Multilevel calcium pyrophosphate dihydrate deposition in cervical ligamentum flavum: clinical characteristics and imaging features

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Abstract
Background: Involvement in cervical ligamentum flavum is a rare manifestation of the calcium pyrophosphate dihydrate deposition disease. Only few cases of this condition have been reported. We revealed eighteen cases of CPPD in cervical ligamentum flavum that diagnosed at a single medical center. In our case series, clinical characteristics and magnetic resonance imaging findings of patients are described.

Methods: We retrospectively reviewed the medical charts and imaging studies of the eighteen patients with pseudogout attack of the cervical ligamentum flavum. In addition, we discussed the differences between this disease and ossification of ligamentum flavum in image manifestations.

Results: There were fourteen men and four women aged between 59 and 87 years. Diabetes mellitus and hypertension were the most common comorbidities. Myelopathy and neck pain were presented in most patients. C4–5 and C5–6 were attacked most frequently, and multiple- rather than single-level involvement could be observed in our series. “Acute on chronic phenomenon” was a specific magnetic resonance image finding in patients whose symptom durations were between 2 to 5 months. Compared to ossification of ligamentum flavum, calcium pyrophosphate dihydrate crystal deposition had different image signs, including morphology, side of the involved ligament, no continuity with the lamina, acute on chronic phenomenon, and presence of retro-odontoid mass.

Conclusions: Nodular calcifications in cervical ligamentum flavum raise highly suspicion for calcium pyrophosphate dihydrate deposition and must be diagnosed by histological examination and polarized light microscopy. This disease is different from ossification of ligamentum flavum, and it could be recognized by specific image features.

Keywords: Calcium pyrophosphate dihydrate, Cervical ligamentum flavum, Magnetic resonance imaging, Ossification of ligamentum flavum

Introduction
Calcium pyrophosphate dihydrate (CPPD) crystal deposition disease, also known as pseudogout, was first described by Zinan and Sitaj in 1958 [1]. It is thought to be a disease of aging and patients younger than 50 years of age are less commonly affected [2]. The pathogenesis of CPPD deposition disease is still unknown, but the disease can be associated with some metabolic or endocrine diseases, including diabetes mellitus, hypertension, hyperparathyroidism, hypothyroidism, hypophosphatasia, hemochromatosis, gout, and rheumatoid arthritis [3]. Idiopathic CPPD deposition disease is the most common form. CPPD crystals deposit commonly at joints of extremities and sometimes at the thoracolumbar spine.
However, the cases with CPPD in cervical spine is a rare entity, and patients were primarily found in Japan and North Africa [4]. The attack of crystals in intervertebral discs, articular cartilages, joint capsules, and ligaments could result in spinal lesions. However, most patients with calcifications of the intervertebral discs and articular cartilages are asymptomatic [5]. Conventional radiography and sonography are main imaging techniques for the diagnosis of peripheral CPPD and the computed tomography is commonly used to detect axial CPPD [6, 7]. In the presented case series, we report 18 patients with calcification of cervical ligamentum flavum who required surgical treatment and finally were confirmed to suffer from CPPD deposition disease by pathologic diagnosis. Compared with previous reports which discussed the findings of pathophysiology, laboratory tests, clinical manifestation, computed tomography (CT) scan, and various pathological studies, our report showed magnetic resonance imaging (MRI) findings and other clinical characteristics. In addition, CPPD deposition disease in the ligamentum flavum is frequently mistaken for ossification of ligamentum flavum which is commonly found in the lower thoracic spine and possibly identified in the cervical spine [8]. To date, there is no study to examine the difference in imaging features between the two diseases. Therefore, we also describe the comparison of imaging findings in this study.

**Materials and methods**

From January 2014 to January 2020, 18 patients with CPPD deposition in cervical ligamentum flavum were retrospectively analyzed and formed the basis of this study. All these cases received surgical treatment and were diagnosed by histological examination at our hospital, a teaching hospital which provides tertiary patient care. In these subjects, only two patients had a previous diagnosis of CPPD and the knee joint was involved. In all the other patients, cervical spine involvement was the first presentation of the disease. Table 1 showed their basic clinical data, including age, gender, comorbidities,

| Case | Age/Sex | Comorbidities | Symptoms | Duration | Type of surgery |
|------|---------|---------------|----------|----------|----------------|
| 1    | 70/M    | DM            | Neck pain Myelopathy | 2 months | Decompression with fusion |
| 2    | 85/M    | Absent        | Myeloradiculopathy    | 10 months| Decompression    |
| 3    | 59/M    | HTN, Arrhythmia| Myeloradiculopathy    | 3 months | Decompression    |
| 4    | 80/M    | DM, HTN       | Myelopathy            | 4 months | Decompression with fusion |
| 5    | 66/M    | HTN, hyperuricemia| Neck pain Myeloradiculopathy| 10 months| Decompression with fusion |
| 6    | 72/M    | DM, HTN       | Neck pain Radiculopathy | 3 months | Decompression    |
| 7    | 86/M    | HTN, COPD     | Neck pain Myeloradiculopathy | 18 months| Decompression    |
| 8    | 79/M    | Absent        | Myelopathy            | 2 months | Decompression    |
| 9    | 87/M    | HTN, COPD     | Myeloradiculopathy    | 2 months | Decompression    |
| 10   | 73/M    | DM, HTN       | Neck pain Myeloradiculopathy | 4 months | Decompression    |
| 11   | 78/F    | HTN CAD       | Myelopathy            | 18 months| Decompression    |
| 12   | 70/F    | DM, CAD       | Neck pain Myeloradiculopathy | 24 months| Decompression    |
| 13   | 77/M    | HTN, ESRD     | Neck pain Myeloradiculopathy | 24 months| Decompression    |
| 14   | 81/M    | HTN           | Neck pain Myelopathy  | 9 months | Decompression    |
| 15   | 67/F    | ESRD          | Neck pain Myelopathy  | 15 months| Decompression with fusion |
| 16   | 81/M    | DM            | Myeloradiculopathy    | 9 months | Decompression    |
| 17   | 78/F    | DM, CAD       | Neck pain Myeloradiculopathy | 5 months| Decompression with fusion |
| 18   | 77/M    | HTN           | Neck pain Radiculopathy | 12 months| Decompression    |

*DM Male, F Female, DM Diabetes mellitus, HTN Hypertension, COPD Chronic obstructive pulmonary disease, CAD Coronary arterial disease, ESRD End stage renal disease*
symptoms, symptom duration, and types of surgery. The data of blood examinations were reviewed if data was available. Anteroposterior and lateral radiographs were obtained in all patients. Computed tomography (CT) scans were available in five patients. MRIs (1.5-T machine, Magnetom Vision, Siemens Medical Systems, Iselin, NJ; Signa, GE Medical Systems, Milwaukee, WI) were performed in all patients preoperatively to evaluate spinal cord compression and confirm abnormal soft tissue structure. We recorded the signal intensity of the lesions in the ligamentum flavum on T1- and T2-weighted images as low, intermediate, high-signal intensity, as compared with that of the nearby bone, disc, or cerebrospinal fluid. Mixed signal intensity meant that the lesion showed both hyperintensity and hypointensity in the ligamentum flavum. The morphology, affected levels and presence of retro-odontoid mass were also recorded.

The patients included in this study were treated initially by pharmacological treatments, such as acetaminophen and non-steroidal anti-inflammatory drugs. All patients received surgical treatment due to failed conservative treatment. The surgical options were consisted of posterior decompressive surgery alone and posterior decompression with posterolateral fusion. In addition, calcified lesions in ligamentum flavum were removed entirely and every surgical specimen was checked by histologic examination and polarized light microscopy. Postoperative results were evaluated according to Frankel scale [9].

Results
The series included fourteen males and four females (cases 11, 12, 15, and 17), and the mean age was 75.9 years (range: 59 to 87 years). The mean age of the women was 73.2 years (range, 67 to 78 years), while the mean age of the men was 76.6 years (range, 59 to 87 years). The mean symptom duration was 9.7 months (range: 2 to 24 months). Comorbidities were present in sixteen patients and consisted of hypertension (n = 10), diabetes mellitus (n = 7), coronary artery disease (n = 3), end stage renal disease (n = 2), chronic obstructive pulmonary disease (n = 2), arrhythmia (n = 1), and hyperuricemia (n = 1). Besides, the patients presented with myelopathy (n = 6), radiculopathy (n = 2), myeloradiculopathy (n = 10) and neck pain (n = 11). In available data of blood examinations, there was no significant difference in the level of serum uric acid and electrolytes. None of the eighteen patients did the plain radiographs of the cervical spine that contributed to the final diagnosis. CT scans that performed in five patients (cases 1, 5, 7, 13, and 18) revealed oval-shaped calcifications located ventrally to the lamina (Fig. 1). These lesions had clear margins and had no connection with the adjacent lamina. They were separated by ligamentum flavum which showed relative low density compared with that of calcified lesions and lamina in the CT images. In addition, the lesions presented bilaterally and paramedially along the midline without extending to posterior facet joint.

MRIs were performed in all patients and the results were listed in Table 2. Nodular lesions were presented in all cases. The mid-cervical spine was often affected, especially at C4–5 (9 in 18 patients, 50%) and C5–6 (10 in 18 patients, 56%) level. C1–2 and C7–T1 were least affected level. Both were just found once among patients. Multilevel involvement was more common than single level involvement, fifteen patients (83%) and three patients (17%) respectively. Furthermore, nine patients (50%) had

![Fig. 1 Cervical computed tomography of case 5. Compression of the spinal cord by two main calcified masses was observed on sagittal and axial sections. There is no calcification or ossification of the posterior longitudinal ligament](image-url)
two levels of involvement, four patients (22%) had three levels of involvement, and two patients (11%) had four levels of involvement. In eight patients (cases 1, 3, 4, 6, 8, 9, 10, and 17), the nodular lesion showed homogenous low-signal intensity and the signal was even lower than cerebrospinal fluid in T1-weighted image. Surrounding each nodular lesion, intermediated-signal change was noted (Fig. 2A, D, Fig. 3 A, and D). In T2-weighted image, the nodular lesions still showed homogenous low-signal intensity, surrounded by intermediated- to high-signal intensity (Fig. 2B, E, Fig. 3 B, and E). The low-signal intensity on both T1- and T2-weighted image referred to a calcified lesion. Around the lesions, the intermediated-signal change on T1-weighted and intermediated- to high-signal change referred to the edematous reaction. We described this condition as “acute on chronic phenomenon” and it might present an acute or subacute inflammatory reaction. The further short tau inversion recovery (STIR) sequence in MRI was used to demonstrate this edematous change (Fig. 2C). In these eight patients with “acute on chronic phenomenon”, their mean duration of symptoms was 3.1 months (range: 2 to 5 months) (Table 1). For patients whose duration of symptoms were longer (10 patients, mean duration: 14.9 months, range: 9 to 24 months), lesions just showed homogenous low-signal intensity on both T1- and T2-weighted images without “acute on chronic phenomenon”. Retro-odontoid lesions were found on thirteen patients (72%). Five patients (case 2, 4, 10, 11, and 13) had no obvious calcification surrounding the peri-odontoid process.

Thirteen patients received posterior decompressive laminectomy alone and five patients (cases 1, 4, 5, 15, and 17) received posterior laminectomy with posterolateral fusion and instrumentation by lateral mass screws. The gross appearance of the calcified ligamentum flavum from a patient who underwent laminectomy with posterolateral fusion was shown in Fig. 4A. Abundant calcified deposits within ligamentum flavum were observed intraoperatively and did not adhere to the dura mater. Nevertheless, these crystals lead to spinal cord compression.

### Table 2 MRIs findings of the lesions in cervical ligamentum flavum

| Case | Morphology | Affected levels | Acute on chronic phenomenon | Retro-odontoid mass |
|------|------------|-----------------|----------------------------|---------------------|
| 1    | Nodular    | C3–4, C4–5, C5–6, C6–7 | Present                    | Present             |
| 2    | Nodular    | C5–6, C6–7       | Absent                     | Absent              |
| 3    | Nodular    | C5–6             | Present                    | Present             |
| 4    | Nodular    | C3–4, C7–T1      | Present                    | Absent              |
| 5    | Nodular    | C3–4, C4–5       | Absent                     | Present             |
| 6    | Nodular    | C2–3, C6–7       | Present                    | Absent              |
| 7    | Nodular    | C1–2, C2–3, C3–4 | Absent                     | Present             |
| 8    | Nodular    | C2–3, C3–4, C4–5 | Present                    | Present             |
| 9    | Nodular    | C3–4, C4–5       | Present                    | Absent              |
| 10   | Nodular    | C3–4, C4–5, C5–6, C6–7 | Present                | Absent              |
| 11   | Nodular    | C4–5, C5–6       | Absent                     | Absent              |
| 12   | Nodular    | C4–5, C5–6, C6–7 | Absent                     | Present             |
| 13   | Nodular    | C2–3             | Absent                     | Absent              |
| 14   | Nodular    | C2–3, C3–4       | Absent                     | Present             |
| 15   | Nodular    | C5–6, C6–7       | Absent                     | Present             |
| 16   | Nodular    | C4–5, C5–6, C6–7 | Absent                     | Present             |
| 17   | Nodular    | C4–5, C5–6       | Present                    | Present             |
| 18   | Nodular    | C5–6             | Absent                     | Present             |

**Fig. 2** Magnetic resonance imaging of case 1. Nodular masses with low-signal intensity in both T1-weighted imaging (A and D) and T2-weighted imaging (B and E) were observed at the interlaminar space of C4–5, C5–6, and C6–7. These lesions were surrounded by areas of intermediate-signal intensity area on T1-weighted imaging and by areas of high-signal intensity area in T2-weighted imaging, and the short tau inversion recovery (STIR) sequence (C) was used to demonstrate the edematous change.
The nodule was composed of fine granules and chalky white in color. Histopathologic examination of the specimen revealed a prominent hyaline degeneration and calcification in the fragment of ligamentum flavum. Endothelial cell, fibroblast, and giant cells were observed in the granulation tissue. There was mild inflammatory change in the selective cases, including cases 1, 3, 4, 6, 8, 9, 10, and 17. No osteoid proliferation or bone formation could be observed. The hematoxylin and eosin stain revealed a basophilic appearance (Fig. 4B). CPPD deposition with rod-shaped crystals was demonstrated under the polarized microscope (Fig. 4C).

Discussion
CPPD deposition in the ligamentum flavum at the cervical spine is extremely rare. According to the English-language literatures, most of case series about the disease were from Japan [10–13]. Nevertheless, the prevalence of this condition in Blacks is still under-recognized. Two patients were confirmed as the diagnosis of CPPD deposition in six Black cases with calcification of cervical ligamentum flavum in the Caribbean island [14]. In this study, we identified 18 cases at a single medical center in recent 8 years, raising the possibility that the incidence of this condition could be definitely underestimated in East Asia. Similar to CPPD deposition in cervical ligamentum flavum, ossification of the ligamentum flavum (OLF) in the cervical spine is also
rare and it is generally understood that they are different diseases histologically.

Kenneth H. Fye [3] declared that CPPD deposition in cervical spine was a disorder of elderly women with tendency of calcium deposition. In their report, 80% of patients were women, but our series showed a predominance in men (14 in 18 patients, 78%). This could be explained by the fact that the mean age of male population is older than that of female population (76.6 and 73.2 years, respectively). Moreover, Miyazawa et al. [15] reported the mean age of the presented patients with OLF in the cervical spine was 56.3 years. However, our study found a mean age of 76 years, suggesting that CPPD deposition in the cervical spine has a later onset than that of the OLF in the cervical spine. In addition to aging and gender, metabolic disorders have been shown to be associated with calcium deposition as well. Diabetes mellitus (incidence from 40 to 60%), hypertension (incidence from 25 to 40%), and atherosclerosis (incidence from 35 to 50%) were most frequently encountered [16] and these comorbidities were consistent with our findings. In various symptoms of CPPD deposition in the cervical spine, cervical myelopathy was the most prevalent one, and it frequently resulted in paresthesia in the upper limbs, deterioration in fine motor skills, and difficulty in walking. A noticeable finding in our case series was that neck pain is a common complaint among patients (11 of 18 patients). Literature has shown that acute neck pain could be caused by CPPD crystal-induced arthritis of the lateral atlantoaxial joint [17]. In addition, Kobayashi et al. [18] reported a case with acute neck pain that resulted from CPPD attack of the cervical ligamentum flavum without facet arthritis. Although we can not conclude that neck pain in our case series is due to CPPD deposition in ligamentum flavum, acute neck pain should be considered a symptom for the diagnosis of CPPD deposition disease.

Markiewitz et al. [19] displayed that 33.8% of patients with CPPD deposition disease had axial involvement, mostly affecting the lumbosacral spine (87%). The cervical spine was less commonly involved. The midcervical spine was most frequently attacked among cervical spine. Hisatoshi et al. [11] reported a case series with 8 patients suffered from CPPD deposition disease, and 81% of the patients’ lesions were located between C4–5 and C6–7 levels. This finding was compatible with our data that the most commonly affected levels were C4–5 and C5–6. C5 segment has the greatest dynamic stress in the cervical spine and the elastic tissue is prone to undergo degeneration [20]. The ligamentum flavum is thinner in the cervical region than that in the thoracic or lumbar region. However, the elastic tissue is more abundant in the cervical spine. Yoshida et al. [21] and S-H Lin et al. [22] reported three cases with CPPD deposition in multilevel of ligamentum flavum. However, most patients (83%, 15 in 18 patients) had at least involvement of two intervertebral levels in our series. We believed that there is no tendency for CPPD deposition to affect just single level. Furthermore, OLF was commonly identified in the thoracic spine or the thoracolumbar junction, and cervical spine was a rare target for OLF [23]. However, when OLF occurs in the cervical spine, there is no common location and single-level involvement is predominant [15].

In order to clearly detect the cervical myelopathy, spinal cord compression, other soft-tissue changes and fluid effusion, we performed MRI for all patients. Sushil et al. [24] reported three diagnostic features of CPPD deposition disease in MR images, including: (1) mostly isointense lesion in the T1-weighted image, (2) a mixed density mass in T2-weighted image, and (3) a peripheral enhancement on the postcontrast MRI images. In term of morphology in sagittal imaging, all lesions had the nodular-type appearance, rather than beat-like or mound-like appearance that commonly observed in cases of ossification [25]. Besides, in T1-weighted image, the nodular lesion showed low-signal intensity, and the signal intensity was even lower than that of the cerebrospinal fluid. In T2-weighted image, the nodular lesion showed low-signal intensity as well. In addition, these lesions were surrounded by areas of intermediate signal intensity in T1-weighted imaging and by areas of high signal intensity in T2-weighted imaging. The different signal intensity in the surrounding areas of lesions was considered an appearance of the edematous reaction. We described this edematous change as “acute on chronic phenomenon”. It meant that there was acute or subacute inflammatory reaction induced by crystal deposition. In this report, the “acute on chronic phenomenon” was found in eight patients whose duration of symptoms was less than 5 months. This unique imaging finding was different from that of the OLF. The images of OLF showed low-signal intensity without edematous change on both T1- and T2-weighted image. Other ten patients whose duration of symptoms more than 9 months did not have the imaging finding of “acute on chronic phenomenon”. However, we could not demonstrate a clear cut-off point for duration of symptoms to determine whether this phenomenon would be present or not.

CT scans were performed in five patients. The calcifications were seen as oval, high attenuation, and symmetric space-occupying lesions ventral to the lamina. They were located on either side of the midline and not observed in the capsular part. This condition could explain why most of patients presented with myelopathy instead of radiculopathy alone. For cervical spine calcifications, CT-scan frequently revealed characteristic “calcified railroad “pattern in the transverse retro-odontoid ligament, fine and
mottled calcifications around the odontoid process, and intervertebral disc calcifications [26]. Unlike calcification, OLF would extent to the posterior facet joint because the initiation of the ossification starts at the joint capsule and the ligament was ossified unilaterally in some cases [27]. CPPD crystal deposits also result in enlarged retro-odontoid masses which should present with calcifications surrounding the apex of the dens. This condition is known as “crowned dens syndrome” [28] when patients present typical clinical features, including acute onset of severe neck pain, neck stiffness (particularly in rotation), and positive inflammatory indicators (fever, increased C-reactive protein level, or elevated white blood-cell count). However, calcification around the dens does not absolutely cause crowned dens syndrome. In addition, tumor and calcified retropharyngeal tendinitis may also have calcium deposition around the dens. Crown dens syndrome should be considered not only in CPPD disease but also in various clinical situations [29]. Salaffi et al. [30] reported that 51% of patients had periodontoid calcifications in known CPPD deposition in cervical ligamentum flavum. Haikal et al. [31] reported that 60% of patients had crowned dens syndrome compatible with their cervical computed tomography findings. They declared the importance of performing cervical computed tomography when evaluating patients with CPPD deposition could not be overemphasized. The incidence of the disease in our study was higher. By contrast, there is no typical retro-odontoid calcified lesion noted in patients with ossification of ligamentum flavum. A comparison between the CPPD deposition disease and ossification of the ligamentum flavum is summarized in Table 3 according to our study and reviewed literatures [10, 15, 23, 25, 27, 32].

The natural history of cervical myelopathy that resulted from CPPD deposition in the ligamentum flavum is not well-known. Prognostic factors include age, comorbidities, rate of deterioration, severity of symptoms, duration of symptoms, and signal change within the spinal cord [33]. It is generally agreed that surgical decompression is necessary for patients with ongoing and progressive symptoms refractory to conservative treatment [34]. However, for long-term effect of surgery, there is no significant difference between operative and non-operative group in a Cochrane Review of randomized controlled trials reported by Fouyas IP et al. [35] In patients with “acute on chronic phenomenon” in MR image, two to three grades of neurologic improvement according to Frankle scale were noted in our series. Two patients (cases 12 and 13) that suffered from severe and prolonged spinal cord compression with persisted and obvious signal change in MR image had shown one grade of improvement after treatment. The purpose of surgery was to prevent further decline in function in two patients. The options of surgical treatment in our case series include posterior decompression alone or decompressive surgery with posterolateral fusion. The long-term follow-up study showed that 91% of patients receiving laminectomy alone were satisfied with the outcomes and had significant improved functional status [36], but fusion with instrumentation still had its role in patients with significant neck pain, cervical lordosis less than 10 degrees, younger age, postoperative radiation, or the presence of instability [37]. For patients with neck pain, we did not routinely performed fusion surgery because neck pain could not always be considered as a predictor of late instability in elderly patients. We performed additional fusion surgery in a patient that demonstrated CPPD deposition in ligamentum flavum of C3–4 and C7-T1 to prevent instability in cervicothoracic junctional level.

This case series highlights important clinical issues as below. First, CPPD attacks of the cervical ligamentum flavum can result in neck pain, radiculopathy, myelopathy, or myeloradiculopathy. Second, nodular calcification of the ligamentum flavum in cervical spine raises high suspicion of CPPD deposition disease. Third, CPPD

| Table 3 | The distinctive imaging findings of CPPD deposition in ligamentum flavum and ossification of ligamentum flavum (OLF) in the cervical spine |
|------------------|------------------|------------------|------------------|
| **CPPD deposition** | **OLF** | **CPPD deposition** | **OLF** |
| 1. Morphology | Nodular | Beaklike or moundlike |  |
| 2. Location | C4–5 and C5–6 | No common location |  |
| 3. Involvement of level | Multilevel predominant | Single level predominant |  |
| 4. Side of the involved ligament | Always bilateral | Unilateral or bilateral |  |
| 5. Extend to the posterior facet joints | Absent | Always present |  |
| 6. Continuity with the lamina | No continuity | Continuity with lamina |  |
| 7. Acute on chronic phenomenon | Depend on symptoms duration | Not reported |  |
| 8. Retro-odontoid lesion | May present | Not reported |  |

CPPD Calcium pyrophosphate dihydrate
deposition can present with single or multilevel involvement. Fourth, CPPD deposition disease and ossification of ligamentum flavum are the different disease and have their own imaging features. We believe that the cases that suffered from CPPD deposition disease in the cervical ligamentum flavum is underreported because specimens are not routinely stained in Hematoxylin and Eosin staining or inspected with polarized-light microscopy. If CPPD deposition is suspected, the specimen should be fixed in 70% ethanol and separated from other sectioning bone or cartilaginous specimens [38]. Most importantly, the pathologist should be informed of the suspicion so that the specimen can be handled appropriately.

Conclusions
On the basis of our study and literatures in these cases and the review of the literature, the imaging features and intraoperative findings of nodular calcifications in cervical ligamentum flavum raise highly suspicion for CPPD deposition. This disease is different from ossification of ligamentum flavum, and it can be recognized by specific image features. An accurate diagnosis can be established through histopathological examination of the specimen. Patients with neurological symptoms may generally require surgical decompression to improve the symptoms and prevent the progression.

Abbreviations
CPPD: Calcium pyrophosphate dihydrate; CT: computed tomography; MRI: magnetic resonance imaging; STIR: short tau inversion recovery; OLF: ossification of the ligamentum flavum.

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Authors’ contributions
Conception and design: Shih-Tien Wang, Chien-Lin Liu, Ming-Chau Chang. Provision of study materials or patients: Hsi-Hsien Lin, Po-Hsin Cho, Ming-Chau Chang. Collections and assembly of data: Yueh-Hsiu Lu, Hsi-Hsien Lin, Provisment of study materials or patients: Hsi-Hsien Lin, Po-Hsin Cho, Ming-Chau Chang. Manuscript writing: all authors. The authors read and approved the final manuscript.

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Availability of data and materials
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Declarations
Ethics approval and consent to participate
This study protocol was approved by Institutional Review Board Committee of Changhua Christian Hospital. The committee’s reference number was CCH IRB No.: 210601. All methods were performed in accordance with the Declaration of Helsinki. Informed consent was obtained from all participants.

Consent for publication
Not applicable.

Competing interests
The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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