Vertebra plana is a radiologic diagnosis that indicates complete compression of the vertebral body [1]. Although eosinophilic granuloma of the bone is the most frequent benign disease responsible for vertebra plana [2], it can be caused by several pathological conditions. Vertebral plana caused by a giant cell tumor (GCT) has been mentioned in only a few case reports in the English medical literature, and those case reports were without a detailed description [1]. To the best of our knowledge, the computed tomography (CT) and magnetic resonance (MR) imaging features of GCT presenting as vertebra plana have seldom been reported in the radiological literature. We report here on the case of

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We report here on the case of a 19-year-old woman who presented with progressive weakness of the lower extremities. The radiographs and CT showed vertebra plana of the first thoracic vertebral body. The mass had low signal intensity on the T1-weighted MR image and intermediate signal intensity on the T2-weighted MR image, and this low signal intensity extended to the spinal canal. Histological examination revealed a giant cell tumor (GCT). MR imaging is the imaging modality of choice for helping to distinguish spinal GCT from other spinal tumors by defining the extent and characteristic signal intensity of the tumor.

Index words: Spine
Tumor
Magnetic Resonance Imaging
Giant Cell Tumors

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a young woman with a GCT involving the thoracic spine and this presented as vertebra plana.

**Case Report**

A 19-year-old woman presented with a 10-day history of progressively worsening paraparesis. She also had a 1-month history of upper back pain that had been managed conservatively. The neurologic examination revealed bilateral hand grasp weakness and paraparesis. The results of the laboratory studies were within the normal limits. A swimmers lateral view radiograph of the cervical spine showed collapse of the first thoracic vertebra, and this was consistent with vertebral plana (Fig. 1). Contrast-enhanced CT showed marked collapse of the vertebra, which was surrounded by a well-enhanced paravertebral mass that extended to the spinal canal. Round bony erosion due to the soft tissue mass was seen in the anterior aspect of the second thoracic vertebral body (Figs. 2A–C).

The mass showed low signal intensity on the T1-weighted image (Fig. 3A), intermediate signal intensity on the T2-weighted image (Fig. 3B) and strong enhancement on the gadolinium-enhanced T1-weighted image (Fig. 3C), and the mass’s signal intensity was similar to that of the spinal cord. The presumptive preoperative diagnosis was eosinophilic granuloma and the alternative diagnosis was Ewing’s sarcoma based on the radiological findings of vertebra plana. The patient underwent corpectomy of C7 and T1 and partial corpectomy of T2. A large extraspinal, intraspinal extradural tumor was identified primarily anterior to the spinal cord at C7 and T1. The tumor was removed subtotally and spine arthrodesis was performed. One month later, a second

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**Fig. 2.** A 19-year-old woman with a spinal GCT. The precontrast (A) and postcontrast axial CT (B) shows a soft tissue mass (arrowheads) that has destroyed the first thoracic vertebral body with extension to the spinal canal (arrow), and this mass shows homogeneous enhancement (arrowheads) on the contrast-enhanced axial CT (B). The sagittal CT reconstruction (C) shows the paper-thin vertebral body (arrowheads), with preservation of the intervertebral disc space. Round bony erosion is seen in the anterior aspect of the T2 vertebral body (arrow).
operation was performed for total removal of the resid-
ual tumor. There has been no recurrence of neurologic
symptoms during 6 months of clinical follow up.

The pathological specimen (Fig. 4) of the tumor re-
vealed proliferation of two cell populations: stromal
cells and giant cells. The mononuclear stromal cells
were round to oval and they lacked cytologic atypia.
The giant cells were large and they contained 50–100
nuclei. Mitotic activity was frequent, while atypical mi-
totic figures were not found.

**Discussion**

GCTs are relatively common, locally aggressive bone
tumors, and they account for 4% to 9.5% of all primary
osseous neoplasms and 18% to 23% of all benign bone
neoplasms [3, 4]. The majority of GCTs occur at the
ends of long bones such as at the knee joint, and only
7% occur in the spine. GCTs in the spine most commonly
involve the sacrum, while the lumbar, thoracic and
cervical spine may be affected in order of decreasing fre-
quency [5].

In several large series, involvement of the thoracic
spine accounted for only 1% to 2% of GCTs [6, 7]. GCTs
of the spine develop in skeletally mature patients during
the second to fourth decades of life, and they are more
frequently seen in females (4, 8). A GCT of the spine can
present with pain (often with a radicular distribution),
weakness and sensory deficits [9, 10].

Like the GCTs in the appendicular bones, spinal GCTs
show no evidence of a mineralized matrix. A GCT in the
spine tends to affect the vertebral body rather than the
posterior elements of the spine. Radiologically, spinal

![Fig. 3. A 19-year-old woman with a spinal GCT. The collapsed first thoracic vertebral body is surrounded by a paravertebral mass with low signal intensity (arrowheads) and the spinal cord is compressed on the T1-weighted image (A), and the mass shows intermediate signal intensity on the T2-weighted image (B) and homogenous enhancement on the contrast-enhanced T1-weighted image (C). MRI shows that the mass involves the anterior aspect of the second thoracic vertebral body (arrows), and this corresponds to the erosive lesion seen on CT (Fig. 2C).](image-url)

![Fig. 4. The histological specimen shows that the tumor consists of mononuclear stromal cells evenly mixed with numerous osteoclast-like giant cells (arrow) (hematoxylin and eosin stain; original magnification, × 200). The mononuclear cells are round to oval or elongated and they lack cytologic atypia. The giant cells are large and they have over 20–30 nuclei.](image-url)
GCTs usually show as an expansile lesion with osteolysis [9]. On CT, a spinal GCT appears as a soft-tissue density mass with well-defined margins that may show a thin rim of sclerosis. It may also have a homogeneous hypervascular appearance with contrast enhancement [6].

This type of tumor usually has low to intermediate signal intensity on T1-weighted MRI. Gadolinium enhancement of the lesion on T1-weighted MRI reflects its vascular supply. Cystic areas, foci of hemorrhage, fluid-fluid levels and a peripheral low-signal-intensity pseudocapsule may also be seen [3]. In addition, GCTs may have signal intensity that is lower than or similar to that of the normal spinal cord on T2-weighted MRI in 63% to 96% of the cases [9]. This feature may be caused by the relative collagen content of the fibrous components and the hemosiderin within the tumor [6, 9].

Although most GCTs show expansile destruction of the vertebra, as was described in this case, a GCT can be seen as vertebra plana, which is defined by the following radiological criteria: collapse of one vertebral body only, normal adjacent intervertebral disks, the height of the intervertebral space is increased by at least one-third compared to normal and the collapsed vertebra shows increased density [1]. Vertebral plana can be seen in various other diseases, including eosinophilic granulomas, myofibromatosis, aneurysmal bone cysts, osteosarcomas, lymphomas and Ewing’s sarcomas [1, 2]. Vertebral plana could be produced by some disease processes that involve most of a vertebral body, and this is followed by spinal fractures. In our case, the initial working diagnosis was eosinophilic granuloma because it is the most common cause of vertebra plana in young patients. However, the radiologic features in our case, such as the presence of a paravertebral mass and involvement of the spinal canal and the adjacent vertebral bodies, were not typical of an eosinophilic granuloma. An eosinophilic granuloma is usually localized to one vertebra and it is contained by the periosteum, with no expansion to the paravertebral soft tissues [3]. A previous report has described the involvement of both the vertebral bodies and posterior osseous elements by eosinophilic granuloma [7].

Ewing’s sarcoma can also present as vertebra plana with extension to the posterior elements of the vertebra and it is one of the most common malignant tumors causing vertebra plana, which led us to include it in the differential diagnosis [1, 2].

Owing to the rarity of a GCT presenting as vertebra plana, it was not considered in the differential diagnosis. Yet in retrospect, the MRI findings suggested the characteristic findings of a GCT tumor of the spine, including the presence of intermediate signal intensity on the T2-weighted image that was similar to that of the spinal cord. Given that this characteristic MRI feature was combined with imaging findings showing the involvement of two adjacent vertebrae and a posterior element, the diagnosis of spinal GCT should have been favored over eosinophilic granuloma and Ewing’s sarcoma.

Surgery is the treatment of choice for giant cell tumor of the spine because it should be completely removed; the recurrence rate is slightly increased in cases with extension of giant cell tumor into the spinal canal and paraspinous space [3].

In conclusion, although rare, a GCT should be considered in the differential diagnosis of vertebra plana, and especially when observing tumor involvement of the adjacent vertebra or paravertebral tumor expansion, as well as when the tumor’s signal intensity is similar to that of the spinal cord on the T2-weighted image.

References

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차장규 ∙ 신원한2 ∙ 박관웅2 ∙ 김희경3 ∙ 박재성 ∙ 이혜경

점진적으로 악화하는 하지의 쇠약감을 호소하는 19세 여자 환자의 증례를 보고한다. 단순사진과 CT 사진에서 첫 번째 흉추의 척추 편평소견이 보였다. 이 종괴는 MR 검사에서 척추관까지 침범하며 T1 저신호강도로 및 T2 강조 영상에서는 중등도 신호강도로 보였다. 조직학적 검사에서 이 종괴는 거대세포종으로 확진되었다. MR 영상은 침범 정도와 특징적인 신호강도를 보여줌으로써 거대 세포종과 다른 척추종양과 감별에 도움을 주는 최선의 영상검사이다.