the head and neck. The original description of KTS included limb hypertrophy, varicose veins and vascular (Port wine) nevus, which were characterised as a clinical triad. Hemangiomas are the most frequent finding in these patients and is usually present at birth.⁴ KTS is also known as angio-osteohypertrophy syndrome, congenital dysplastic angiopathy or klippel trenaunay weber syndrome.

KEYWORDS: Klippel trenaunay syndrome (KTS), vascular malformation, soft tissue hypertrophy, venous varicosity.

CASE REPORT: A four year old female child presented to our OPD with complaints of progressive swelling of whole of right lower limb since birth. Swelling was present since birth and has slowly progressed to present size. She had no history of tuberculosis, fever with chills and rigor, trauma, and any other bleeding manifestation.

On examination, the patient was afebrile, moderate pallor was present. No clubbing, cyanosis or lymphadenopathy was present. Heart rate = 110 bpm, blood pressure = 96/60 mmHg, respiratory rate= 20 /min. Cardiopulmonary examination was unremarkable except tachycardia. Patient was cooperative but was little shy. All milestones were normal for age. No history of similar complaints in siblings. She had marked hypertrophy of her right lower limb. Her right calf and thigh were swollen and non-tender. Superficial varicose veins were palpated over her right calf muscles. No discrepancy in length was noted in two lower extremities. Slight pigmentation was noted near the tortuous veins, however no venous ulcers were present.

On lab investigations, her hemoglobin was 7.6 gm%, PBS- microcytic hypochromic anemia was present. Bleeding time, clotting time, prothrombin time, activated partial thromboplastin time were all normal. LFT and RFT were all normal. Chest x-ray and ECG were normal. Plain X-ray of the right hip and right leg revealed hypertrophy of soft tissues and bones. USG abdomen showed no organomegaly. Colour doppler of right lower limb demonstrated significant reflux at saphenofemoral junction. Superficial veins showed increased flow. Deep venous system was not present in right lower extremity. But in the left lower extremity superficial and deep venous system were normal. Echocardiography was done to rule out any possibility of cardiac overload, failure or any associated heart defects. However, echocardiography revealed no significant abnormality. Due to financial constraints chromosomal analysis was not done in this patient. The patient is now on regular follow up.

DISCUSSION: This case is a variant of KT Syndrome. KTS is a rare congenital malformation characterised by the triad of capillary malformation, venous malformation and bone or soft tissue hypertrophy.
hypertrophy. The most frequent clinical aspect of KTS is a dermal capillary stain associated with
dilated marginal vein of Servelle often identified in the subcutaneous fat of the lateral calf and thigh.
The capillary malformation, venous varicosities and limb hypertrophy has been found to occur in
98%, 72% and 67% of children with KTS respectively (Jacob et al).\(^5\) It should be distinguished from
Parkes weber syndrome by absence of underlying atriovenous malformation. Prominent superficial
varicose veins are present in majority of patients.\(^6\) Extremity pain, spontaneous hemorrhages,
venous insufficiency or thrombophlebitis are commonly encountered.\(^6\)

The cause of KTS remains unclear, however many workers believe it to be caused by
mesodermal abnormality during fetal development leading to vascular and soft tissue malformations
in the affected limb (Baskerville et al, 1985).\(^7\) The de novo translocation t(8;14) (q22. 3;q13) has
also been reported (Wang et al, 2001).\(^8\) Most cases of KTS are sporadic. But there are certain cases
reported which suggests an autosomal dominant pattern of inheritance.\(^9\) Happle (1993) suggested a
paradominant inheritance, caused by single gene defect. The trait is only expressed when a somatic
mutation occurs in the normal allele at an early stage of embryogenesis, giving rise to a clonal
proliferation of cells either homozygous or hemizygous for KTS mutation.\(^10\)

Ultrasonography, Doppler study, contrast enhanced MRI, Venogram are required for
definitive diagnosis of KTS. Management of KTS is dependent upon individual symptoms. Although
both non operative and surgical approaches are used, management is mainly supportive and
symptomatic. Compressive garments are advised for chronic venous insufficiency, lymphedema,
recurrent cellulitis and recurrent bleeding from the capillary or venous malformation. Pain
medication, antibiotics and limb elevation are all used to manage cellulitis. Many studies have given
positive results in patients using compression therapy (Stringel and Dastous, 1987). Laser therapy
indicated when there is ulceration and breakdown of lymphatic blebs. Surgery includes vein ligation,
vein resection and in rare cases, removal of problematic area of abnormal tissue (debunking surgery)
and amputation. Rapamycin has also shown to improve quality of life in these patients by improving
pain, decreasing bleeding from lymphatic blebs.

**Fig. 1:** Doppler image demonstrating significance reflux at sapheno-femoral junction.
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**Fig. 2:** Colour Doppler showing absent deep venous system.

**Fig. 3:** Doppler image showing continuous reflux in sapheno–femoral junction.
REFERENCES:


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