Acute Knee Pain in a Child Due to Pigmented Villonodular Synovitis

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What to Learn from this Article?

PVNS should be considered as one of the differentials in acute irritable knee in children. Prompt and proper treatment can lead to an excellent clinical result.

Abstract

Introduction: Pigmented villonodular synovitis (PVNS) is a proliferative condition of the synovium, which is composed of nodules and/or villi and has an abundant number of hemosiderin-laden macrophages [1]. It is rare in children. A child with acute irritable knee mimicking infective arthritis of the knee joint was presented. Arthroscopy revealed a vascular mass with active contact bleeding, with histology confirmed a diagnosis of PVNS. The presence of hemarthrosis suggested that acute bleeding of the lesion into the joint may be the cause of such an atypical acute presentation. PVNS, although rare, should be considered as one of the differential diagnoses of an acute irritable knee presenting in children.

Case report

A 10-year-old boy had sudden onset of left knee pain and swelling 1-day before admission. There was no recent injury reported. He was unable to move the knee or walk due to the pain. There was no other joint pain, fever, or systemic upset. Clinical examination showed an irritable boy with marked left knee effusion. The knee joint was extremely painful with diffuse tenderness and increase in local temperature. The joint was not erythematous. The range of movement of the knee was markedly limited from 30° to 40°. Knee aspiration yielded 40 ml of blood-stained fluid. Radiographs of the knee showed local soft tissue swelling without any bone lesion (Fig. 1). Emergency arthroscopy was performed. Arthroscopic findings showed a 4 cm × 3 cm multi-lobulated vascular mass at the intercondylar notch with active contact bleeding. The mass was so big that it essentially occupied the arthroscopic view of the intercondylar notch (Fig. 2). There was no other similar lesion in any other area of the joint. Complete resection of the mass was performed through open arthrotomy. Histological examination of the resected mass confirmed the diagnosis of PVNS with areas of necrosis (Fig. 3). All specimens had negative bacterial culture. Knee pain and swelling improved gradually after surgery. By 2 months post operation, there was no more knee pain and the patient could run and squat as usual. Magnetic resonance imaging performed at 3 months post operation showed no residual lesion (Fig. 4). The latest
follow-up at 1-year post operation was normal with no sign of local recurrence.

Discussion

Acute, non-traumatic, irritable knee in children is not uncommon in daily practice. The presence of effusion should alert the physician to a diagnosis of infective arthritis and consideration of arthrocentesis. Other differential diagnoses include reactive arthritis, transient synovitis, hemophilia, inflammatory arthritis, and malignancy, e.g., leukemia. The common causes of inflammatory arthritis include rheumatic fever, Reiters syndrome and vasculitis (e.g. Henoch-Schonlein Purpura) [2]. To the best of our knowledge, our patient is the first case in English literature in which PVNS presented as an acute irritable knee.

First described by Jaffa et al. in 1941, PVNS is a rare inflammatory granulomatous condition of unknown etiology [3]. It can classified into diffuse or localized/nodular type, with knee joint being the most commonly affected joint. The World Health Organization classification now defines localized PVNS under the group of giant cell tumor of the tendon sheath, which encompasses a group of lesion arising from the synovium of joints, bursae, and tendon sheath. While diffuse PVNS with diffuse intra-articular involvement is under the group of diffuse-type giant cell tumor [4].

PVNS has estimated the incidence of 1.8 per million [5]. It is even rarer in children and has only been presented as a case report or small case series [6-10]. Among the reported cases, there was a considerable delay in diagnosis and subsequent surgery. The delay ranged from a few months to more than 1-year [6]. The typical presentation was that of a child presenting with chronic insidious onset of pain, swelling, and stiffness in the involved joint; while the local PVNS more commonly presented with locking, catching, and instability [11-16].

The clinical presentation of our case was atypical. The patient presented with acute irritable and swollen knee, which clinically was highly suggestive of infective arthritis. In our practice, MRI is not readily available in such an acute setting. Under such clinical scenario, emergency arthroscopy is a justified choice for early diagnosis and treatment of the suspected infection, as a delay in treatment of septic arthritis can lead to the long-term morbidity of the patient. As a result, emergency arthroscopy was arranged, ultimately revealing the causative pathology. Fortunately, a complete excision of the lesion was achieved which lead to complete recovery.

To the best of our knowledge, this is the first reported case of PVNS mimicking infective arthritis in children. Only one similar case has been reported in an adult, in which a pregnant patient presented with acute knee arthritis mimicking infective arthritis [17]. Microscopic evaluation revealed hemorrhage and necrosis, which may suggest torsion or bleeding of the tumor as the cause of the acute symptom.

Area of necrosis was also noted in the lesion in our case. Since the stalk of the lesion was quite narrow, this might suggest acute torsion or strangulation of the lesion as the cause of the acute severe knee pain. The lesion was noted to be quite vascular with contact bleeding during arthroscopic manipulation. Together with the presence of hemarthrosis, this might suggest an episode of spontaneous hemorrhage from the lesion into the joint leading to acute massive hemarthrosis, which can mimic clinically an infective arthritis.

![Figure 1: Radiographs of the patient’s left knee showed soft tissue swelling without bony lesion.](image1)

![Figure 2: (a) Arthroscopic view showed a multi-lobulated vascular mass at the intercondylar notch, (b) resection of the mass.](image2)

![Figure 3: Histologic pictures of the lesion, (a) Low power field (×100) showing multinucleate giant cells, (b) high power field (×200) showing multinucleate giant cells, (c) hemosiderin deposits in macrophages, (d) area of necrosis at the left side of the picture](image3)

![Figure 4: Post-operative magnetic resonance imaging showed no residual lesion.](image4)
The treatment of pediatric PNVS of the knee follows that of the adult counterparts. For the local/nodular form of PVNS in adult, it is well-reported in many studies that local excision can lead to excellent or very good result with complete and persistent regression of the pathology [18-21]. Although no comprehensive data are available for the pediatric group in view of the small case load, the similar result has been obtained from the available literature [6-9].

Concerning the diffuse type of PVNS, studies from the adult age group had shown a higher risk of recurrence and morbidity [19,22]. Subtotal or total synovectomy, either arthroscopic or open, can reduce the risk of recurrence [19,23,24]. In contrary to adult, the role of radiotherapy as an adjunctive therapy is not recommended in children in view of potential damage of epiphyseal growth plate and post-radiation sarcomas [25,26].

Conclusion

PVNS of the knee in children is a rare entity. It can be one of the differentials in acute irritable knee in children. Prompt and proper treatment can lead to an excellent clinical result.

Clinical Messege

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