

Klippel-Trénaunay syndrome associated with chronic disseminated intravascular coagulation and massive osteolysis

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The Klippel-Trénaunay syndrome is characterized by a hemangiomatous lesion of a limb with varices and limb hyperplasia. Recently a boy was evaluated with this syndrome in association with two other rare conditions, chronic disseminated intravascular coagulation, and massive osteolysis. These disorders are discussed and their possible interrelationships explored.

Case report

A 15-year-old boy was first examined at the Cleveland Clinic in March 1972. He was the second of two children; his mother had had a normal pregnancy, labor, and delivery. As a neonate, a large hemangioma involving the left buttock and part of the left thigh was noted. Circumcision at that time was uneventful. No medical problems were described until the patient was 13 months old when he reportedly had a pathologic fracture of the left tibia which healed poorly despite extensive therapy. Consultants concluded that the hemangioma over the left limb involved the tibia causing delayed healing of the fracture. Following that event, little growth of the left leg was attained, although the limbs were described as being equal in length prior to that time.

The patient did well until March 1972, when he was first examined for continued oozing from the site of plantar wart surgery. No history of epistaxes, bleeding gums, or petechiae was elicited, although

he did complain of always having bruised easily. Physical examination revealed a healthy looking boy without evidence of bruising or petechiae. A large cavernous hemangioma involving the left buttock and extending onto the thigh of a shortened left limb was the only physical finding of note (*Fig. 1A and B*).

Laboratory studies (*Table*) revealed a low fibrinogen level and mild thrombocytopenia, slightly prolonged thrombin time, abnormally accelerated euglobulin lysis time, and the presence of fibrin split

products as measured by the Fi test,¹ protamine sulfate paracoagulation test,² and Thrombo-Wellcotest test.³ Peripheral blood smears demonstrated morphologic changes compatible with a microangiopathic hemolytic anemia. The condition was diagnosed as disseminated intravascular coagulation, but no specific therapy was advised because of the patient's relatively asymptomatic state. Wound healing progressed normally without further bleeding.

In October 1972 the patient was exam-

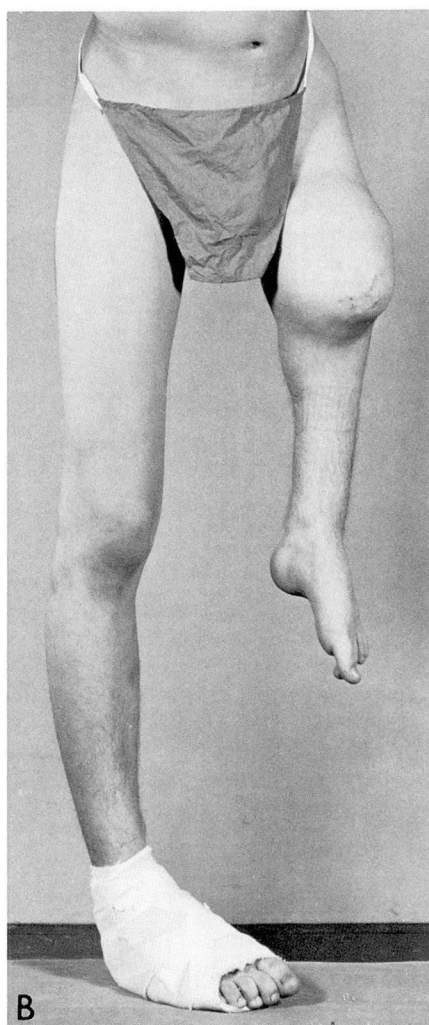
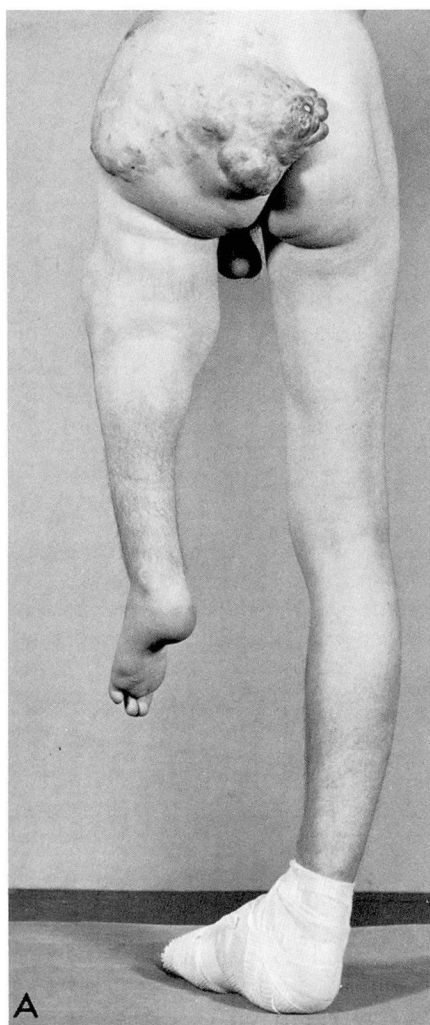


Fig. 1. **A**, Posterior view of patient at age 15 demonstrating diffuse hemangioma involving the left buttock and thigh with limb shortening. **B**, Anterior view.

Table. Results of laboratory studies

Test	Date		
	March 1972	October 1972	May 1975
Platelet count	136,000/mm ³	100,000/mm ³	110,000/mm ³
Fibrinogen	71 mg/dl	50 mg/dl	64 mg/dl
Prothrombin time	14 sec (12 sec)*	13 sec (12 sec)*	17 sec (12 sec)*
Activated partial thromboplastin time	37 sec	...	34 sec
Factor V	75%
Factor VII	80%
Thrombin time	23 sec (15 sec)*	...	20 sec (12 sec)*
Euglobulin lysis	2 hr	50 min	30 min
Fibrin split products			
Fi test	+
Protamine sulfate	+	+	+
Thrombo-Wellcotest	40 µg/ml
Bleeding time, Ivy	15 min

* Controls.

ined for the sudden onset of pain in the left thigh following a minor fall. Roentgenograms showed a pathologic fracture of the left femoral head, total resorption of the midshaft of the left femur, and demineralization of the bones of the left hemipelvis, knee, fibula, and foot.

In May 1975 he returned with a chief complaint of severe left hip pain of 1 week's duration. He had experienced intermittent dull left hip pain once or twice a year for the previous 6 years. Physical examination revealed a well-developed 17-year-old boy in no distress. The general physical and neurologic examinations were normal except for several small petechiae, a considerably shortened left limb covered by a varicose cavernous hemangioma, and a mild lumbosacral scoliosis. No bruits were heard over the hemangioma. The patient walked well with a left limb brace and without apparent pain at that time. Roentgenographic examinations of the left extremity showed massive osteolysis of the femur with disappearance of all but the femoral head. The left hemipelvis, left tibia, and left fibula were also affected by the osteolytic process (Fig. 2). Results of laboratory tests showed a coagulation pattern and red blood cell morphology essentially un-

changed from that identified 3 years previously. The patient experienced no further pain while hospitalized nor at follow-up 2 weeks later.

Discussion

The triad of cavernous heman-gioma of a limb associated with var-ices and soft tissue and bone hyper-trophy was first described by Klippel and Trénaunay⁴ in 1900. More recent observers⁵⁻⁹ have demonstrated the presence of a malformed deep ve-nous system within the angioma. The formation of varices and subsequent hypertrophy of the involved extrem-ity are believed to follow the devel-opment of venous stasis within the abnormal vessels. Children with the typical manifestation of this disorder frequently do not develop varices or limb hypertrophy until several years after the hemangioma is first discov-ered.⁹⁻¹² In fact, not only may this disorder be present without limb hy-pertrophy, but it also may be associ-ated with hypoplasia of the involved extremity.^{7, 9} Therefore, the limb hy-poplasia in this case is consistent with

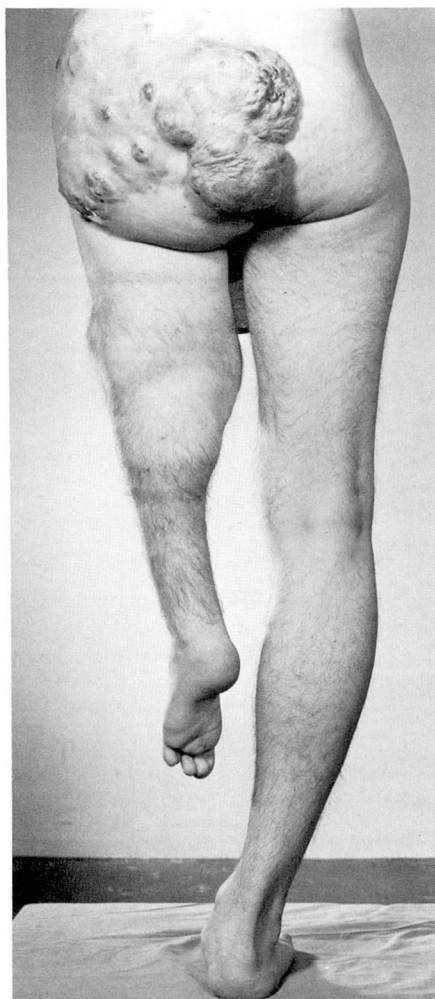


Fig. 2. Posterior view of patient at age 17 showing increase in size of the hemangioma.

previous reports of the Klippel-Trénaunay syndrome. The possibility exists, however, that this patient's limb hypoplasia may have been related to osteolytic changes of the bones rather than secondary to the vascular malformation.

The association of thrombocytopenia with large hemangiomata was first reported in 1940 by Kasabach and Merritt,¹³ but it was not until more recent times that aberrations in

the coagulation mechanism itself were described.¹⁴⁻¹⁷ The main characteristics of this coagulopathy are moderate reduction in platelets, reduction in fibrinogen and other plasma coagulation factors, evidence of fibrinolysis, and the presence of fibrin split products. In studying this phenomenon, Gilon et al¹⁶ demonstrated a relative decrease in the peripheral blood platelet count as opposed to that within the hemangiomatous mass in patients with this disorder, a finding that was later substantiated by Petit et al¹⁸ using chromium-labeled platelets. Gilon et al¹⁶ theorized that the tortuosity of the vessels within the angioma resulted in the trapping and subsequent destruction of platelets which, in turn, led to the activation of the coagulation mechanism and a localized intravascular coagulopathy. This theory is supported by the discovery that although the prothrombin and partial thromboplastin times are usually normal or only slightly prolonged in the peripheral blood of these patients, samples from within the hemangioma itself show greatly prolonged values.¹⁹ Heparinization, high doses of corticosteroids, selective surgical removal of the hemangiomatous mass, and local radiation therapy have been used in an attempt to control this disorder; however, only the latter two methods have been shown to be of any long-term value.^{15, 17}

Massive osteolysis is a disease of bone characterized by progressive demineralization and replacement of bone by fibrous connective tissue.²⁰ Minor trauma frequently heralds its onset. Jackson²¹ first described this disease in 1838 when he reported the case of a young man in whom osteol-

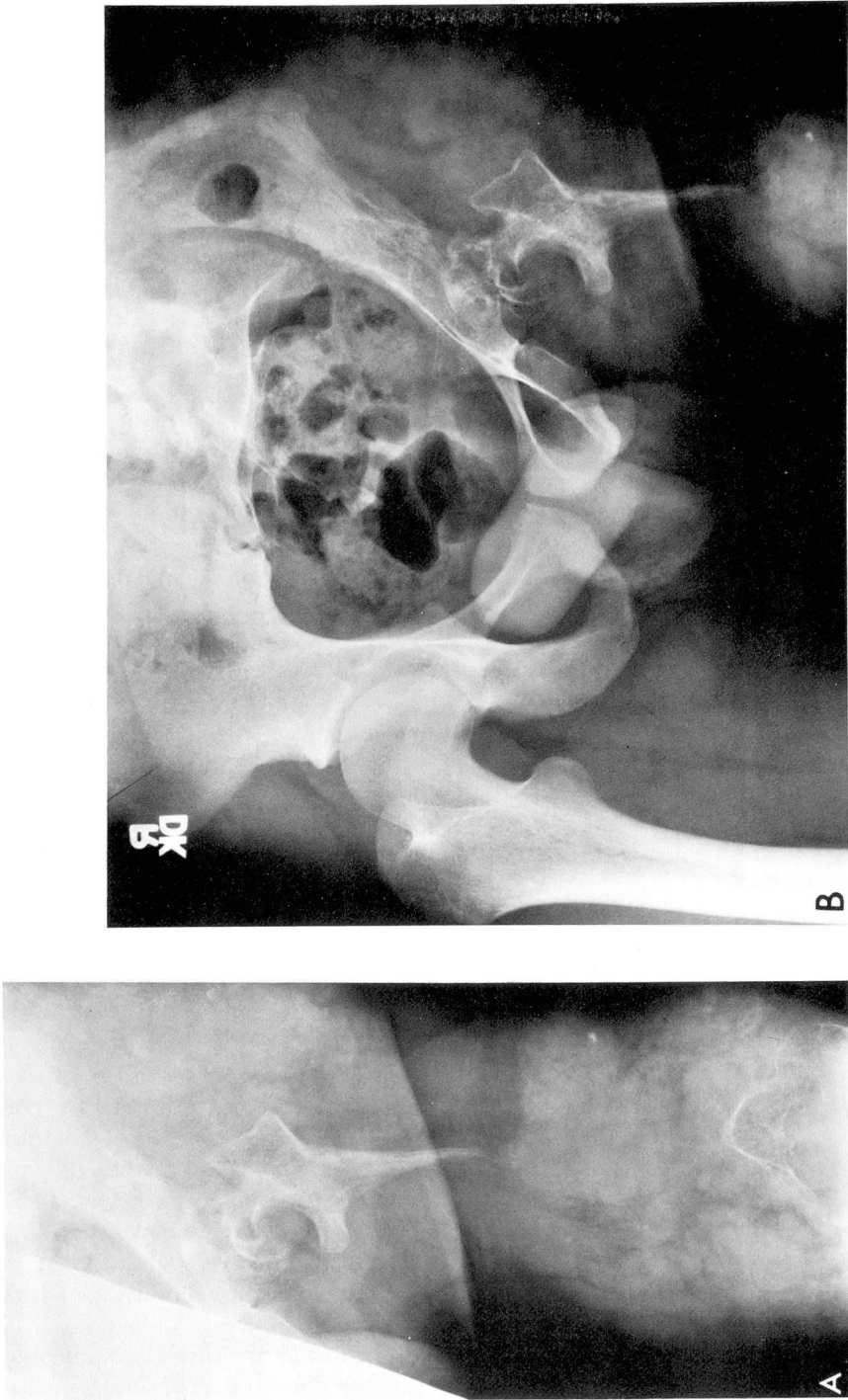


Fig. 3. A, Roentgenogram of left femur at age 17 showing the tapering characteristic in massive osteolysis. B, Roentgenogram of pelvis demonstrating osteolytic involvement of the left hemipelvis.

ysis of the humerus developed following a simple fracture. The osteolytic process progressed over a period of years until the humerus was totally replaced by a thin fibrous band.

Massive osteolysis may be isolated to the originally involved structure or may go on to involve contiguous bones. In more than 130 reported cases, males appear to be equally as affected as females, and the average age of onset is in late adolescence with a range from 18 months to 58 years. The rapidity and extent of bony resorption is unpredictable, showing any pattern from persistent, relentless destruction to mild demineralization with spontaneous remission. This process usually ends in the development of a flail limb; however, if the chest wall is involved, massive chylothorax and death frequently ensue. The bones most commonly affected are the ribs and scapulae followed by the legs, arms, pelvis, head, and neck.²² Biopsy specimens of the bone and its surrounding tissues show numerous thin-walled, widely dilated vessels.²³

This histologic finding and the fact that systemic hemangiomatosis has been reported in association with massive osteolysis^{12, 24-26} led to the theory that an angiomatous malformation was the cause of this curious bone lesion.²³ The question arises, however, whether the hemangiomatosis observed clinically and apparently histologically is directly related to the development of massive osteolysis. Hemangiomata with their origin in bone are well-known entities, but their roentgenographic and histologic features are distinct and quite unlike those described in massive osteolysis.²⁷ However, some hemangiomata of soft tissue possess vascular

patterns that promote localized hyperoxygenation, a factor known to increase bone resorption.²⁸ To investigate this possibility, angiography has been performed on extremities involved in massive osteolysis; however, a change in circulation time compatible with a state of hyperoxygenation was not demonstrated.^{25, 29}

Some investigators believe that the dilated vessels present in biopsy specimens of massive osteolysis may not be related to hemangiomatosis, but may actually represent a compensatory increase in the size of the bone marrow's rich vascular plexus following resorption of its bony matrix.^{29, 30} Although more data would be necessary before any conclusions could be reached, the evidence at present does not appear to support the theory that the bony resorption of massive osteolysis is secondary to an intrinsic vascular malformation.

Another possible cause of massive osteolysis is a reversal of the normal osteoblastic to osteoclastic ratio.²⁸ Some researchers have reported increased numbers of osteoclasts in specimens of massive osteolysis,^{29, 31, 32} but this has not been a consistent finding.³³ Therefore, despite several theories, the cause of this condition is still not determined.

The roentgenographic stages of osteolysis have been described by Torg and Steel.³³ They include (1) nondescript patchy osteoporosis, (2) concentric shrinkage (tapering of the involved ends of tubular bones), (3) complete resorption, and (4) progression to contiguous bony structures (*Fig. 3A and B*).

The initial differential diagnosis of massive osteolysis should include such disorders as Sudek's atrophy, atrophy of disuse, and bony changes

secondary to infectious and metabolic derangements. However, after stage 1, the roentgenographic appearance of this disease is so characteristic as to exclude most other diagnostic entities. The only disorder that could be confused with massive osteolysis in later stages is that of acroosteolysis, an autosomal dominant disease with osteolysis of the carpal and tarsal bones. The main differentiating features are that acroosteolysis is hereditary, affects only the distal bones of the hands and feet rather than those of the proximal extremities, and characteristically involves noncontiguous bones.³⁴

Attempts at definitive therapy of massive osteolysis have included radiation therapy, high-dose corticosteroids, bone transplants, vitamin D, parathyroid hormone, and calcium.²² Successfully treated cases have been reported, but there is little to suggest that any one therapy is curative.

Although one cannot draw any definite conclusions on the role of this patient's hemangioma in the development of chronic disseminated intravascular coagulation and massive osteolysis, experimental evidence does support a direct relationship between the vascular malformation of the Klippel-Trénaunay syndrome and the appearance of a consumption coagulopathy. The same cause and effect relationship between hemangiomatous lesions and massive osteolysis has not been conclusively proved to date; yet the occurrence of these three rare disease states in the same patient leads to speculation concerning their possible interrelationships.

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