Pseudogout: A Rare Cause of Acute Carpal Tunnel Syndrome and Acute Guyon Canal Syndrome

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Case Report

Pseudogout is an acute inflammatory arthropathy that often presents as a hot, swollen, painful joint. Rarely, the inflammatory response caused by pseudogout has led to acute neuropathic symptoms of the hand. We present a case of pseudogout causing acute neuropathic symptoms in the median and ulnar nerves, ultimately necessitating urgent surgical decompression. The patient’s symptoms were alleviated after surgical decompression. Pseudogout should be considered a rare cause of acute neuropathic compression of the hand. Its management may require surgical intervention and should involve postoperative follow-up with a rheumatologist.

Crystal arthropathies affecting the hand often present as chronic pathology with intermittent flares of pain and inflammation. Few case studies have shown pseudogout flares associated with neuropathic symptoms, specifically with compression of the median nerve, resulting in acute carpal tunnel syndrome (CTS). Acute deposition of hydroxyapatite crystals has also been implicated in acute CTS, along with tumoral calcium pyrophosphate dihydrate (CPPD) causing compression of the median and ulnar nerves. Here, we present an uncommon case of new onset of pseudogout, presenting in the hand as both acute carpal tunnel syndrome and acute Guyon canal syndrome. The patient provided written informed consent for the use of deidentified health information and images for research and educational purposes.

Case

A 55-year-old right-handed woman with no history of rheumatologic disease presented to the emergency department with 2 days of increased pain, swelling, and progressive numbness in the right hand and all 5 fingers. She denied any trauma, injections, or penetrating injuries to her hand. She had chills and night sweats, but no fever. She denied any similar events in the past. Her medical history was notable for hepatitis C; distant intravenous drug use (heroin, last used 25 years before); and osteoarthritis of the knees, hips, and spine. She denied a personal history of gout, pseudogout, autoimmune disorders, or kidney stones. She reported a family history consistent with rheumatoid arthritis in her paternal grandmother and paternal great aunt.

On physical examination of her hand, there was erythema and swelling most prominent in the hypothenar region, extending proximally into the volar wrist. She was diffusely tender to palpation in the distal volar forearm, wrist, and hand. The volar wrist, along with the thenar and hypothenar compartments, was profoundly taut. She had numbness and paresthesia in the median and ulnar nerve distributions with diminished 2-point discrimination and normal sensation in the radial nerve distribution. Her motion was severely limited because of pain; however, she could weakly fire all intrinsic and extrinsic hand muscles. She had a palpable radial pulse, and her hand was warm and well-perfused.

The patient was afebrile and hemodynamically stable. The laboratory results revealed white blood cell count of 9,300 cells per cubic millimeter (reference, 4,300–10,800 per cubic
There was a considerable hourglass shape down to the transverse carpal ligament. Upon release of the volar carpal ligament, there was an immediate release of pressure on the volar wrist, thenar, and hypothenar compartments, and tension-free primary closure was performed. Rheumatology and infectious disease departments were consulted after surgery because of concern for crystal arthropathy versus possible atypical mycobacterial infection. The patient was discharged on sulfamethoxazole-trimethoprim while the results of the intraoperative cultures were pending. The final tissue pathology results showed dystrophic calcification suggestive of pseudogout (Fig. 2). Aerobic, anaerobic, and mycobacterial cultures were negative. The patient was seen at her 10-day follow-up and was noted to have complete resolution of pain with improvement in her range of motion and paresthesia. She plans to follow-up at the rheumatology department as an outpatient.

Discussion

Pseudogout, which describes the acute deposition of CPPD crystals into articular cartilage, triggers a proinflammatory immune response involving articular phagocytic cells and a resulting cytokine storm, propagating leukocyte and mononuclear cell migration. This results in a warm, painful, swollen joint that can mimic other musculoskeletal pathologies, including septic arthritis, acute calcific tendinitis, gout, rheumatoid arthritis, or primary osteoarthritis.

The first case of acute CTS attributed to pseudogout was reported in 1976 by Spiegel et al. Since then, there have been 4 other case reports in the published literature. The average age of patients in these reports was 69.8 years; 4 women and 1 man were included. Notably, the range of symptom onset before presentation for most patients was 3–10 days. Interestingly, our patient was relatively young and had a rapid onset of symptoms over 2 days. Most remarkably, she had signs and symptoms of acute ulnar and median nerve compression. Acute compression of the median and ulnar nerves is exceedingly rare, usually attributed to trauma or burns. Yamazaki et al. presented a case of median and ulnar nerve compression caused by tumoral CPPD disease, with both nerves affected by a 6 × 4-cm cystic mass of the volar wrist that had developed over the course of 1 year. Our patient did not have tumoral CPPD disease; however, she had considerable acute median and ulnar neuropathy. Previous reports of pseudogout causing acute CTS have suggested that CPPD deposits cause direct compression of the median nerve within the carpal tunnel. Chiu et al. reported direct compression of the median nerve by an outpocketing of the anterior capsule of the radiocarpal joint filled with a chalky, whitish substance. Goodwin et al. found “caseous-like deposits” under the median nerve. Interestingly, no CPPD deposits or gelatinous material were identified in the carpal tunnel of our patient. We hypothesize that the CPPD material found intraoperatively at the radial side of the Guyon canal caused direct compression of the ulnar nerve, with the resultant inflammatory response causing an acute compartment syndrome with concomitant acute median nerve compression.

There is 1 case report of acute calcific tendinitis of the flexor carpi ulnaris tendon causing acute compression of the ulnar nerve. Notably, the patient described in this case report had a...
presentation similar to ours, with the tenseness of the volar wrist compartment and ulnar paresthesia that progressed to involve the median nerve as well. Intraoperative findings within Guyon canal revealed a mass of soft, chalky material associated with the tendon of the flexor carpi ulnaris was extracted piecemeal. However, the pathologic analysis was more consistent with acute calcific tendinitis of the flexor carpi ulnaris and not crystalline pseudogout. Pathologic analysis of our sample revealed calcium phosphate deposits within the fibrous tissue, with no clear pattern of deposition around vessels or other structures. Interestingly, the deposits were not associated with accompanying inflammation or areas of necrobiosis. According to the pathologist, this was likely because of the lack of adjacent tissue included in the sample and inflammation and fibrosis in the adjacent affected tissue may be present. There was little to no birefringence under polarized light microscopic examination. Although the preservation of polarization can be accomplished by alcohol fixation or nonaqueous methods of fixation, the specimen here was processed routinely with formalin-fixation and staining with hematoxylin-eosin, leading to the absence of appreciable polarization.11 Calcific tendinitis can be difficult to differentiate from pseudogout clinically. Calcific tendinitis is caused by synovial deposition of basic calcium phosphate (hydroxypatite), which is associated with fibrous and papillary thickening of the synovium. In joint fluids, the aggregated hydroxypatite crystals often form irregularly shaped microspheroids. Pathological analysis often reveals a pale, laminated psammomatous appearance on decalcified tissue sections. Calcium pyrophosphate dihydrate typically appears dark blue to purple with a distinctly granular appearance on nondecalcified hematoxylin-eosin—stained slides. Decalcified pyrophosphate leaves oval to rounded pools of basophilic material that results in a fibrillar or feathery appearance with sharp and pointed edges, not typically seen in the rounded, psammomatous calcifications in calcific tendinitis.12 The pathologic findings, in conjunction with the overall histologic and gross findings, were most consistent with acute pseudogout as the primary explanation for the patient’s symptomatology.

Surgical decompression and removal of the material promptly alleviated the patient’s pain and neuropathy symptoms, similar to the findings of previous reports.2–4 With regard to further management of patients with pseudogout, current recommendations include starting a nonsteroidal anti-inflammatory drug (naproxen) at 500 mg twice daily for 5–7 days in patients that present with >24 hours of symptoms. Colchicine dosed at 0.6 mg is recommended for patients who present within the first 24 hours of symptoms onset, with no more than 1.8 mg taken within the first 24 hours. This is followed by 0.6 mg taken twice daily until the attack resolves. Other treatment options for refractory cases include steroid injections and oral glucocorticoids; however, care must be taken to rule out the presence of infection before initiating these treatments.13 Our patient was started on intravenous antibiotics before intraoperative culture collection due to concern for a possible infection; she was discharged on 10 days of oral sulfamethoxazole-trimethoprim. Although the culture results were negative, a previous study has shown that patients who receive preoperative antibiotics before the debridement of upper extremity infections are significantly more likely to have negative cultures at the time of debridement, with even 1 dose of antibiotics reaching significance (P < .01).14 Clinicians should be wary of possible concomitant infection with pseudogout given the similar presentations, along with the increase in negative cultures that can occur with the preoperative initiation of antibiotics.

The acute inflammatory response caused by pseudogout can present as a challenging diagnostic scenario and has been implicated in acute nerve compression necessitating surgical intervention. This is a unique case of nontumoral pseudogout causing acute compartment syndrome with neuropathic symptoms of the median and ulnar nerves. Pseudogout should be included in the differential diagnosis of a patient presenting with atraumatic, acutely neuropathic symptoms associated with swelling and pain that do not appear to have an infectious etiology.

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Figure 2. A 5-μm hematoxylin-eosin—stained section of formalin-fixed, paraffin-embedded tissue with no decalcification. White arrows indicate dense fibrous and fibrovascular tissue with coarse basophilic deposits of calcium phosphate. B A 5-μm hematoxylin-eosin—stained section of formalin-fixed, paraffin-embedded tissue with no decalcification. White arrows indicate calcium phosphate deposits within the fibrous tissue, with no clear pattern of deposition around vessels or other structures (Hematoxylin-eosin stain; magnification: A × 100, B × 200).
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