Intraspinal choristoma in the lumbar region
A case report
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Abstract
Rationale: Intraspinal choristoma is a relatively uncommon intervertebral canal tumor. Prior to our reports, only 2 cases of intraspinal choristoma had been reported. Because this disease is not common and looks like a mass of fatty tissue on the magnetic resonance imaging (MRI), intraspinal choristoma can be easily misdiagnosed as teratomas or lipomas (like the case of this article presenting) without a pathology report. So if a lumber intraspinal lesion is discovered in a clinical examination, intraspinal choristoma should be considered as a differential diagnosis. We present a case of intraspinal choristoma that is unlike any other reported case.

Patient concerns: A 35-year-old woman with left lower extremity hypoesthesia and burning-like pain in the lumbar region for 1 month visited the local hospital for plain lumbar spine MRI. The patient was diagnosed with a lumbar space-occupying lesion. A second plain lumbar spine MRI scan and a MRI scan with enhancement were performed in our hospital to confirm the presence of a congenital lipoma in the spinal canal. A postoperative biopsy of the lumbar spinal mass indicated that the mass was an intraspinal choristoma located in the spinal canal.

Diagnosis: Intraspinal choristoma.

Intervention: The lesion was surgically removed, and follow-up plain and enhanced MRI images of the patient’s lumbar spine were obtained.

Outcomes: After surgery, the patient no longer experienced the burning pain in her lumbar region or the left lower extremity hypoesthesia when the patient was discharged. And there was no evidence of recurrence 2 years after the surgery.

Lessons: The MRI presentation of intraspinal choristoma is similar to intraspinal lipoma. Therefore, a pathological assessment is critical to provide an accurate diagnosis.

Abbreviations: JOA = Japanese Orthopaedic Association Scores for Assessment of Cervical Myelopathy, MRI = magnetic resonance imaging.

Keywords: choristoma, congenital intraspinal tumours, intraspinal choristoma

1. Introduction
Choristoma is a tumor-like structure composed of normal tissue located at an abnormal site.[1] While it can occur anywhere in the body, it is more frequently observed in the head and neck region, especially in the pituitary gland[2] and oral cavity.[3] Intraspinal choristomas are rare and, to the best of our knowledge, only 2 cases have been reported.[4,5] Thus, it is pretty rare for choristoma to occur in the spinal canal. The postoperative pathological assessment of an intravertebral occupying lesion in our patient showed a case of intraspinal choristoma. In addition, this patient also had a combination of spinal cord deformities such as spinal cord tethering syndrome which makes this case unique.

2. Case Report
2.1. Clinical summary
A 35-year-old woman was diagnosed with an intraspinal space-occupying lesion after a magnetic resonance imaging (MRI) scan of her lumbar spine was performed at a local hospital. The presenting complaint was hyperalgesia of the left lower extremity and burning pain in the lumbar region for 1 month. The patient was admitted to our hospital for further diagnosis and treatment. The preoperative physical examination revealed scoliosis, pain elicited with positive pressure on both sides of the fourth lumbar vertebra, and bilaterally diminished Achilles tendon and knee tendon reflexes. The straight-leg raising test...
was positive. The extension-hallucis test was positive, and the tension test for the femoral nerve was negative.

MRI of the lumbar spine revealed that a near-circular short T1 and a long T2 signal shadow with clear borders were observed at the level of the inferior border of the fourth lumbar vertebral body. A low signal of Short Time Of Inversion Recovery demonstrated a rounded, non-enhanced area at the level of the lower edge of the fourth lumbar vertebral body. The area exhibited clear borders and was approximately 1.0 cm in diameter within the spinal canal (Figs. 1 and 2). In addition to intraspinal choristoma, other spinal cord deformities were observed on the sagittal MRI images in this patient: the spinal cord extended to the level of the lower edge of the fourth lumbar vertebra, which was distinctly abnormal because the normal adult spinal cord does not extend beyond the lower edge of the first lumbar vertebra, and the spinal cord was irregularly divided into 2 parts beginning at the level of the lower border of the 11th thoracic vertebral body, ending at the lower border of the fourth lumbar vertebral body. These images indicated the presence of a tethered spinal cord, resulting in shortened spinal cord end filaments. Furthermore, the corresponding region of the spinal canal and the subarachnoid space were widened significantly, suggesting the presence of a possible developmental abnormality associated with the spinal canal. Therefore, this patient also was diagnosed with tethered spinal cord syndrome and spinal cord deformities.

The pre-surgical Japanese Orthopedic Association Score (JOA score) was 14. Preoperatively, the lesion was thought to be a lumbar intraspinal lipoma. The mass was surgically removed under general anesthesia. During the surgery, the tumor appeared yellow and exhibited a tough texture. The tumor size was 1.5 cm × 1 cm × 0.4 cm; the color was primarily gray to yellow; the appearance was nodule-like, soft, with a slightly tough localized region of gray-white color. A small amount of bone tissue was present. The microscopic examination revealed the following information: mature fibrous and fatty tissues, blood vessels, and epithelium were present; small sweat gland ducts also were seen. The histological pattern resembled an intraspinal choristoma (Fig. 4). Immunohistochemical features were assessed based on immunohistochemical staining, including positive staining for S-100 (Fig. 5), GCDFP-15 (Fig. 6), CK7 (Fig. 7), CK8 (Fig. 8), P63 (Fig. 9), and AR (Fig. 10). We summarized the comparison of results of the immunohistochemical features of choristoma and hamartoma in Table 1.

### 2.2. Pathological findings

The histological examination provided the following information: the tumor size was 1.5 cm × 1 cm × 0.4 cm; the color was primarily gray to yellow; the appearance was nodule-like, soft, with a slightly tough localized region of gray-white color. A small amount of bone tissue was present. The microscopic examination revealed the following information: mature fibrous and fatty tissues, blood vessels, and epithelium were present; small sweat gland ducts also were seen. The histological pattern resembled an intraspinal choristoma (Fig. 4). Immunohistochemical features were assessed based on immunohistochemical staining, including positive staining for S-100 (Fig. 5), GCDFP-15 (Fig. 6), CK7 (Fig. 7), CK8 (Fig. 8), P63 (Fig. 9), and AR (Fig. 10). We summarized the comparison of results of the immunohistochemical features of choristoma and hamartoma in Table 1.

### 3. Discussion

Based on the pathological diagnosis, the features of choristoma and hamartoma can be confounding. Both are rare, and many clinicians and pathologists find it challenging to identify them accurately. In addition, many articles have mentioned the need to be able to differentiate between choristoma and hamartoma. However, none of the previous reports have clarified the critical features that facilitate differentiation. The term choristoma is derived from the Greek root “choristos,” meaning separated and also means mass or tumor. The term is synonymous with heterotopia and ectopia, which are defined as the proliferation of normal cells in an abnormal location and an ectopic, histologically normal mature tissue. Hamartoma is defined as a lesion containing mature and well-differentiated ectoderm and mesoderm derivatives that appear in abnormal locations; usually, 1 component dominates. Hamartoma is similar to choristoma in that, both are benign proliferations of mature tissue, but a hamartoma is a malformation resembling the tissue at its site of origin, while the choristoma contains components that do not usually appear at the anatomic location where it is located. Therefore, choristoma can be considered to be a subtype of hamartoma. Thus, a “hamartoma” confirmed histologically...
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To contain ectopic tissue is actually a choristoma. Castillo et al noted in an article on midline spinal cord hamartomas that only hamartomas that contained local elements in a disorganized fashion could be considered pure hamartomas, while those with normal but ectopic tissues (including glands, lymphoid tissue, and urinary tract tissues) should be considered choristomas. Malela et al reported a sporadic case of “Congenital Midline Spinal Hamartoma in a 5-Month-Old Infant in 2020.” The similarity with our case is that it also contained numerous adipocytes, however, the case reported by Malela et al did not exhibit cartilage, skeletal muscle, mature collagen fibers, nerve and vascular components in the normal tissue at the location of the choristoma. The histologic examination of our case revealed mature fibrous tissue and squamous epithelium, as well as small sweat gland ducts. These 3 mature tissue components should not be present in the spinal canal, so the histologic examination indicated that a choristoma was present in the patient in this report.

Since choristoma is extremely rare, it is not usually suspected before the surgery and pathology. Alternative differential diagnoses include teratoma and intraspinal lipoma. It was noted that

Figure 3. (F and G) Images taken 2 y after the surgery. The lumbar spine plain MRI scan in sagittal section (F) and cross-section (G). MRI = magnetic resonance imaging.

Figure 4. (H) Microscopic image of the tumor tissue with H&E staining (×100 magnification). (I) Another microscopic image at a magnification of ×200. (H and I) Mature fibrous and adipose tissue, blood vessels, epithelium, small sweat gland ducts, and a small amount of squamous-like tissue.

Figure 5. Immunohistochemical staining for S-100 protein (H&E ×100).
the preoperative imaging presentation of this patient was similar to intraspinal lipoma. Teratomas contain cell populations from at least 2 of the 3 primary germ layers, endoderm, mesoderm, and ectoderm. They form during embryonic development and may be diagnosed at any age. Teratomas are thought to result from the failure of primitive germ cells to migrate and usually occur along the body midline, anywhere from the coccyx to the pineal gland. The primitive germ cells fail to degenerate and subsequently proliferate in abnormal locations in the midline to form tumors. Thus, the pluripotent capacity of these cells gives rise to multiple tissues consisting of immature cells that have the potential for malignant degeneration. There are 4 histological teratoma variants: mature teratoma, immature teratoma, malignant teratoma, and monodermal teratoma. Intraspinal lipoma also is a rare benign tumor, accounting for <1% of all spinal cord tumors. Lesions often are located in the lumbosacral region, while occurrence within the dura of the cervical and thoracic medulla is less common. Though the pathogenesis of intradural lipomas remains unclear but occasional association with other midline anomalies points to a developmental origin. Intraspinal lipomas often occur in conjunction with a tethered spinal cord, and the lesion may contain other types of tissues such as bone, cartilage, or even bone marrow. Ammerman et al had described the clinicopathological characteristics of intramedullary lipomas: the tumors tend to be subpial in origin, involve multiple spinal cord segments as well as adjacent nerve roots, and have no clear plane demarcating themselves from adjacent neural parenchyma. Histologic investigation classically demonstrates normal adipose cells with intervening collagen. The collagen is often in continuity with
Table 1
Summary of immunohistochemical features of reported choristoma and hamartoma cases in the literature.

| Diagnosis                                      | Phakomatous choristoma 1 | Phakomatous choristoma 2 | Cartilaginous choristoma | Lacrimal gland choristoma | Palatal osseous choristoma | Mammary choristoma | Intraspinal choristoma | Pancreatic endocrine heterotopia | Leiomyomatous hamartoma | Hamartoma of mature cardiac myocytes | Pulmonary hamartoma | Schwann cell hamartoma | Sinonasal seromucinous hamartoma |
|------------------------------------------------|--------------------------|--------------------------|--------------------------|---------------------------|---------------------------|---------------------|------------------------|---------------------------------|-----------------------------|---------------------------------|-------------------------|-------------------------|-----------------------------|
| Reference                                      | S D et al[6]             | SM R et al[7]            | GW Pereira et al[9]      | A T-E et al[10]           | R S, et al[11]            | Chen X, et al[12]  | PF C, et al[13]         | Zhang M, et al[14]               | JC M-H, et al[15]               | Li Y, et al[16]                  | YW H, et al[17]              |
| S-100                                          | Positive                 | Positive                 | Positive                 | Negative                  | Positive                 |                     | Positive                | Positive                         | Positive                     | Positive                          | Positive               | Positive               | Positive                       |
| EMA                                            | Negative                 | Negative                 | Positive                 | Negative                  | Positive                 |                     | Positive                | Positive                         | Positive                     | Positive                          | Positive               | Positive               | Positive                       |
| desmin                                         |                          |                          |                          |                           |                          |                     | Positive                | Positive                         |                            | Some cases positive             |                       |                       |                              |
| AE1/AE3                                        | Negative                 | Positive                 | Positive                 | Positive                  |                          |                     | Positive                | Negative                         |                            | Some cases positive             |                       |                       |                              |
| SMA                                            | Negative                 | Positive                 | Positive                 | Positive                  |                          |                     | Positive                | Positive                         |                            | Some cases positive             |                       |                       |                              |
| CO34, p40, p63                                 | Negative                 | Positive                 | Positive                 | Positive                  |                          |                     | Positive                | Positive                         |                            | Focal positivity               |                       |                       |                              |
| Synaptophysin                                   | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Cytokeratin                                    | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Synaptophysin                                   | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Cytokeratin                                    | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| CK5/6                                          | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Ck7                                            | Positive                 | Positive                 | Positive                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Vimentin                                       | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Glial fibrillary acidic protein Melan-A         | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Actinin                                        | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| CAM5.2                                         | Positive                 | Positive                 | Positive                 | Positive                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| NSE                                            | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |
| Actin muscle-specific antibody Calretinin      | Negative                 | Negative                 | Negative                 | Negative                  |                          |                     | Positive                | Positive                         |                            |                               |                       |                       |                              |

(Continued)
| Diagnosis                                      | Reference                        | CK27 | TTF-1 (SPT24) | Trpsin | Insulin | Glucagon | Somatostatin | CD31 | ERG |
|-----------------------------------------------|----------------------------------|------|----------------|--------|---------|----------|--------------|------|-----|
| Phakomatous choristoma①                       | S D et al[6]                    |      | Negative       | Negative| Positive| Positive   | Negative      |      |     |
| Phakomatous choristoma②                       | SM R et al[7]                   |      | Negative       | Positive| Positive| Positive   | Negative      |      |     |
| Cartilaginous choristoma                      | GW Pereira et al[8]             |      | Positive       | Negative|          |          |               |      |     |
| Lacrimal gland choristoma                     | A T-E et al[9]                  |      | Negative       | Positive| Positive| Negative   | Negative      |      |     |
| Palatine osseous choristoma                   | R S et al[10]                   |      | Negative       | Positive| Positive| Negative   | Negative      |      |     |
| Mammary choristoma                            | PF C, et al[11]                 |      | Negative       | Positive| Positive| Negative   | Negative      |      |     |
| Intraspinal choristoma                        | DJ R et al[12]                  |      | Negative       | Positive| Positive| Negative   | Negative      |      |     |
| Pancreatic endocrine heterotopia               | Zhang M et al[13]               |      | Negative       | Positive|          |          |               |      |     |
| Leiomyomatous hamartoma                       | JW M-H et al[14]                |      | Negative       | Positive|          |          |               |      |     |
| Hamartoma of mature cardiac myocytes          | Y L et al[15]                   |      | Negative       | Positive|          |          |               |      |     |
| Pulmonary hamartoma                           | YW H et al[17]                  |      | Negative       | Positive|          |          |               |      |     |
| Schwann cell hamartoma                        |                                 |      | Negative       | Positive|          |          |               |      |     |
| Sinonasal seromucinous hamartoma              |                                 |      | Negative       | Positive|          |          |               |      |     |

| Diagnosis                                      | Reference                        | Pan-CKs | MUC5AC | Napasin A | MUC1 | MUC2 | MUC6 | CD7 | CD19 | Proliferating cell nuclear antigen | Estrogen receptor | Progesterone receptor | K67 | GSA | CK (glandular) | Neurofilament protein | Claudin-1 | Glucose | Melan-A | HM3-45 | c-Kit | Chromogranin A |
|-----------------------------------------------|----------------------------------|--------|--------|------------|------|------|------|-----|-----|----------------------------------|------------------|---------------------|-----|-----|-------------|---------------------|----------|--------|----------|--------|-------|----------------|
| Phakomatous choristoma①                       | Kurman et al[6]                 |        | Positive| Positive   | Negative| Negative| Negative| Positive| Positive| Positive |
| Phakomatous choristoma②                       | Seyed et al[7]                  |        | Positive| Negative   | Negative| Negative| Negative| Positive| Positive| Positive |
| Cartilaginous choristoma                      | GW P et al[8]                   |        | Positive| Negative   | Negative| Negative| Negative| Positive| Positive| Positive |
| Lacrimal gland choristoma                     | A T-E et al[9]                  |        | Positive| Negative   | Positive| Positive| Negative| Positive| Positive| Positive |
| Palatine osseous choristoma                   | R S, et al[10]                  |        | Positive| Positive   | Positive| Negative| Negative| Positive| Positive| Positive |
| Mammary choristoma                            | Chen et al[11]                  |        | Positive| Positive   | Positive| Negative| Negative| Positive| Positive| Positive |
| Intraspinal choristoma                        | PF C, et al[11]                 |        | Positive| Positive   | Positive| Negative| Negative| Positive| Positive| Positive |
| Pancreatic endocrine heterotopia               | M S, et al[11]                  |        | Positive| Positive   | Positive| Positive| Positive| Positive| Positive| Positive |
| Leiomyomatous hamartoma                       | M Z, et al[13]                  |        | Positive| Positive   | Positive| Positive| Positive| Positive| Positive| Positive |
| Hamartoma of mature cardiac myocytes          | JW M-H, et al[14]               |        | Positive| Positive   | Positive| Positive| Positive| Positive| Positive| Positive |
| Pulmonary hamartoma                           | G R[15]                        |        | Positive| Negative   | Negative| Negative| Negative| Positive| Positive| Positive |
| Schwann cell hamartoma                        | Y L et al[15]                   |        | Positive| Negative   | Negative| Negative| Negative| Positive| Positive| Positive |
| Sinonasal seromucinous hamartoma              | YW H, et al[17]                 |        | Positive| Negative   | Negative| Negative| Negative| Positive| Positive| Positive |

EMA = epithelial membrane antigen, HMWK = high molecular weight keratin, NSE = neuron-specific enolase, SMA = smooth muscle actin.
approximately 16 months, without evidence of recurrence or residual cases in children, the tumor was incompletely excised. The mass is more likely to be a benign tumor. Therefore, the results that were obtained confirmed the diagnosis of choristoma. We also suggest that it was not a coincidence that the patient also exhibited spina bifida and spinal cord tethering, as an association between spinal dysraphism and the occurrence of a lipoma is possible. The etiology of congenital spina bifida is not fully understood. However, most scholars believe that abnormal embryonic development of the neural tube is responsible for spina bifida, and abnormalities in the fusion process result in its formation. Specifically, the neural tube separates from the ectoderm through a process of detachment. After separation, midline fusion of the ectoderm above the neural tube occurs.[21]

Besides, a case of autograft-derived spinal cord mass following olfactory mucosal cell transplantation in a spinal cord injury patient[22] enlightened us that the possible etiology of this intraspinal choristoma is a malformation during the embryonic development, and the malformation results in a transplantation of the regional disordered embryonic cells in lumbar region. Therefore, we believe that the abnormal developmental process of spinal bifida is most likely the reason for the development of intraspinal choristoma in our case. Based on this case, if a space-occupying lesion in the spinal cord with a predominantly fatty signal appears on the MRI, surgeons should carefully distinguish whether the boundaries between the occupying lesion and the surrounding tissues are distinct. If the lesion has definitive boundaries and the patient has congenital spina bifida or spinal cord embolism, a possible diagnosis of intraspinal choristoma should be considered. Surgical options include complete resection, and the prognosis for patients with intraspinal choristoma is generally considered to have a lower recurrence rate than an intraspinal lipoma.

In general, the clinical course and histopathological features of intraspinal choristoma indicate its benign nature. In 2 reported cases in children, the tumor was incompletely excised. One child was followed for 7.5 years and the other for approximately 16 months, without evidence of recurrence or residual tumor[20]. Because choristoma is classified as a benign tumor, many studies do not discuss recurrence. The case reported in this study showed benign tumor characteristics based on the preoperative MRI findings and the intraoperative observations. We followed the patient in this case for 2 years, and the patient did not exhibit any recurrence, which indicated that surgical excision is curative, with a low incidence of recurrence. However, a case of osseous choristoma that exhibited recurrence was reported in another article, which might have been due to the tumor having a greater degree of malignancy. Typically, tumors that are multiple and have no pedunculated mass surrounded by soft tissue, such as within the buccal mucosa or muscle, might have the possibility of recurrence.[21]

4. Conclusion

Intraspinal choristoma is a rare congenital intraspinal tumor that might correlate significantly with congenital spinal deformities. Intraspinal choristomas are not easily distinguishable from intraspinal lipomas and intraspinal hamartomas based on MRI. However, if the lesion has distinct boundaries and the patient has congenital spina bifida or spinal cord embolism, a possible diagnosis of intraspinal choristoma should be considered. With the presence of distinct tumor boundaries, surgical options for complete resection should be considered. The prognosis of a lower recurrence rate for an intradural psammoma is generally better than for an intraspinal lipoma.

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